

TURN "ON" YOUR LONGEVITY GENES

Life Extension

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The ULTIMATE Source For New Health And Medical Findings From Around The World

February 2010

Neutralize the Lethal Effects of Excess Calories

**Reverse
Mitochondrial
Dysfunction**

**Delay the
Absorption
Of Sucrose**

**Cutting-Edge
Prostate
Defense**



PLUS-

**THE MOST COMPREHENSIVE WEIGHT LOSS PROGRAM
ELEVATED HOMOCYSTEINE DOUBLES ALZHEIMER'S RISK
SOY MAY PREVENT COLON CANCER**

SUPER SALE

The annual **SUPER SALE** enables members to obtain premium grade supplements at prices substantially **below** what commercial companies charge.

When members buy products from the **Life Extension Foundation Buyers Club**, they know that the **quality** of the products are backed by the organization's commitment to achieving an indefinitely extended life span. What follows are a few examples of the **savings** members enjoy during the **SUPER SALE**.

Retail	Member SUPER SALE Discount Price Per Bottle
Super Omega-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract 120 softgels Super-refined highly concentrated EPA/DHA fish oil plus sesame lignans and more potent olive fruit extract. Item # 01482	\$32 \$16.81 (10-bottle purchase)
Life Extension Mix™ 315 tablets High-potency multi-nutrient formula now with twice as much vitamin D and iodine, plus pterostilbene to favorably regulate gene expression. Item # 01455	\$98 \$46.91 (10-bottle purchase)
Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™ 100 mg, 60 softgels Preferred "ubiquinol" combined with shilajit, an organic compound shown to double levels of CoQ10 in the mitochondria. Item # 01426	\$62 \$35.10 (10-bottle purchase)
Super Booster Softgels with Advanced K2 Complex 60 softgels Critical, oil-based nutrients including gamma-tocopherol, sesame lignans, lycopene, lutein, ginkgo, chlorophyllin, selenium, and both forms of vitamin K2. Item # 01380	\$42 \$25.65 (four-bottle purchase)
Super Zeaxanthin with Lutein & Meso-Zeaxanthin plus Astaxanthin 60 softgels Zeaxanthin, lutein, meso-zeaxanthin to support macular density, and astaxanthin to reduce eye fatigue. Item # 01286	\$42 \$25.65 (four-bottle purchase)
Bone Restore 150 capsules High-potency bone protection formula with FruiteX B® OsteoBoron®. Item # 00811 FruiteX B® and OsteoBoron® are registered trademarks of VDF Pharmaceuticals, Inc. US patent #5,962,049.	\$22.50 \$13.16 (four-bottle purchase)
Super Bio-Curcumin® 60 vegetarian capsules Absorbs up to seven times better than conventional curcumin. Item # 00407	\$30 \$17.89 (four-bottle purchase)
Ultra Natural Prostate Formula* 60 softgels Now with more potent Saw Palmetto to support normal urinary flow and help support healthy prostate function. Item # 01475 *Not available for export	\$38 \$21.60 (12-bottle purchase)
Calorie Restriction Mimetic Formula 60 vegetarian capsules Provides in two capsules, 250 mg of trans resveratrol , the highest dose (3 mg) of pterostilbene , more quercetin , plus beneficial grape seed and black tea polyphenols , all to help favorably support healthy gene expression observed in response to calorie restriction . Item # 01419	\$36 \$22.28 (four-bottle purchase)

These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

The **SUPER SALE** extends to **February 1, 2010**. Members traditionally take advantage of the **SUPER SALE** to stock up on a year's supply of their favorite supplements. To place your order, **call 1-800-544-4440** or visit www.lef.org (**SUPER SALE** pricing available only to members in the US, Canada, and England.)

REPORTS



30 ACTIVATE YOUR LONGEVITY GENES

Modern science has only recently begun to unravel the mystery behind how calorie restriction extends life span and improves overall health. One of the most astounding revelations about calorie restriction is its power to turn on "youthful" genes and turn off "senescent" genes. Most people, however, find it difficult to sufficiently reduce the amount of food they consume. The incredible news is that **five natural compounds** have been identified that simulate many of the favorable **gene expression** changes observed during **calorie restriction**.



42 BLOCK ABSORPTION OF KILLER CARBOHYDRATES

An estimated **60 million** American adults suffer from **abnormally high blood sugar**—a pre-diabetic state that can sharply increase degenerative disease risk. Many of these people have *lost* their **metabolic capacity** to process even a low carbohydrate diet. A novel group of natural carbohydrate enzyme-inhibitors offers aging individuals a practical way to control glucose levels, improve blood markers of health, and regain glycemic balance.



54 REVERSE MITOCHONDRIAL DYSFUNCTION

Age-related deterioration of the *mitochondria* in our cells is associated with an array of deadly conditions, ranging from senility to diabetes and heart failure. Researchers have recently discovered that this deadly process can be *reversed* with cellular energizers that help restore and preserve **mitochondrial function**.



72 HIGH-TECH PROSTATE DEFENSE

Prostate disorders remain a modern scourge, with *half* of adult men eventually suffering benign prostatic hyperplasia (enlarged prostate), and **186,000** developing prostate cancer each year. An advanced extraction technology now delivers even higher concentrations of the active ingredients in saw palmetto. Here we review the most effective natural compounds to support the aging prostate gland.

DEPARTMENTS



23 IN THE NEWS

Antioxidants could offer protection to lungs during flu season; long-term high dose vitamin D well tolerated; green tea may help protect against oral cancer; soy shows promise for colon cancer prevention; and more.

SUPERFOODS: STRAWBERRIES 85

Strawberries are a nutrition power-house with major anti-inflammatory properties. They also have potent cancer protection and heart health benefits.

WELLNESS PROFILE 89

Ann Fonfa's personal quest to get answers about effective complementary alternative medicines to battle cancer resulted in the creation of The Annie Appleseed Project, a world renowned cancer education center.



7 ON THE COVER

AS WE SEE IT

ADD YEARS TO YOUR LIFE: THREE WAYS TO ACHIEVE THE ANTI-AGING BENEFITS OF CALORIC RESTRICTION

Research conducted over the last 75 years in mammals confirms that consuming fewer calories induces *extraordinary* anti-aging effects, from radically extended life span to reduced disease incidence and improved blood markers of health.

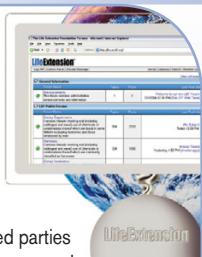
A recently completed 20-year primate study provides the most compelling evidence to date that even *modest* caloric restriction may bring about these benefits to humans. In this issue, we offer *practical* ways to trigger the same physiological mechanisms as caloric restriction—without having to endure drastic dietary deprivation.



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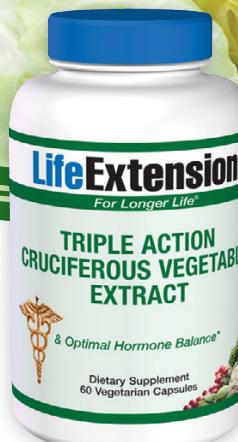
NEW TRIPLE ACTION CRUCIFEROUS Vegetable Extract with Apigenin

Scientists have identified specific extracts from **cruciferous vegetables**—such as broccoli, cauliflower, cabbage and Brussels sprouts—that help maintain healthy hormone levels. Maintaining optimal hormone balance is essential to any anti-aging strategy. **Triple Action Cruciferous Vegetable Extract** combines these plant extracts into the most comprehensive food-based, plant compilation for cell protection yet.

I3C (indole-3-carbinol) and **DIM (di-indolyl-methane)** favorably modulate estrogen metabolism and induce liver detoxification enzymes to help neutralize potentially harmful estrogen metabolites and xenoestrogens (potentially toxic, estrogen-like environmental chemicals).¹⁻⁴

Extracts of **broccoli**, **watercress**, and **rosemary** provide **glucosinolates**, **isothiocyanates**, **carnosic acid**, and **carnosol**—bioactive compounds that have a multitude of favorable effects on estrogen metabolism and cell division.⁵⁻⁸ **Apigenin**, a powerful plant flavonoid found in plants such as **parsley** and **celery**, is also added to the formula to boost cell protection,⁹ while 25 mg of a natural source of benzyl isothiocyanate (BITC), are included to maintain cell health.¹⁰

For those weighing less than 160 pounds, just **one** capsule a day provides optimal potencies. Those weighing over 160 pounds should consider taking two capsules a day. A 60-capsule bottle of **Triple Action Cruciferous Vegetable Extract** retails for \$24. If a member buys four bottles during **Super Sale**, the price is reduced to only **\$14.85 per bottle**.



ITEM #01468



ITEM #01469

The new Triple Action Cruciferous Vegetable Extract provides the following concentrates in just one capsule:

Broccoli Super Concentrate.....	400 mg
[standardized to 4% glucosinolates (16 mg)]	
Watercress 4:1 extract.....	50 mg
Indole-3-Carbinol (I3C).....	80 mg
Rosemary Extract.....	50 mg
Cat's Claw Extract.....	50 mg
Cabbage Extract.....	25 mg
DIM (di-indolyl-methane).....	14 mg
Apigenin.....	25 mg

Those who want to obtain the benefits of **resveratrol** can order **Triple Action Cruciferous Vegetable Extract with Resveratrol**. Each capsule provides **20 mg** of **resveratrol** in addition to the **vegetable extracts** and retails for \$32 per 60-capsule bottle. When a member buys four bottles during **Super Sale**, the price is reduced to only **\$19.98 per bottle**.

REFERENCES:

1. *Biochem Pharm*. 2002; 64:393-404.
2. *Toxicol Appl Pharm*. 2001 Jul 15;174(2):146-52.
3. *J Natl Cancer Inst*. 1997 May 21;89(10):718-23.
4. *Cancer Detect Prevent*. 2004;28:72-9.
5. *Carcinogenesis*. 2002 Apr;23(4):581-6.
6. *Mol Cancer Ther*. 2003 Oct;2(10):1045-52.
7. *Carcinogenesis*. 1998 Oct;19(10):1821-7.
8. *Carcinogenesis*. 1995 Sep;16(9):2057-62.
9. *J Clin Biochem Nutr*. 2009 May;44(3):260-5.
10. *Food Chem Toxicol*. 2008 Jul;46(7):2358-64.

To order the New Triple Action Cruciferous Vegetable Extract, call 1-800-544-4440 or visit www.LifeExtension.com

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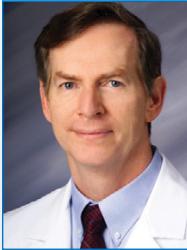
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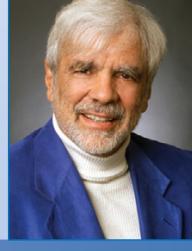
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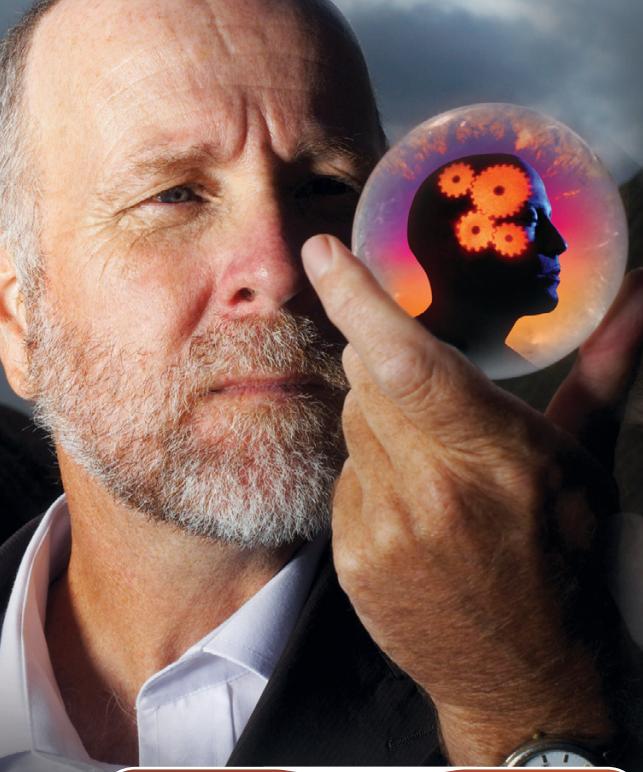
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Phosphatidylserine-DHA (PS-DHA)	100 mg
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Pregnenolone	50 mg
Vinpocetine	20 mg
Phospholipid-Grape Seed Extract	150 mg
Wild Blueberry	150 mg
(Vaccinium angustifolium) 130:1 Extract	
Sensoril® Ashwagandha Extract	125 mg
(Withania somnifera)	
Uridine-5'-Monophosphate (disodium)	50 mg
Proprietary NeuroProtection Complex Blend	125 mg
Perluxan® Hops Extract (Humulus lupulus)	
Ginger (Zingiber officinale) Extract	
Rosemary (Rosmarinus officinalis) Extract	



Item #00921



Item #00922

Diminished levels of neurotransmitters and other brain compounds profoundly affect cognition and memory in aging adults. **Cognitex** was developed in 1982 to increase brain levels of **acetylcholine** — a neurotransmitter that enables neurons to communicate. Over the years, **Cognitex** has been improved with the addition of nutrients used in **Europe** to protect and enhance neurological function.



Cognitex with Pregnenolone & NeuroProtection Complex provides the following **scientifically validated** nutrients to provide broad-spectrum neurological support:

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- **Sharp-PS® GOLD**, a rich **phosphatidylserine** compound bound to **DHA**, promotes already-normal neuronal cell membrane function and structure.
- **Alpha-glyceryl phosphoryl choline (A-GPC)** boosts acetylcholine, a critical neurotransmitter that declines with age.
- **Vinpocetine** enhances circulation and oxygenation to brain cells, improves neural electrical conductivity, and protects against neuron-destroying excitotoxicity.
- **Phospholipid-grape seed extract** bound together results in a compound that is better absorbed into the bloodstream where it improves blood vessel tone and elasticity, thus enhancing blood circulation and oxygen flow to the brain.
- **Wild blueberry extract** protects against free-radical damage in the brain and helps maintain fluid balances already within the normal range.
- **Sensoril® ashwagandha extract** helps alleviate mental fatigue by inhibiting an enzyme (acetylcholinesterase) that degrades acetylcholine in the aging brain.
- **Proprietary NeuroProtection Complex Blend** contains standardized extracts of **hops**, **ginger**, and **rosemary** — proven to help with inflammation.
- **Pregnenolone** is a hormone that may be especially beneficial to the brain.

The retail price for 90 softgels of Cognitex (with or without pregnenolone) is \$74 (item #00922) and \$72 (item #00921), respectively. If a member orders four bottles of either version during **Super Sale**, the price per bottle is reduced to just **\$44.96** and **\$43.20**, respectively.

Caution: Cognitex is also available without pregnenolone for those with existing steroid hormone-sensitive cancer.

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To order Cognitex, call 1-800-544-4440 or visit www.LifeExtension.com

Reap Benefits of Calorie Restriction without Chronic Hunger



BY WILLIAM FALOON

The most significant **anti-aging discovery** in history was made in **1935**. In that year, rats fed a **calorie restricted** diet achieved radically-extended mean and maximum life spans, along with a delayed onset of age-related diseases.¹

Since this finding was published 75 years ago, dozens of experiments in mammals have validated that **undernutrition without malnutrition** induces **profound** anti-aging effects.²⁻¹⁰ Not only do calorie restricted animals live much longer, but they remain far healthier than normally-fed controls.

When a group of humans consumed a similar calorie restricted diet, their conventional blood markers of aging (excess glucose, cholesterol, triglycerides, LDL) plummeted to much lower levels.^{11,12}

In the most significant finding to date, two groups of Rhesus monkeys were studied for **twenty years**. The group placed on a **moderately** restricted diet reduced their incidence of age-related disease by a **factor of three!** Cancers and cardiovascular disease were **less than half** in the **moderate** calorie restricted group compared to controls. None of the moderate calorie restricted group developed **diabetes** or impaired glucose tolerance, despite a usually high prevalence in these monkeys. An interesting finding showed that *moderate* calorie reduction preserved **brain volume** in certain regions.¹³ Normal aging is accompanied by brain **shrinkage** as neurons are lost.

This issue of **Life Extension®** is dedicated to providing practical options to attain many of the beneficial effects of **calorie restriction**. Our objective is to help induce the favorable changes in our members' bodies that may be responsible for the remarkable age-delaying and disease-protecting effects of reduced calorie intake.



A Time to Make Personal Commitments

After reading this article, I hope each of you will reduce some of the excess calories you ingest each day. I can almost promise that if you do it right, you'll never miss these surplus **age-accelerating** sugars and fats. Even if you are unable to cut a single calorie, we are going to show how you can benefit nonetheless.

A study of those making New Year's resolutions showed that **46%** are compliant after six months.¹⁴ During this time of year, when people strive for a healthier and longer life, I believe most of you will succeed in cutting back at least *some* excess food intake.

We All Eat Too Much

Calorie-dense toxic foods are abundant, cheap and heavily advertised. It is no wonder that so many children and adults eat far **more** than what their bodies require.



Excess calorie intake causes our bloodstream to be chronically 'bloated' with glucose, insulin, cholesterol, fat, homocysteine, and other pro-inflammatory inducers. Persistent bloodstream overload predisposes us to cancer, stroke, heart attack, senility, painful inflammation and virtually every other age-related ailment.¹⁵⁻²⁷

Consuming excess calories shortens our *average* life span by facilitating the development of *age-related diseases* that preclude us from attaining healthy longevity.

Overconsumption of calories has another insidious effect. Many scientists believe that our *life span* is largely controlled by **genes** that program our bodies to function in a youthful-healthy state. When we consume excess calories, we cause some of these genes to turn against us and contribute to *accelerated aging and death*.

If we reduce the amount and/or effects of ingested calories, the result is a more *favorable* gene

expression profile. Such a positive effect helps us to live longer, healthier lives.

Controlling the impact of *ingested* calories becomes an essential component of a science-based longevity program, as excess calorie exposure reduces life span by increasing degenerative disease risk and accelerating aging.

So now that you understand that most of us are eating too much, let's talk about **practical** approaches to doing something about it.

Reduce the Number of Calories You Absorb

Foods eaten must first be broken down by *digestive enzymes* before they are absorbed into our bloodstream. Dietary fats, for instance, are broken down by **lipase** enzymes in the stomach and small intestines. Taking 120 mg of the *lipase-inhibiting* drug **orlistat** before meals reduces dietary fat absorption by **30%**.²⁸

Green and **black tea polyphenols** also inhibit **lipase**, thus enabling one to eat more calories without **absorbing** all of the fats.^{29,30} In fact, when theaflavins from black tea were administered to rats, there was an immediate **suppression** of post-meal (post-prandial) triglyceride elevation in the blood. The scientists attributed these results to the "*inhibition of pancreatic lipase activity*."³⁰

You might think that a drug like **orlistat** that reduces fat absorption by **30%** would induce significant **weight loss**. The harsh reality is that study subjects must reduce their dietary fat intake and take orlistat to lose just **20.5 pounds** in one year. These data reveal the frightening degree to which most of us overeat.³¹ In other words, even when dietary **fat** absorption



is reduced by 30%, we still take in too many fat calories.

Studies show that in response to consumption of **green** or **black tea polyphenols** (or orlistat), substantial reductions in blood glucose, triglycerides, cholesterol, and other vascular risk factors occur.³²⁻⁴¹ These same changes happen when one reduces calorie intake,^{11,12} suggesting that those who continue to eat too much should take steps to block **digestive enzymes** that enable excess calories to be absorbed.

Most Western diets contain too many refined **carbohydrates** that add to the calorie burden. Ingested **carbohydrates** are broken down for **absorption** by the enzymes **sucrase**, **amylase**, and **glucosidase**. One may obtain some of the effects of following a “low-carb” diet by taking 50-100 mg of a drug called **acarbose** (glucosidase inhibitor) before each meal.^{42,43}

Nutrients that may slow carbohydrate *absorption* include **white kidney bean** (amylase inhibitor),⁴⁴⁻⁴⁷ **InSea™** (containing amylase and glucosidase inhibitors from special seaweeds),⁴⁸⁻⁵¹ **Irvingia** (amylase inhibitor)⁵²⁻⁵⁵ and **L-arabinose** (sucrase inhibitor).⁵⁶ These carbohydrate-blunting **nutrients** are all contained in the new Optimized Irvingia formula.

Clinical studies confirm significant reductions in glucose, insulin, and triglyceride levels in response to taking **acarbose** alone.⁵⁷⁻⁵⁹ These effects also occur when one restricts their calorie intake. Clinical studies show dramatic reductions in heart attack rate in response to acarbose.⁵⁹ This same kind of vascular disease risk reduction has been observed in experimental studies where food intake is restricted.⁶⁰⁻⁶² What this indicates is that one may obtain some of the benefits of following a **calorie restricted** diet by inhibiting **lipase**, **amylase**, **glucosidase**, and **sucrase** enzymes.

Consuming soluble dietary fibers before meals slows the absorption of carbohydrates, thus blunting the postprandial insulin spike.⁶³⁻⁶⁶ This is another method of impeding calorie absorption and helping to mimic the effects of calorie restriction.

Improve Your Gene Expression

In response to consuming fewer calories, gene expression favoring youthful vigor is improved. Overeating, on the other hand, induces pathologic gene expression favoring the onset of diseases and accelerated aging.^{72,73}

Until recently, the only way of achieving a favorable gene expres-

sion profile was through **caloric restriction**. Back in the late 1990s, research funded by the **Life Extension Foundation®** enabled scientists to compare the effects of various compounds (such as resveratrol) to the **gene expression** changes that occur during calorie restriction.

These experiments have been used to identify **nutrients** that mimic many of the beneficial gene expression changes observed during **calorie restriction**. As you will read in this month's issue, many of the nutrients **Life Extension®** members have been taking for decades, and a few new ones, favorably influence **gene expression** changes that occur during calorie restriction.

Once scientists gain total control over the expression of the genes that affect longevity, mankind may achieve biological immortality. Until that time arrives, it is comforting to know that we can now exert at least some control over the expression of genes that influence our health and longevity.

With the discovery of **novel plant extracts** that mimic some of the **gene expression** changes observed in response to **calorie restriction**, one may enjoy many of the longevity benefits of under-eating by taking the proper supplements daily.

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Moderate Calorie Restriction—It's Not as Difficult as You Think!

For too many people, copious food intake has become an addictive drug. I am so gratified that the nutrient **Irvingia** and the drug **metformin** keep my appetite under control and enable me to eat a lot less than I used to. There are other nutrients and drugs that effectively curb appetite, but these are all I have needed.

I don't miss gorging on surplus calories. One reason is my regular review of scientific articles that discuss reasons why people develop lethal diseases. For many medical conditions, **overeating** (described as "excess energy intake") is an **independent risk factor** for developing anything from cancer, to vascular disease, to arthritis, and even to senility.⁷⁴⁻⁷⁹

I regularly speak with individuals who practice what many would say is *extreme* calorie restriction. The media views these individuals as being so unique that they have been favorably featured on CBS News' *60 Minutes*, the *Oprah Winfrey Show*, and in *Newsweek* magazine.

When I asked two of the most famous calorie restrictors how many calories they consume a day, Paul McGlothin told me he eats around **1,900**, while Meredith Averill's intake is around **1,600** daily. Based on these numbers, I did not think Paul or Meredith are overly depriving themselves. People who go on crash diets, sometimes consuming fewer than 1,000 calories a day, are absolutely miserable, and rapidly regain the weight they lost. Paul and Meredith, on the other hand, say they are activating their "happiness biochemistry" and would never consider eating more food that would compromise their mental state of well being and physical state of health.



Combating Carbohydrate Overload

Older people are more likely to suffer from **impaired glucose tolerance**, meaning their bodies are unable to properly utilize the sugars they ingest. This condition often manifests clinically as unwanted **weight gain, metabolic syndrome** and/or **type 2 diabetes**.^{67,68}

Though people try to adopt healthier eating patterns, their aging bodies let them down by losing cellular sensitivity to **insulin**. The result is that even when carbohydrate intake is limited, too much tissue-damaging **glucose** remains in the blood and then converts to excess body fat.

Other than those who aggressively practice calorie restriction, most aging people lack the **metabolic capacity** to process the carbohydrate calories they ingest each day. Fortunately, there are practical solutions to combating the lethal effects of carbohydrate overload.

We know that by interfering with the activity of certain carbohydrate-digesting **enzymes**, blood **glucose** levels **plummet**, along with vascular risk factors such as **triglycerides** and **C-reactive protein**.

The American diet is chronically overloaded with **carbohydrates** that play a role in virtually every age-related disease. Many people need to inhibit the activity of all **three** of the following **carbohydrate digestive enzymes** to regain glycemic control:

- 1) **Amylase**
- 2) **Glucosidase**
- 3) **Sucrase**

Fortunately, a low-cost drug called **acarbose** effectively inhibits glucosidase.^{42,43} A patented nutrient compound called **InSea™** inhibits amylase and glucosidase,⁴⁸⁻⁵¹ **white kidney bean extract**⁴⁴⁻⁴⁷ and **Irvingia** suppress amylase,⁵²⁻⁵⁵ while a new patented ingredient containing **L-arabinose** suppresses sucrase.⁵⁶ These carbohydrate enzyme inhibiting nutrients are available in the **new Optimized Irvingia** formula. Acarbose is available as a prescription drug.

Those who want to be even more aggressive in reducing excess glucose and insulin levels should ask their doctor to prescribe **metformin**.⁶⁹⁻⁷¹ This drug not only acts as an insulin sensitizer, but functions via additional mechanisms to reduce blood glucose loads. I personally take **850 mg** of metformin two to three times a day, though most people take only **250-500 mg** three times a day (before meals). Metformin is available as a very low-cost generic drug.

Your objective in combating **carbohydrate overload** is to drive fasting glucose levels below 86 mg/dL of blood. I suggest that you have your blood tested within two months of initiating a comprehensive program to make sure the nutrients, drugs, and/or lifestyle changes are working for you.

At the beginning of this article I described the Rhesus monkey study that utilized only *modest* calorie restriction to achieve remarkable longevity benefits. From everything we know today, it would appear that many of you could migrate towards a routine of at least **modest** calorie reduction and feel better doing so.

So much of our appetite craving is psychological. I was at a business dinner a while back and ordered an 8-ounce filet mignon. When I finished, I thought that I should have ordered a larger portion. Several nights later, I ordered a 12-ounce size at the same restaurant and felt uncomfortably

full. This made me realize how we “think” we need large portions of food to be satiated, when we really don’t. (I rarely eat beef by the way, usually only small portions a few times a month.)

It is hard for me to dine with regular people and not be appalled by the enormous amount of calories they ingest at a single meal. I know I feel better at the meal’s conclusion, while these individuals are bloated from gorging themselves with so many excess calories.

Once one accepts the fact that surplus food ingestion is **poison** to the body, it’s a lot easier to practice at least a **moderate** degree of caloric cutback.

Calorie Restriction without Hunger!

In a study published in the **Journal of the American Medical Association (JAMA)**, the effects of caloric restriction were measured in a group of overweight adults over a six-month period.⁸⁰

The findings showed that in response to reduced food intake, fasting **insulin** levels plummeted. Reducing excess insulin is important because “insulin overload” increases the risk of heart disease, cancer, blindness, stroke, Alzheimer’s, and other age-related diseases.⁸¹⁻⁸⁴

The most exciting finding of the JAMA study was the amount of weight lost in the groups that restricted their calorie intake. The **moderate** caloric-restriction group experienced a **24%** reduction in body fat mass, while the **very low**-calorie group achieved a **32%** reduction in fat mass.⁸⁰ You may wonder if you can replicate these findings, since hunger is the factor that precludes most people from staying on a low-calorie diet.

Fortunately, a plant extract discovered in Europe called **pinolenic acid** has been shown to suppress appetite without causing any stimulatory effect. This plant extract attacks the underlying mechanisms involved in hunger so effectively that study participants reduced their food intake by **36%**.⁸⁵

Pinolenic acid is a polyunsaturated fatty acid derived from pine nuts. It stimulates the secretion of the hunger suppressing hormones **cholecystokinin (CCK)**⁸⁶ and **glucagon-like peptide-1 (GLP-1)**. The results from human clinical studies using pinolenic acid reveal a reduced “desire to eat” along with early satiety.⁸⁵

Those seeking to curb their appetite to reduce daily calorie intake should take **three capsules** of a pinolenic acid-containing supplement called **Natural Appetite Control** before most meals.

Practical Options to Emulate Calorie Restriction

The scientific data solidly document that surplus calorie intake shortens your life span.^{87,88} Everyone responds differently when reducing food intake, but most people can at least modestly decrease the number of calories they consume each day.

Most individuals, however, will still eat more than they should. Fortunately, there are multiple ways to inhibit the activity of *digestive enzymes*, and thus spare most of the body from the lethal effects of excess calorie absorption. An aggressive three-month plan to reduce the number of absorbed calories involves taking the following nutrients and drugs before most meals:

- **Orlistat** 120 mg
(Inhibits *lipase* enzyme^{*28,31,35,40,41})
- **Acarbose** 50-100 mg
(Inhibits *alpha-glucosidase* enzyme^{42,43,57,59})
- **Optimized Irvingia** 2 caps
(Inhibits *amylase*, *alpha-glucosidase*, and *sucrase* enzymes, functions by additional mechanisms to facilitate calorie reduction^{29,44,48,52})
- * **Black tea theaflavins**
300-350 mg (Also inhibit *lipase*, but not as potently as orlistat³⁰)

For some individuals, drugs like **orlistat** that inhibit *lipase* will cause gastrointestinal discomfort after eating too much. One objective of suggesting a 90-day trial using this drug is to forcibly educate you about healthier lifelong dietary patterns. For example, if you continue to consume excess **fat** calories and take orlistat, the result will be lots of lipid-laden feces in the toilet after a bowel movement. Understanding that

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this fat appearing in your toilet was destined to be absorbed into your bloodstream (which could shorten your life span) may motivate you to eat less. Likewise, digestive discomfort associated with excess carbohydrate intake in the presence of *alpha-glucosidase*, *amylase*, and *sucrase* inhibitors may entice you to reduce intake of dangerous simple carbohydrates. In case you have not figured this out yet, a side benefit to doing all this is a loss of surplus body fat.

To mimic some of the favorable gene expression changes that occur in response to *calorie restriction*, a new nutrient formula called **Calorie Restriction Mimetic Formula** has been developed that consists of the following nutrients in the daily dose:

Black tea extract (<i>Camellia sinensis</i>)	300 mg
Quercetin	150 mg
Trans-Pterostilbene	3 mg
Trans-Resveratrol	250 mg
Oligomeric proanthocyanidins from grape seeds	50 mg

These and a number of other nutrients that health-conscious people already take, such as high dose **vitamin D3** and **fish oil**, have been shown to favorably alter gene expression.⁸⁹⁻¹⁰¹

We Live in an Exciting Era!

Scientists continue to validate the health benefits of nutrients that **Life Extension** members supplement with every day. Only recently, however, have these favorable effects been linked to the ability of these nutrients to mimic *gene expression* changes that occur in response to *calorie restriction*.^{72, 101-105}

An explosive volume of newly published research indicates that aging humans can exert tremendous degrees of **control** as to whether they maintain **youthful** or **senescent** gene expression patterns in their cellular DNA.¹⁰⁶⁻¹⁰⁸

Some youthful gene expression can be maintained by reducing intake of calories to a level **20-40% lower** than is typical, while still obtaining all the necessary nutrients and vitamins.^{13,109-113} The most recent study indicates even a **modest** calorie restricted diet produces **huge reductions** in degenerative disease in monkeys.¹³

Those who are unable to sufficiently cut their food intake can still inhibit the **absorption** of ingested calories by taking compounds (such as **orlistat**, **acarbose**, and/or the **new Optimized Irvingia** formula) that block *digestive enzyme* activity.

Finally, and most exciting of all, there are nutrients (such as **pterostilbene** and **resveratrol**) that

have been shown to mimic many of the favorable effects induced by **calorie restriction**.^{102-104,114-116} These low-cost nutrients provide humans with an unprecedented power to determine whether their cellular DNA expresses **youth-promoting** or **senescent-inducing** genes. Humans have never enjoyed so much **control** over the rate that they age!

Advanced Nutritional Formulas at the Year's Lowest Prices

For more than two decades, **Life Extension** members have taken advantage of the annual **Super Sale** to acquire their favorite nutrient formulations.

Life Extension **upgrades** its formulas almost every year to provide more *effective* health-sustaining nutrients. In the last few months, we introduced a new form of **ubiquinol CoQ10** that is significantly better absorbed into



the cell's mitochondria, standardized **apigenin** to maintain healthy cell division, and a novel plant compound that protects against persistent **inflammatory joint** conditions.

In this month's issue, you'll learn about nutrients that help arouse youthful **gene expression** changes, along with a new **carbohydrate blocker** (sucrase) that diminishes the amount of unwanted glucose absorbed into your bloodstream.

Every time you purchase a **Life Extension®** product, you contribute to research aimed at radically extending the healthy human lifespan. Last year, the **Life Extension Foundation** funded a record number of scientific projects.

During the traditional winter Super Sale, all **Life Extension** formulas are discounted so that members can obtain up-to-date versions at the **lowest prices** of the year.

Until February 1, 2010, members take advantage of **Super Sale discounts** to stock up on the world's most *avant-garde* life-extending formulations.

For longer life,



William Faloon

Lethal Business and Family Rituals

Many of you were alive when the Surgeon General proclaimed the lethal effects of cigarette smoking in **1964**. I recall smokers back then saying that they "enjoyed" their cigarettes too much to ever quit. Most of these individuals did eventually quit, after perishing from agonizing smoking-induced illnesses.

Americans today ingest oversized breakfasts, lunches, dinners, and snacks as substitutes for tobacco and other addictive substances, sometimes as business or family rituals. To cure drug addiction, one important step is to remove other drug addicts from one's environment. With food addiction, this is often *impossible* as business associates and family members provide constant **temptations** to help destroy your health and longevity by overeating.

I tell those who want to discuss a business matter with me over a meal that I don't hold discussions over a meal or in the presence of food, unless it happens to be at a regularly scheduled mealtime for me. I do not hesitate to accuse well wishing individuals that they are seeking to shorten my life span by inviting me to consume food I would not normally eat.

I realize that many of you may find it difficult to practice moderate calorie restriction. That is why we provide you the option of blocking the absorption of surplus calories by inhibiting digestive enzymes AND reducing the lethal gene expression changes caused by overeating, using nutrients that have been shown to *mimic* some of the gene expression effects of calorie restriction.

References

1. McCay CM, Crowell MF, Maynard LA. The effect of retarded growth upon the length of life span and upon the ultimate body size. 1935. *Nutrition*. 1989 May-Jun;5(3):155-71.
2. Weindruch R, Walford RL. Dietary restriction in mice beginning at 1 year of age: effect on life span and spontaneous cancer incidence. *Science*. 1982 Mar 12;215(4538):1415-8.
3. Weindruch R, Walford RL, Fligiel S, Guthrie D. The retardation of aging in mice by dietary restriction: longevity, cancer, immunity and lifetime energy intake. *J Nutr*. 1986 Apr;116(4):641-54.
4. Raffoul JJ; Guo Z; Soofi A; Heydari AR. Caloric restriction and genomic stability. *J Nutr Health Aging*. 1999;3(2):102-10.
5. Roth GS, Ingram DK, Black A, Lane MA. Effects of reduced energy intake on the biology of aging: the primate model. *Eur J Clin Nutr*. 2000 Jun;54 Suppl 3:S15-20.
6. Lane MA, Ingram DK, Roth GS. Nutritional modulation of aging in nonhuman primates. *J Nutr Health Aging*. 1999; 3(2):69-76.
7. Weed JL, Lane MA, Roth GS, Speer DL, Ingram DK. Activity measures in rhesus monkeys on long-term calorie restriction. *Physiol Behav*. 1997 Jul;62(1):97-103.
8. Hsieh EA, Chai CM, Hellerstein MK. Effects of caloric restriction on cell proliferation in several tissues in mice: role of intermittent feeding. *Am J Physiol Endocrinol Metab*. 2005 May;288(5):E965-72.
9. Weindruch R. Effect of caloric restriction on age-associated cancers. *Exp Gerontol*. 1992 Sep-Dec;27(5-6):575-81.
10. Weindruch R. Dietary restriction, tumors, and aging in rodents. *J Gerontol*. 1989 Nov;44(6):67-71.
11. Martins C, Morgan LM, Robertson MD. Effects of restrained eating behaviour on insulin sensitivity in normal-weight individuals. *Physiol Behav*. 2009 Mar 23;96(4-5):703-8.
12. Lefevre M, Redman LM, Heilbronn LK, et al. Caloric restriction alone and with exercise improves CVD risk in healthy non-obese individuals. *Atherosclerosis*. 2009 Mar;203(1):206-13.
13. Colman RJ, Anderson RM, Johnson SC, et al. Caloric restriction delays disease onset and mortality in rhesus monkeys. *Science*. 2009 Jul 10;325(5937):201-4.
14. Norcross JC, Mrykalo MS, Blagys MD. Auld Lang Syne: Success predictors, change processes, and self-reported outcomes of New Year's resolvers and nonresolvers. *J Clin Psych*. 2002;58(4).
15. Rush EC, Chandu V, Plank LD. Reduction of abdominal fat and chronic disease factors by lifestyle change in migrant Asian Indians older than 50 years. *Asia Pac J Clin Nutr*. 2007;16(4):671-6.

AS WE SEE IT

16. Calabro P, Yeh ET. Intra-abdominal adiposity, inflammation, and cardiovascular risk: new insight into global cardio-metabolic risk. *Curr Hypertens Rep.* 2008 Feb;10(1):32-8.
17. Lakka HM, Laaksonen DE, Lakka TA, et al. The metabolic syndrome and total and cardiovascular disease mortality in middle-aged men. *JAMA.* 2002 Dec 4;288(21):2709-16.
18. Kopp W. The atherogenic potential of dietary carbohydrate. *Prev Med.* 2006 May;42(5):336-42.
19. Zhang X, Zhang G, Zhang H, Karin M, Bai H, Cai D. Hypothalamic IKK β /NF- κ B and ER stress link overnutrition to energy imbalance and obesity. *Cell.* 2008 Oct 3;135(1):61-73.
20. Lacqueman C, Vasseur F, Lepretre F, Froguel P. Adipocytokins, obesity and development of type 2 diabetes. *Med Sci (Paris).* 2005 Dec;21 Spec No10-8.
21. Danielsson A, Fagerholm S, Ost A, et al. Short-term overeating induces insulin resistance in fat cells in lean human subjects. *Mol Med.* 2009 Jul-Aug;15(7-8):228-34.
22. Galili O, Versari D, Sattler KJ, et al. Early experimental obesity is associated with coronary endothelial dysfunction and oxidative stress. *Am J Physiol Heart Circ Physiol.* 2007 Feb;292(2):H904-11.
23. Home P. Contributions of basal and post-prandial hyperglycaemia to micro- and macrovascular complications in people with type 2 diabetes. *Curr Med Res Opin.* 2005 Jul;21(7):989-98.
24. Ardigo D, Valtuena S, Zavaroni I, Baroni MC, Delsignore R. Pulmonary complications in diabetes mellitus: the role of glycemic control. *Curr Drug Targets Inflamm Allergy.* 2004 Dec;3(4):455-8.
25. Eckel RH, Grundy SM, Zimmet PZ. The metabolic syndrome. *Lancet.* 2005 Apr 16-22;365(9468):1415-28.
26. Grattagliano I, Palmieri VO, Portincasa P, Moschetta A, Palasciano G. Oxidative stress-induced risk factors associated with the metabolic syndrome: a unifying hypothesis. *J Nutr Biochem.* 2008 Aug;19(8):491-504.
27. Reeves GK, Pirie K, Beral V, et al. Cancer incidence and mortality in relation to body mass index in the Million Women Study: cohort study. *BMJ.* 2007 Dec 1;335(7630):1134.
28. Henness S, Perry CM. Orlistat: a review of its use in the management of obesity. *Drugs.* 2006 Jun;66(12):1625-56.
29. Juhol C, Armand M, Pafumi Y, Rosier C, Vandermander J, Lairol D. Green tea extract (AR25) inhibits lipolysis of triglycerides in gastric and duodenal medium in vitro. *J Nutr Biochem.* 2000 Jan;11(1):45-51.
30. Kobayashi M, Ichitani M, Suzuki Y, et al. Black-tea polyphenols suppress postprandial hypertriacylglycerolemia by suppressing lymphatic transport of dietary fat in rats. *J Agric Food Chem.* 2009;57(15):7131-6.
31. Pinkston MM, Poston WS, Reeves RS, Haddock CK, Taylor JE, Foreyt JP. Does metabolic syndrome mitigate weight loss in overweight Mexican American women treated for 1-year with orlistat and lifestyle modification? *Eat Weight Disord.* 2006 Mar;11(1):e35-41.
32. Hosoda K, Wang MF, Liao ML, et al. Antihyperglycemic effect of oolong tea in type 2 diabetes. *Diabetes Care.* 2003 Dec;26:1714-8.
33. Fukino Y, Ikeda A, Maruyama K, Aoki N, Okubo T, Iso H. Randomized controlled trial for an effect of green tea-extract powder supplementation on glucose abnormalities. *Eur J Clin Nutr.* 2008 Aug;62(8):953-60.
34. Lin CL, Huang HC, Lin JK. Theaflavins attenuate hepatic lipid accumulation through activating AMPK in human HepG2 cells. *J Lipid Res.* 2007 Nov;48(11):2334-43.
35. Filippatos TD, Gazi IF, Liberopoulos EN, et al. The effect of orlistat and fenofibrate, alone or in combination, on small dense LDL and lipoprotein-associated phospholipase A2 in obese patients with metabolic syndrome. *Atherosclerosis.* 2007 Aug;193(2):428-37.
36. Hodgson JM, Puddey IB, Burke V, Beilin LJ, Jordan N. Effects on blood pressure of drinking green and black tea. *J Hypertens.* 1999 Apr;17(4):457-63.
37. Riemersma RA, Rice-Evans CA, Tyrrell RM, Clifford MN, Lean ME. Tea flavonoids and cardiovascular health. *QJM.* 2001 May;94(5):277-82.
38. Nagao T, Hase T, Tokimitsu I. A green tea extract high in catechins reduces body fat and cardiovascular risks in humans. *Obesity (Silver Spring).* 2007 Jun;15(6):1473-83.
39. Yung LM, Leung FP, Wong WT, et al. Tea polyphenols benefit vascular function. *Inflammopharmacology.* 2008 Oct;16(5):230-4.
40. Rössner S, Sjöström L, Noack R, Meinders AE, Noseda G. Weight loss, weight maintenance, and improved cardiovascular risk factors after 2 years treatment with orlistat for obesity. *Obes Res.* 2000 Jan;8(1):49-61.
41. Shi YF, Pan CY, Hill J, Gao Y. Orlistat in the treatment of overweight or obese Chinese patients with newly diagnosed Type 2 diabetes. *Diabet Med.* 2005 Dec;22(12):1737-43.
42. Oyama T, Saiki A, Endoh K, et al. Effect of acarbose, an alpha-glucosidase inhibitor, on serum lipoprotein lipase mass levels and common carotid artery intima-media thickness in type 2 diabetes mellitus treated by sulfonylurea. *J Atheroscler Thromb.* 2008 Jun;15(3):154-9.
43. Hanefeld M, Chiasson JL, Koehler C, Henkel E, Schaper F, Temelkova-Kurktschiev T. Acarbose slows progression of intima-media thickness of the carotid arteries in subjects with impaired glucose tolerance. *Stroke.* 2004 May;35(5):1073-8.
44. Uddani J, Singh BB. Blocking carbohydrate absorption and weight loss: a clinical trial using a proprietary fractionated white bean extract. *Altern Ther Health Med.* 2007 Jul-Aug;13(4):32-7.
45. Celleno L, Tolaini MV, D'Amore A, Perricone NV, Preuss HG. A Dietary supplement containing standardized Phaseolus vulgaris extract influences body composition of overweight men and women. *Int J Med Sci.* 2007 Jan 24;4(1):45-52.



AS WE SEE IT



46. Zhang XQ, Yang MY, Ma Y, Tian J, Song JR. Isolation and activity of an alpha-amylase inhibitor from white kidney beans. *Yao Xue Xue Bao*. 2007 Dec;42(12):1282-7.

47. Udani J, Hardy M, Madsen DC. Blocking carbohydrate absorption and weight loss: a clinical trial using Phase 2 brand proprietary fractionated white bean extract. *Altern Med Rev*. 2004 Mar;9(1):63-9.

48. Lamela M, Anca J, Villar R, Otero J, Calleja JM. Hypoglycemic activity of several seaweed extracts. *J Ethnopharmacol*. 1989 Nov;27(1-2):35-43.

49. Iwai K. Antidiabetic and antioxidant effects of polyphenols in brown alga Ecklonia stolonifera in genetically diabetic KK-A(y) mice. *Plant Foods Hum Nutr*. 2008 Dec;63(4):163-9.

50. Zhang J, Tiller C, Shen J, et al. Antidiabetic properties of polysaccharide- and polyphenolic-enriched fractions from the brown seaweed Ascophyllum nodosum. *Can J Physiol Pharmacol*. 2007 Nov;85(11):1116-23.

51. Available at: <http://www.naturalproductsinsider.com/news/2009/12/insea2-reduces-glycemic-response.aspx#>. Accessed December 2, 2009.

52. Omoruyi F, Adamson I. Digestive and hepatic enzymes in streptozotocin-induced diabetic rats fed supplements of dikanut (Irvingia gabonensis) and cellulose. *Ann Nutr Metab*. 1993; 37(1):14-23.

53. Oben JE, Ngondi JL, Momo CN, Agbor GA, Sobgui CS. The use of a Cissus quadrangularis/Iringia gabonensis combination in the management of weight loss: a double-blind placebo-controlled study. *Lipids Health Dis*. 2008 Mar 31;7:12.

54. Ngondi JL, Etoundi BC, Nyangono CB, Mbofung CM, Oben JE. IGOB131, a novel seed extract of the West African plant Irvingia gabonensis, significantly reduces body weight and improves metabolic parameters in overweight humans in a randomized double-blind placebo controlled investigation. *Lipids Health Dis*. 2009 Mar 2;8:7.

55. Oben JE, Ngondi JL, Blum K. Inhibition of Irvingia gabonensis seed extract (OB131) on adipogenesis as mediated via down regulation of the PPARgamma and leptin genes and up-regulation of the adiponectin gene. *Lipids Health Dis*. 2008 Nov 13;7:44.

56. Osaki S, Kimura T, Sugimoto T, Hizukuri S, Iritani N. L-Arabinose feeding prevents increases due to dietary sucrose in lipogenic enzymes and triacylglycerol levels in rats. *J Nutr*. 2001;131:796-9.

57. Zeymer U. Cardiovascular benefits of acarbose in impaired glucose tolerance and type 2 diabetes. *Int J Cardiol*. 2006 Feb 8;107(1):11-20.

58. Aronson JK. *Meyler's Side Effects of Endocrine and Metabolic Drugs*. New York, NY: Elsevier Science; 2009.

59. Hanefeld M, Cagatay M, Petrowitsch T, Neuser D, Petzinna D, Rupp M. Acarbose reduces the risk for myocardial infarction in type 2 diabetic patients: meta-analysis of seven long-term studies. *Eur Heart J*. 2004 Jan;25(1):10-6.

60. Ungvari Z, Parrado-Fernandez C, Csiszar A, de Cabo R. Mechanisms underlying caloric restriction and lifespan regulation: implications for vascular aging. *Circ Res*. 2008 Mar 14;102(5):519-28.

61. Fontana L, Villareal DT, Weiss EP, et al. Calorie restriction or exercise: effects on coronary heart disease risk factors. A randomized, controlled trial. *Am J Physiol Endocrinol Metab*. 2007 Jul;293(1):E197-202.

62. Fontana L, Meyer TE, Klein S, Holloszy JO. Long-term calorie restriction is highly effective in reducing the risk for atherosclerosis in humans. *Proc Natl Acad Sci U S A*. 2004 Apr 27;101(17):6659-63.

63. McCarty MF. Glucomannan minimizes the postprandial insulin surge: a potential adjuvant for hepatothermic therapy. *Med Hypotheses*. 2002 Jun;58(6):487-90.

64. Liu S, Willett WC, Manson JE, et al. Relation between changes in intakes of dietary fiber and grain products and changes in weight and development of obesity among middle-aged women. *Am J Clin Nutr*. 2003 Nov;78(5):920-7.

65. Reyna-Villasmil N, Bermudez-Pirela V, Mengual-Moreno E, et al. Oat-derived beta-glucan significantly improves HDLC and diminishes LDLC and non-HDL cholesterol in overweight individuals with mild hypercholesterolemia. *Am J Ther*. 2007 Mar;14(2):203-12.

66. Poppitt SD, van Drunen JD, McGill AT, Mulvey TB, Leahy FE. Supplementation of a high-carbohydrate breakfast with barley beta-glucan improves postprandial glycaemic response for meals but not beverages. *Asia Pac J Clin Nutr*. 2007;16(1):16-24.

67. Bhansali A, Dutta P. Pathophysiology of prediabetes. *J Indian Med Assoc*. 2005 Nov;103(11):594-5.

68. Petersen KF, Shulman GI. Etiology of insulin resistance. *Am J Med*. 2006 May;119(5 Suppl 1):S10-6.

69. Emral R, Koseoglulari O, Tonyukuk V, et al. The effect of short-term glycemic regulation with gliclazide and metformin on postprandial lipemia. *Exp Clin Endocrinol Diabetes*. 2005 Feb;113(2):80-4.

70. Deutsch JC, Santhosh-Kumar CR, Kolhouse JF. Efficacy of metformin in non-insulin-dependent diabetes mellitus. *N Engl J Med*. 1996 Jan 25;334(4):269-70.

71. Paolisso G, Amato L, Eccellente R, et al. Effect of metformin on food intake in obese subjects. *Eur J Clin Invest*. 1998 Jun;28(6):441-6.

72. Ordovas J. Diet/genetic interactions and their effects on inflammatory markers. *Nutr Rev*. 2007 Dec;65(12 Pt 2):S203-7.

73. Raffoul JJ; Guo Z; Soofi A; Heydari AR. Caloric restriction and genomic stability. *J Nutr Health Aging*. 1999 3(2):102-10.

74. Ardigo D, Valtuena S, Zavaroni I, Baroni MC, Delsignore R. Pulmonary complications in diabetes mellitus: the role of glycemic control. *Curr Drug Targets Inflamm Allergy*. 2004 Dec;3(4):455-8.

75. Chang SC, Ziegler RG, Dunn B, et al. Association of energy intake and energy balance with postmenopausal breast cancer in the prostate, lung, colorectal, and ovarian cancer screening trial. *Cancer Epidemiol Biomarkers Prev*. 2006 Feb;15(2):334-41.

76. Fujita A, Hashimoto Y, Nakahara K, Tanaka T, Okuda T, Koda M. Effects of a low calorie vegan diet on disease activity and general conditions in patients with rheumatoid arthritis. *Rinsho Byori*. 1999 Jun;47(6):554-60.

77. Andersson SO, Wolk A, Bergström R, et al. Energy, nutrient intake and prostate cancer risk: a population-based case-control study in Sweden. *Int J Cancer*. 1996 Dec 11;68(6):716-22.

78. Pan SY, DesMeules M, Morrison H, Wen SW, et al. Obesity, high energy intake, lack of physical activity, and the risk of kidney cancer. *Cancer Epidemiol Biomarkers Prev*. 2006 Dec;15(12):2453-60.

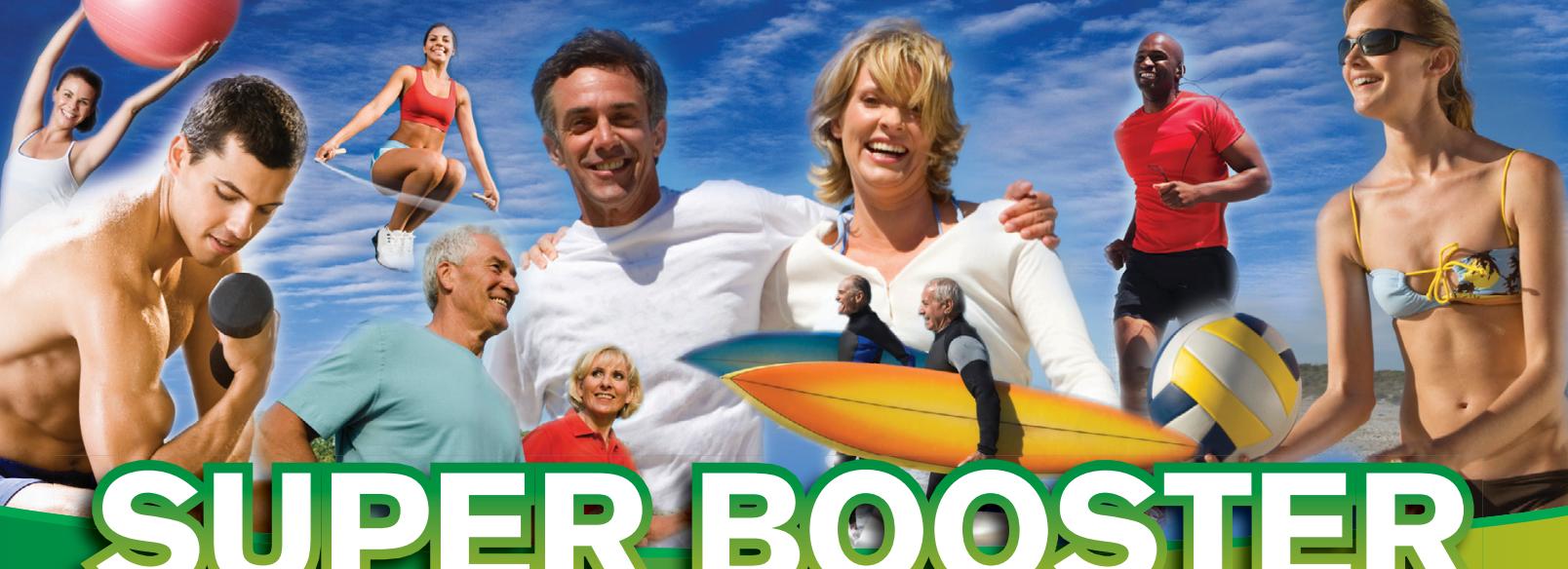
79. Dahl A, Hassing LB, Fransson E, et al. Being Overweight in Midlife Is Associated With Lower Cognitive Ability and Steeper Cognitive Decline in Late Life. *J Gerontol A Biol Sci Med Sci*. 2009 Apr 6.

80. Heilbronn LK, de Jonge L, Frisard MI, et al. Effect of 6-month caloric restriction on biomarkers of longevity, metabolic adaptation, and oxidative stress in overweight individuals: a randomized controlled trial. *JAMA*. 2006 Apr 5;295(13):1539-48.

AS WE SEE IT



81. Lakka HM, Laaksonen DE, Lakka TA, et al. The metabolic syndrome and total and cardiovascular disease mortality in middle-aged men. *JAMA*. 2002 Dec 4;288(21):2709-16.
82. Dekker JM, Girman C, Rhodes T, et al. Metabolic syndrome and 10-year cardiovascular disease risk in the Hoorn Study. *Circulation*. 2005 Aug 2;112(5):666-73.
83. Calle EE, Kaaks R. Overweight, obesity and cancer: epidemiological evidence and proposed mechanisms. *Nat Rev Cancer*. 2004 Aug;4(8):579-91.
84. Messier C, Teutenberg K. The role of insulin, insulin growth factor, and insulin-degrading enzyme in brain aging and Alzheimer's disease. *Neural Plast*. 2005;12(4):311-28.
85. Pasman WJ, Heimerikx J, Rubingh CM, et al. The effect of Korean pine nut oil on in vitro CCK release, on appetite sensations and on gut hormones in post-menopausal overweight women. *Lipids Health Dis*. 2008 Mar 20;7:10.
86. Little TJ, Horowitz M, Feinle-Bisset C. Role of cholecystokinin in appetite control and body weight regulation. *Obes Rev*. 2005 Nov;6(4):297-306.
87. Flegal KM, Graubard BI, Williamson DF, Gail MH. Excess deaths associated with underweight, overweight, and obesity. *JAMA*. 2005 Apr 20;293(15):1861-7.
88. Fontaine KR, Redden DT, Wang C, Westfall AO, Allison DB. Years of life lost due to obesity. *JAMA*. 2003 Jan 8;289(2):187-93.
89. Bouwens M, van de Rest O, Dellschaft N, et al. Fish-oil supplementation induces antiinflammatory gene expression profiles in human blood mononuclear cells. *Am J Clin Nutr*. 2009 Aug;90(2):415-24.
90. Eyles D, Almeras L, Benech P, Patatian A, Mackay-Sim A, McGrath J, Féron F. Developmental vitamin D deficiency alters the expression of genes encoding mitochondrial, cytoskeletal and synaptic proteins in the adult rat brain. *J Steroid Biochem Mol Biol*. 2007 Mar;103(3-5):538-45.
91. Lagouge M, Argmann C, Gerhart-Hines Z, et al. Resveratrol improves mitochondrial function and protects against metabolic disease by activating SIRT1 and PGC-1alpha. *Cell*. 2006 Dec 15;127(6):1109-22.
92. Chen HW, Lee JY, Huang JY, et al. Curcumin inhibits lung cancer cell invasion and metastasis through the tumor suppressor HLJ1. *Cancer Res*. 2008 Sep 15;68(18):7428-38.
93. Vanoorbeek E, Eelen G, Verlinden L, et al. Microarray analysis of MCF-7 breast cancer cells treated with 1,25-dihydroxyvitamin D3 or a 17-methyl-D-ring analog. *Anticancer Res*. 2009 Sep;29(9):3585-90.
94. Aneja R, Odoms K, Denenberg AG, Wong HR. Theaflavin, a black tea extract, is a novel anti-inflammatory compound. *Crit Care Med*. 2004 Oct;32(10):2097-103.
95. Chen YC, Liang YC, Lin-Shiau SY, Ho CT, Lin JK. Inhibition of TPA-induced protein kinase C and transcription activator protein-1 binding activities by theaflavin-3,3'-digallate from black tea in NIH3T3 cells. *J Agric Food Chem*. 1999 Apr;47(4):1416-21.
96. Lin YL, Tsai SH, Lin-Shiau SY, Ho CT, Lin JK. Theaflavin-3,3'-digallate from black tea blocks the nitric oxide synthase by down-regulating the activation of NF-kappaB in macrophages. *Eur J Pharmacol*. 1999 Feb 19;367(2-3):379-88.
97. Lin JK. Cancer chemoprevention by tea polyphenols through modulating signal transduction pathways. *Arch Pharm Res*. 2002 Oct;25(5):561-71.
98. Bode AM, Dong Z. Signal transduction pathways: targets for chemoprevention of skin cancer. *Lancet Oncol*. 2000 Nov;1:181-8.
99. Lyn-Cook BD, Rogers T, Yan Y, et al. Chemopreventive effects of tea extracts and various components on human pancreatic and prostate tumor cells in vitro. *Nutr Cancer*. 1999;35(1):80-6.
100. Beltz LA, Bayer DK, Moss AL, Simet IM. Mechanisms of cancer prevention by green and black tea polyphenols. *Anticancer Agents Med Chem*. 2006 Sep;6(5):389-406.
101. Lee CK, Pugh TD, Klopp RG, Edwards J, Allison DB, Weindruch R, Prolla TA. The impact of alpha-lipoic acid, coenzyme Q10 and caloric restriction on life span and gene expression patterns in mice. *Free Radic Biol Med*. 2004 Apr 15;36(8):1043-57.
102. Pearson KJ, Baur JA, Lewis KN, et al. Resveratrol delays age-related deterioration and mimics transcriptional aspects of dietary restriction without extending life span. *Cell Metab*. 2008 Aug;8(2):157-68.
103. Baur JA, Pearson KJ, Price NL, et al. Resveratrol improves health and survival of mice on a high-calorie diet. *Nature*. 2006 Nov 16;444(7117):337-42.
104. Barger JL, Kayo T, Vann JM, et al. A low dose of dietary resveratrol partially mimics caloric restriction and retards aging parameters in mice. *PLoS One*. 2008 Jun 4;3(6):e2264.
105. Crujeiras AB, Parra D, Milagro FI, et al. Differential expression of oxidative stress and inflammation related genes in peripheral blood mononuclear cells in response to a low-calorie diet: a Nutrigenomics Study. *OMICS*. 2008 Dec;12(4):251-61.
106. Caramia G. Omega-3: from cod-liver oil to nutrigenomics. *Minerva Pediatr*. 2008 Aug;60(4):443-55.
107. Jump DB. N-3 polyunsaturated fatty acid regulation of hepatic gene transcription. *Curr Opin Lipidol*. 2008 Jun;19(3):242-7.
108. Kornman K, Rogus J, Roh-Schmidt H, et al. Interleukin-1 genotype-selective inhibition of inflammatory mediators by a botanical: a nutrigenetics proof of concept. *Nutrition*. 2007 Nov;23(11-12):844-52.
109. Kennedy BK, Steffen KK, Kaeberlein M. Ruminations on dietary restriction and aging. *Cell Mol Life Sci*. 2007 Jun;64(11):1323-8.
110. Guarente L, Picard F. Calorie restriction—the SIR2 connection. *Cell*. 2005 Feb 25;120(4):473-82.
111. Dilova, I., Easlon, E., and Lin, S. Calorie restriction and the nutrient sensing signaling pathways. *Cell Mol Life Sci*. 2007;64:752-67.
112. Chen D, Guarente L. SIR2: a potential target for calorie restriction mimetics. *Trends Mol Med*. 2007;13:64-71.
113. Longo VD. Linking sirtuins, IGF-I signaling, and starvation. *Exp Gerontol*. 2009;44:70-4.
114. Rimando AM, Cuendet M, Desmarchelier C, Mehta RG, Pezzuto JM, Duke SO. Cancer chemopreventive and antioxidant activities of pterostilbene, a naturally occurring analogue of resveratrol. *J Agric Food Chem*. 2002 Jun 5;50(12):3453-7.
115. Joseph JA, Fisher DR, Cheng V, Rimando AM, Shukitt-Hale B. Cellular and behavioral effects of stilbene resveratrol analogues: implications for reducing the deleterious effects of aging. *J Agric Food Chem*. 2008 Nov 26;56(22):10544-51.
116. Cichocki M, Paluszczak J, Szafer H, Piechowiak A, Rimando AM, Baer-Dubowska W. Pterostilbene is equally potent as resveratrol in inhibiting 12-O-tetradecanoylphorbol-13-acetate activated NFκB, AP-1, COX-2, and iNOS in mouse epidermis. *Mol Nutr Food Res*. 2008 Jun;52 Suppl 1:S62-70.



SUPER BOOSTER

with the most effective form of Vitamin K2, Gamma Tocopherol, Ginkgo, and much more!

Despite abundant scientific validation, many people still do not take vital nutrients because they don't want to swallow so many pills. This problem has been solved with a one-per-day softgel that includes multiple health-promoting nutrients in just one supplement. The **Life Extension® Super Booster** contains critical oil-based nutrients that cannot be incorporated into dry-powder based formulas like the **Life Extension Mix™**.

The **Super Booster** has been upgraded to provide higher doses of the most effective form of **vitamin K** known as **menaquinone-7**. Just one **Super Booster** softgel provides:

➔ **Gamma Tocopherol** If one consumes only alpha tocopherol, the critically important gamma tocopherol is displaced from cells within the body. While alpha tocopherol vitamin E inhibits *lipid peroxidation*, the gamma tocopherol form also quenches the dangerous *peroxynitrite* free radical. It is especially important for those who take vitamin E supplements to make sure they consume at least 200 mg a day of gamma tocopherol.

➔ **Sesame Lignans** Sesame lignans augment the antioxidant effects of both alpha- and gamma-tocopherol. In a human study conducted at Life Extension, gamma tocopherol plus sesame lignans was 25% more effective in suppressing measurements of free-radical damage than gamma tocopherol and tocotrienols.

➔ **Vitamin K2** Vitamin K1 from dietary plant sources is poorly absorbed and only a small fraction gets into the bloodstream. Vitamin K2 is absorbed much more efficiently. Scientific studies show K2 provides superior benefits for the bones, arteries, and other tissues. The **MK-4** form of vitamin K2 is the most rapidly absorbed and is now routinely used in Japan to maintain healthy bone density. **MK-4**, however,

Just one softgel of Super Booster supplies:

Gamma tocopherol	230 mg
Ginkgo extract	120 mg
Chlorophyllin	100 mg
Vitamin K2 (as menaquinone-7)	100 mcg
Vitamin K2 (as menaquinone-4)	1000 mcg
Vitamin K1	1000 mcg
Sesame lignans	20 mg
Lycopene	10 mg
Lutein	2 mg
Se-methylselenocysteine	67 mcg
Selenomethionine	67 mcg
Sodium selenite	67 mcg
Vitamin B12	300 mcg
Vitamin C	90 mg
Ascorbyl palmitate	50 mg
Zinc	10 mg
Mixed tocopherols	130 mg

only remains active in the blood for a few hours. The **MK-7** form of K2, on the other hand, remains bioavailable to the human body over a sustained **24-hour period**. Super Booster now provides more **MK-7** than ever before — to keep calcium in the bone and out of the arteries.

➔ **Lycopene** Evidence suggests that people who ingest the carotenoid lycopene enjoy healthier prostate function. Lycopene also helps guard against LDL oxidation.

➔ **Lutein** The carotenoid lutein helps maintain healthy cell division, supports the macula of the eye, and protects the endothelial lining of the arteries.

➔ **Ginkgo** Hundreds of studies substantiate the multifaceted effects of *ginkgo biloba* in promoting healthy circulatory and neurological function.

➔ **Chlorophyllin** Scientific studies indicate that chlorophyllin may protect against environmentally induced damage to DNA.

➔ **Selenium** Some scientific evidence suggests that consumption of selenium may reduce the risk of certain forms of cancer. However, the FDA has determined that this evidence is limited and not conclusive. Selenium's effects in boosting glutathione are well-established.

A bottle of 60 Super Booster softgels retails for \$42. If a member buys four bottles during **Super Sale**, the price is reduced to just \$25.65 per bottle.

The **Super Booster** saves consumers **huge dollars** by combining a wide variety of costly nutrients into one daily softgel. If you add up the price of the individual ingredients contained in the **Super Booster**, you would spend **two to three times more** for this potency if taken separately.



Item #01380

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www.LifeExtension.com

Caution: Those individuals currently taking anticoagulants such as Coumadin® (warfarin) should consult their personal physician before taking supplemental vitamin K.

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3 WAYS TO BENEFIT FROM CALORIE RESTRICTION

1

EAT LESS

Controlled studies in primates reveal that *modest* calorie reduction (only **30%**) yields a **huge** reduction in the risk of issues associated with aging. For example, a recent study in *Science* showed that modest calorie restriction in primates provided a 50% improvement in cardiovascular support.¹ Studies on humans show that a calorie reduction of only **20%** dramatically improves biomarkers associated with longevity.²⁻⁴

For those who find it difficult to ingest **20-30%** fewer calories, consider one or more of the following natural approaches:

1. Pinolenic acid is a polyunsaturated fatty acid derived from pine nuts. It stimulates secretion of the hunger-suppressing hormones **cholecystokinin** (CCK) and **glucagon-like peptide-1** (GLP-1). Human studies showed that participants taking pinolenic acid reduced their food intake by **36%**...more than enough to accomplish *modest* calorie restriction. Pinolenic acid is the active ingredient in a supplement called **Natural Appetite Control**. Those seeking to reduce daily calorie intake should take **three** capsules of **Natural Appetite Control** before most meals.

Natural Appetite Control was developed for adults seeking to lower their calorie intake and maintain a successful, long-term weight management program. Each softgel of **Natural Appetite Control** provides **1,000 mg** of standardized extract of Korean pine nuts containing the highest concentration of pinolenic acid found in any pine nut species, which stimulate the release of two of the body's most powerful hunger-suppressing hormones.

A bottle of 90 softgels of **Natural Appetite Control** retails for \$28. If a member buys four bottles during **Super Sale**, the price is reduced to **\$17.01** per bottle. **Item # 00891**



NOTE: Supplements should be taken in conjunction with a healthy diet and regular exercise program. Results may vary.

2. Ingest **six grams** of **soluble fiber** before each meal. Studies show that pre-meal fiber intake helps induce early satiety. One or more of the following low-cost fiber supplements can be used:

Fiber Food is a natural bulk-producing soluble fiber consisting of psyllium, guar gum, and apple pectin that is available both in capsule and powder forms.

A bottle of 200 capsules of **Fiber Food Capsules** retails for \$15. If a member buys four bottles during **Super Sale**, the price is reduced to **\$8.44** per bottle. **Item# 00229**

A bottle of 300 grams of **Fiber Food Powder** retails for \$15. If a member buys four bottles during **Super Sale**, the price is reduced to **\$8.44** per bottle. **Item#00228**

CocoaGold™ with Beta-Glucan

This combination powder provides **standardized cocoa extract** plus oat **beta-glucans**. The **CocoaGold™** polyphenols support healthy insulin sensitivity, while **beta-glucan** fiber delays **carbohydrate** absorption. **Beta-glucans** have a slower and more sustained effect on blood glucose and energy.

A jar of 180 grams of **CocoaGold™ with Beta-Glucan** retails for \$20. If a member buys four jars during **Super Sale**, the price is reduced to **\$12.15** per jar. **Item# 01285**



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**CALORIE RESTRICTION
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or any of the other supplements that
help protect against chronically bloated
bloodstreams, call **1-800-544-4440**
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REDUCE THE EFFECT OF EXCESS CALORIES

Famine prevailed throughout most of mankind's evolutionary history. Those who efficiently absorbed and stored scarce calories are our direct ancestors. With abundant quantities of calorie-dense foods, it is no wonder that problems associated with excess food intake are so prevalent in modern-day societies.

Excess consumption of dietary fat and carbohydrate results in rapid spikes in levels of fat (triglyceride) and sugar (glucose) in the blood. Regrettably, many people are not aware of the age-accelerating consequences of **postprandial overload**. Elevated levels of post-meal blood fat (triglycerides) and blood sugar (glucose) *accelerate* the aging process in our cardiovascular system.

Ingested foods are broken down by *digestive enzymes* before they are *absorbed* into the bloodstream.

Green and black tea polyphenols target the fat-digesting enzyme *lipase*. **Theaflavins** from **tea polyphenols** have been shown in experimental studies to help reduce the post-meal impact of ingested fat calories in a dose-dependent manner.⁶

Most of us eat too many refined *carbohydrates* that add to the calorie burden. The enzymes *amylase*, *glucosidase*, and *sucrase* break down ingested carbohydrates to smaller starch and sugar particles. Nutrients such as **white kidney bean** (targets amylase activity),⁷⁻⁸ **InSea™** (targets amylase and glucosidase activity),⁹ **Irvingia** (targets amylase activity)¹⁰⁻¹¹ and **L-arabinose** (targets sucrase activity)¹² have been shown in published scientific studies to help reduce the metabolic impact of ingested carbohydrate calories from dietary starch and/or sugar. These *nutrients* are all contained in the new **Optimized Irvingia** formula.

Optimized Irvingia with Phase 3™ Calorie Control Complex contains *Green Tea Phytosome Extract* to slow down the breakdown and absorption of dietary fat, **Phase 2® White Kidney Bean** extract to moderate *alpha-amylase* activity, **InSea™** seaweed extract, which can help support a *slowing* of the rate of carbohydrate absorption from the intestines, **Irvingia gabonensis** extract for *leptin sensitivity* support, and **Phase 3™ Sucrase Modulator** to reduce the number of absorbed calories from sugar. Directions are to take two capsules before the two heaviest meals of the day. A bottle of **120** capsules of **Optimized Irvingia with Phase 3™ Calorie Control Complex** retails for \$78. If a member buys four bottles during **Super Sale**, the price is reduced to **\$48.60 per bottle**. Item# **01492**



REFERENCES

- Science. 2009 Jul 10;355(5937):201-4.
- Circ Res. 2008 Mar 14;102 (5): 519-28.
- Am J Physiol Endocrinol Metab. 2007 Jul;293(1):E197-202.
- Proc Natl Acad Sci USA. 2004 Apr 27;101(17):6659-63.
- Chemical Society National Meeting & Exposition; March 26-30, 2006; Atlanta, GA.
- Agric. Food Chem. 2009, 57 (15), pp 7131–6.
- Yao Xue Xue Bao. 2007 Dec;42(12):1282-7.
- Altern Ther Health Med. 2007 Jul-Aug;13(4):32-7.
- J Ethnopharmacol. 1989 Nov;27(1-2):35-43.
- Lipids Health Dis. 2009 Mar 2;8:7.
- Lipids Health Dis. 2008 Nov 13;7:44.
- J Nutr. 2001 Mar 131:796-9.
- Cell. 2006 Dec 15;127(6):1109-22.
- Endocrinology. 2008 Jan;149(1):84-92.
- Crit Care Med. 2004 Oct;32(10):2097-103.
- J Agric Food Chem. 1999 Apr;47(4):1416-21.
- Arch Pharm Res. 2002 Oct;25(5):561-71.
- Nutr Cancer. 1999;35(1):80-6.
- Anticancer Agents Med Chem. 2006 Sep; 6(5):389-406.
- Nature. 2006 Nov 16;444(7117):337-42.
- PLOS One. 2008 Jun 4;3(6):e2264.
- OMICS. 2008 Aug 7.

CONTROL YOUR GENE EXPRESSION

Restricting the amount of calories allowed into your blood stream *increases* healthy life span via several mechanisms. Cardiovascular support can be maintained by blunting the post-meal surge of glucose, insulin, triglycerides, and inflammatory-inducing compounds that engorge the bloodstream following eating binges.

Scientists, however, are focused on an even more important mechanism to explain how *calorie restriction* so radically extends life span. They have uncovered favorable alterations in **gene expression** in response to reduced calorie intake. It turns out that caloric restriction slows aging by **activating** beneficial "youth" genes while disabling detrimental "senescence" genes.

The incredible news is that a select group of nutrients has been discovered that trigger many of the favorable mechanisms (including more youthful **gene expression** patterns) as **caloric restriction**.

Among the most promising of these **caloric restriction mimics** and enhancers are **resveratrol**, **pterostilbene**, **quercetin**, and **grape seed polyphenols**, along with **black tea extract**. These nutrients have been shown to generate many of the same effects in the body as caloric restriction, without significant dietary modification.¹³⁻²⁰ In particular, they help "mimic" caloric restriction's favorable impact on **genes** that influence the aging process.¹³⁻²⁰

Genes have the capacity to directly affect life span by regulating a broad spectrum of aging factors. Calorie restriction exerts a beneficial effect on the *activity of gene expression*, supporting healthy cellular function through numerous physiological pathways.²¹ Compounds that mimic caloric restriction bring about favorable changes in *gene expression* and improve the primary *biomarkers* of aging.²²

Those taking high-dose vitamin D, along with coenzyme Q10 and fish oil are already favorably altering many of their gene expression patterns. A new **Calorie Restriction Mimetic Formula** includes **resveratrol**, **pterostilbene**, **quercetin**, **grape seed polyphenols**, and **black tea extract** to provide even broader-spectrum gene expression support in one nutritional compound. Two capsules of the new **Calorie Restriction Mimetic Formula** provides:



Item #01419

Trans-Resveratrol	250 mg
Trans-Pterostilbene	3 mg
Quercetin	150 mg
Black tea extract	300 mg
Grape seed polyphenols	50 mg

A bottle containing 60 vegetarian capsules of the new **Calorie Restriction Mimetic Formula** retails for \$36. If a member buys four bottles during the **Super Sale**, the cost per bottle is reduced to **\$22.28**. Item# **01419**

While the new **Calorie Restriction Mimetic Formula** can enable aging individuals to *simulate* many of the beneficial **gene expression** effects associated with caloric restriction, **Life Extension** members are still urged to reduce their pathologic calorie burden by following steps #1 and #2 as outlined herein.



Heart Healthy SUPER ome

Fish Oil Blend with Sesame Lignans and a **MORE** Concentrated Olive Fruit Extract

A Comprehensive Essential Fatty Acid Formula

An abundance of scientific research substantiates the wide-ranging health benefits, including promoting a healthy heart, of **omega-3 fatty acids** in fish oil and **monounsaturated fatty acids-polyphenols** in olive fruit.¹⁻⁹

SUPER OMEGA-3 from Life Extension® uses a patented EPA/DHA extraction process that results in a pure, stable and easy-to-tolerate **fish oil extract**.

While most fish oil is distilled to decrease contaminants such as mercury and PCBs, the **Pure +™** fish oil used in **Super Omega-3** is produced with a patented method that purifies the oil with a highly advanced *distillation process* to reduce pollutants to virtually undetectable levels. The result is an improved **fish oil** that exceeds the standards set by international rating agencies.

Sesame Lignans Enhance Fish Oil's *In Vivo* Effects

The *unstable* nature of fatty acids like fish oil limits their biological efficacy in the body. Scientific studies show that when **sesame lignans** are supplemented with fish oil, the beneficial effects are augmented.¹⁰

Sesame lignans help guard against **lipid peroxidation**, thereby extending the stability of **DHA** in the body. These **lignans** also direct fatty acids toward pathways which can help with inflammatory reactions. **Super Omega-3** provides standardized **sesame lignans** to enhance the overall benefits of the improved EPA/DHA fish oil blend.

New Higher Potency Standardized Olive Fruit

To emulate a Mediterranean diet, **Super Omega-3** provides a standardized **olive fruit extract** to deliver the polyphenol **hydroxytyrosol**, a powerful antioxidant that protects normal **LDL** from oxidation and counters dangerous free radicals.¹¹⁻¹⁴ Research shows that a combination of olive oil and fish oil supplements helps with inflammation better than a placebo or fish oil alone.¹⁵ Therefore, Super Omega-3 provides the equivalent polyphenol content of **6 ounces of Extra Virgin Olive Oil**.

Super Omega-3 provides a new, higher concentration **olive fruit extract** standardized for **hydroxytyrosol**, **tyrosol** and **oleuropein** polyphenols. Research shows the value of **oleuropein** in favorably altering gene expression, delaying senescence in specialized skin cells, and helping maintain normal platelet activation.

Life Extension's **Super Omega-3** is a fish oil concentrate that contains a *full-spectrum blend* of synergistic nutrients, providing a product of the utmost quality to produce optimal effects.

Super Omega-3 is certified to contain no detectable levels of mercury, arsenic, lead, cadmium, and other toxic metals by the International Fish Oil Standards (IFOS™). This product meets or exceeds IFOS™ standards for PCBs, dioxins, and other contaminants, and thus has received its highest 5-star rating.

IFOS™ certification mark is a registered trademark of Nutrasource Diagnostics, Inc. These products have been tested to the quality and purity standards of the IFOS™ program conducted at Nutrasource Diagnostics, Inc.

References:

1. *Public Health Nutr.* 2006 Dec;9(8A):1136-40.
2. *Am J Prev Med.* 2005 Nov;29(4):335-46.
3. *J Am Diet Assoc.* 2005 Mar;105(3):428-40.
4. *Mini Rev Med Chem.* 2004 Oct;4(8):859-71.
5. *Nurs Stand.* 2004 Aug 11-17;18(48):38-42.
6. *Cleve Clin J Med.* 2004 Mar;71(3):208-10, 212, 215-8 *passim*.
7. *J Nutr Health Aging.* 2001;5(3):144-9.
8. *Inflamm Res.* 2001 Feb;50(2):102-6.
9. *Arch Intern Med.* 2000 Mar 27;160(6):837-41.
10. *Biochem Biophys Acta.* 2004 Jun 1;1682(1-3):80-91.
11. *Anal Chim Acta.* 2007 Feb 5;583(2):402-10.
12. *J Agric Food Chem.* 2007 Sep 5;55(18):7609-14.
13. *Lipids.* 2001 Nov;36(11):1195-202.
14. *Eur J Cancer.* 2000 Jun;36(10):1235-47.
15. *Nutrition.* 2005 Feb;21(2):131-6.

omega-3 EPA/DHA

Two softgels should be taken twice daily with meals. A bottle of 120 softgels of **SUPER OMEGA-3 EPA/DHA with Sesame Lignans and Olive Fruit Extract** retails for \$32. If a member buys four bottles during **Super Sale**, the price is reduced to **\$18.90 per bottle**. If 10 bottles are purchased during **Super Sale**, the price per bottle is reduced to only **\$16.81**. **Item# 01482**

For those with a sensitive stomach, Super Omega-3 is now also available with enteric coating and retails for \$34. If a member buys four bottles during **Super Sale**, the price is reduced to **\$20.93 per bottle**. If 10 bottles are purchased during **Super Sale**, the price per bottle is reduced to only **\$18.90**. **Item# 01484**

To order the most advanced fish oil supplement, **Super Omega-3 EPA/DHA with Sesame Lignans and MORE Olive Fruit Extract ... with or without enteric coating ... call 1-800-544-4440 or visit www.LifeExtension.com**

Supportive but not conclusive evidence shows that **consumption of EPA and DHA omega-3 fatty acids may reduce the risk of coronary heart disease.**

Important note: If you are taking anticoagulant drugs, use this product only under the supervision of your physician. Blood tests that measure clotting can be used to ensure these nutrients are not reducing the clotting factors in your blood to abnormal levels.

Upgraded Formula!

Just four softgels of **SUPER OMEGA-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract** provide:

EPA Pure+™ Extract 1400 mg
(eicosapentaenoic acid)

DHA Pure+™ Extract 1000 mg
(docosahexaenoic acid)

Olive Fruit Extract 600 mg
[standardized to 1.5% hydroxytyrosol (9 mg),
0.7% oleuropein (4.2 mg), 0.5% verbascoside
(3 mg) and 0.23% tyrosol (1.4 mg)]

Sesame seed lignan extract 20 mg



Item #01482

Item #01484



POWERFUL PROTECTION AGAINST LDL Oxidation and Proinflammatory Cytokines

An increasing number of scientists recognize the critical need to protect the arterial wall against low-density lipoprotein (LDL) **oxidation** and **inflammatory** insults.

Research has shown that unique extracts present in black tea called **theaflavins** can have multiple applications for arterial health.¹

New Theaflavin Standardized Extract contains a number of beneficial flavonoids found naturally in tea leaves that help support levels of cholesterol that are already within the normal range.¹

Theaflavins have been shown in human studies to protect against **LDL oxidation** and favorably affect **endothelial function**,² thus helping to maintain healthy circulation.

Scientists have also found that black tea flavonoids possess strong **antioxidant** properties, which can help mitigate oxidative damage to cells and tissues from free radicals.³ In addition, theaflavins have been found to be helpful in regulating key *inflammatory* mediators in the body, thus helping to preserve cellular integrity.^{4,5}

A bottle of 30 350-mg vegetarian capsules of **Theaflavin Standardized Extract** retails for \$18. If a member buys four bottles during **Super Sale**, the cost is only **\$10.80 per bottle**.

References

1. J Nutr. 2003;133:3298S-3302S.
2. J Nutr. 2003;133:3293S-3297S.
3. Prev Med. 2005;40:910-8.
4. Folia Biol (Praha). 2007;53(5):164-72.
5. Crit Care Med. 2004;32:2097-103.

This product contains a black tea extract which is licensed from Applied Food Sciences, Inc. and is protected by US Patent Nos. 6,811,799 and 6,602,527.

Item #01304



To order **Theaflavin Standardized Extract Capsules**, call **1-800-544-4440** or visit www.lifeextension.com



IN THE NEWS

Obesity Responsible for 100,000 Cancer Cases Annually

A recent report from the *American Institute of Cancer Research* (AICR) states that excess body fat is now seen as a major cause of cancer.* Researchers at AICR studied seven cancers known to have links with obesity and calculated the actual case counts that were likely to have been caused by obesity. The numbers are staggering, including 49% of endometrial cancers, 35% of esophageal cancers, 28% of pancreatic cancers, 24% of kidney cancers, 21% of gallbladder cancers, 17% of breast cancers, and 9% of colorectal cancers.

Laurence Kolonel, MD, PhD, Deputy Director of the Cancer Research Center of Hawaii and AICR/WCRF expert panel member, presented the new preventability estimates and reviewed the evidence linking obesity to cancer risk. "We now know that carrying excess body fat plays a central role in many of the most common cancers," he said. "And it's clearer than ever that obesity's impact is felt before, during and after cancer—it increases risk, makes treatment more difficult and shortens survival."

Recent findings also suggest that excess body fat lowers immune function and increases oxidative stress, which can lead to DNA damage.

—Jon Finkel

* Available at: http://www.aicr.org/site/News2?abbr=pr_&page=NewsArticle&id=17333&news_iv_ctrl=1102. Accessed December 1, 2009.



Elevated Homocysteine Associated with Doubling of Women's Alzheimer's Disease Risk

A thesis composed by Dr. Dimitri Zylberstein at the University of Gothenburg, Sweden links high levels of homocysteine in women with twice the risk of developing Alzheimer's disease compared to those with low levels.*

Dr. Zylberstein reports the results of a study involving participants in the Prospective Population Study of Women in Gothenburg. The women were followed for 35 years, making it the longest study to evaluate the relationship between homocysteine level and dementia.

"Alzheimer's disease was more than twice as common among the women with the highest levels of homocysteine than among those with the lowest, and the risk for any kind of dementia was 70% higher," revealed Dr. Zylberstein.

Although elevated homocysteine can be the result of vitamin B12 and folate deficiencies, it can also occur when vitamin status is considered normal according to current standards.

Editor's note: Homocysteine can be reduced by taking folic acid, vitamin B12, and vitamin B6 supplements. Those with stubbornly high homocysteine can sometimes lower it by taking **1,000-3,000 mg of trimethylglycine (TMG)** each day. Those with intractably high homocysteine should take the metabolically active form of folic acid called **5-methyltetrahydrofolate** in the daily dose of **2,000 mcg** to over **10,000 mcg**. Optimal homocysteine blood levels are below **7-8 µmol/L**.

—Dayna Dye

* Dr. Dimitri Zylberstein thesis presented at the University of Gothenburg.

IN THE NEWS

Long-term High Dose Vitamin D Well Tolerated

A letter published in the *Archives of Internal Medicine* revealed that treatment with 50,000 IU of vitamin D per week was safe over an 8-week period, and could also be used every other week as maintenance.*

The researchers reviewed the records of 86 patients treated for vitamin D insufficiency. Forty-one subjects received 50,000 IU vitamin D2 weekly for 8 weeks followed by a maintenance dose of 50,000 IU every other week for up to 6 years. The remainder of the patients received every-other-week maintenance therapy.

For patients who received the starting therapy, 25-hydroxyvitamin D levels rose to 37.2 ng/mL after 8 weeks. Maintenance therapy increased these levels to 46.9 ng/mL. For those who received only maintenance therapy, vitamin D levels increased to 47 ng/mL.

"While treating and preventing vitamin D deficiency, these large doses of vitamin D2 do not lead to vitamin D toxicity," researcher Michael Holick concluded.

Editor's note: It is interesting to see that even when these high doses of vitamin D are used, the average study subject did not attain optimal vitamin D blood levels of greater than **50 ng/mL**. This further validates the importance for most people to take at least **5,000 IU** of vitamin D3 each day for 60-90 days and then have their blood tested for **25-hydroxyvitamin D**. Dosage adjustment can then be made to achieve optimal **25-hydroxyvitamin D** status, which based upon the current scientific literature appears to be **50-80 ng/mL**.

—Dayna Dye

* *Arch Int Med.* 2009 Oct 26;169(19).

Vitamin E Beats Drug Treatment for Liver Disease



At the 60th Annual Meeting of the American Association for the Study of Liver Diseases, Arun Sanyal, MD of Virginia Commonwealth University reported the results of a trial which found that vitamin E was more successful than the antidiabetic drug pioglitazone at treating non-alcoholic steatohepatitis (NASH).* Non-alcoholic steatohepatitis is a progressive liver disease associated with fatty liver, insulin resistance, and obesity.

The study included 247 patients whose liver biopsies confirmed NASH within six months of the trial. Participants were randomized to receive 30 mg per day of pioglitazone, 800 IU per day vitamin E, or a placebo for 96 weeks.

While disease activity scores improved in 19% of the placebo patients, 34% of those who received pioglitazone and 43% of those who received vitamin E had improved scores.

The trial is the first large study to demonstrate a benefit for vitamin E in non-alcoholic steatohepatitis treatment.

Editor's note: Non-alcoholic steatohepatitis (fatty liver disease) can be effectively treated by losing weight and suppressing chronic inflammatory reactions in the body. The best nutrient to consider is **polyenylphosphatidylcholine** in the dose of **1,800 to 2,700 mg** a day.

—Dayna Dye

* 60th Annual Meeting of the American Association for the Study of Liver Diseases.

Green Tea May Help Protect Against Oral Cancer

An article published in a recent issue of *Cancer Prevention Research* describes a protective effect for green tea against the progression of oral lesions that are at high risk of developing into cancer.*

Forty-one patients with oral premalignant lesions were randomized to receive one of three concentrations of a green tea extract or a placebo 3 times daily for 12 weeks. Among participants who received the highest concentrations of tea extract, 58.8% showed partial clinical responses, defined as improvement in degree of maturation of epithelium with no new lesions or progression, while complete or partial responses were observed in 36.4% of those who received the lowest extract dose and 18.2% of those who received the placebo.

The trial is the first to test the effects of green tea as a cancer preventive among those with premalignant oral lesions.

—Dayna Dye

* *Cancer Prev Res.* 2009 Nov; 2(11):919-21.



Heart Disease, Stroke, Heart Failure, and Premature Death All Linked to Insufficient Vitamin D Levels

The results of a study presented at the American Heart Association's Scientific Conference in Orlando confirmed a strong association between the presence of reduced vitamin D levels and a greater risk of coronary artery disease, stroke, heart failure, and dying over follow-up in men and women 50 years of age and older.*

Brent Muhlestein, MD and his colleagues followed 27,686 subjects with no history of heart disease for an average of 1.2 years. Those with very low vitamin D levels were 45% likelier to develop heart disease, twice as likely to develop heart failure, 78% more likely to experience a stroke, and 77% likelier to die than those with normal levels.

"This was a unique study because the association between vitamin D deficiency and cardiovascular disease has not been well-established," Dr. Muhlestein commented.

—Dayna Dye

* American Heart Association's Scientific Conference in Orlando.

Vitamin Treatment Improves ED in Men with Elevated Homocysteine

An article published online in the *Journal of Sexual Medicine* describes research conducted at the University of Rome which found that it may be necessary to reduce homocysteine before treatment for erectile dysfunction (ED) can be effective.*

The study included 75 men with erectile dysfunction who were treated with sildenafil citrate (Viagra®) for 2 months. Nonresponders to the drug were treated with 600 mg vitamin B6 per week and 15 mg folic acid per day along with sildenafil for 6 weeks.

All of the 18 patients who initially failed to respond to drug treatment had high levels of homocysteine and low folic acid levels. Subsequent to the 6 week course of vitamin therapy, all but two participants experienced improvement in ED.

The association of hyperhomocysteinemia with vascular disease supports the mechanism for homocysteine-reducing agents in improving ED, which is caused primarily by the same factors that affect the coronary arteries.

—Dayna Dye

* *J Sex Med.* 2009 Aug 17.

Omega-3 Fatty Acids Boost Glucosamine's Arthritis Benefit

In a recent issue of the journal *Advances in Therapy*, German researchers report that the addition of omega-3 polyunsaturated fatty acids to glucosamine sulfate resulted in improved alleviation of symptoms compared to glucosamine alone.*

The researchers enrolled 177 men and women moderate-to-severe hip or knee osteoarthritis. Participants were randomized to receive glucosamine sulfate with or without the omega-3 fatty acids EPA and DHA for 26 weeks.

When an at least 80% reduction in pain was evaluated, 44% of those in the combination group compared to 32% of those who received only glucosamine were categorized as responders. Combination therapy was received by twice as many patients who reported a 90 to 100% reduction in pain compared to those who received glucosamine alone.

The authors remark that while glucosamine sulfate improves cartilage metabolism, EPA and DHA further reduce degradation by suppressing inflammation, which lowers swelling and pain.

Editor's note: **Life Extension** members have long been advised to use a combination of glucosamine and/or chondroitin sulfate along with omega-3 fatty acids from fish oil to help manage arthritis. Life Extension suggests 1,400 mg EPA and 1,000 mg DHA per day, such as are supplied by two Super Omega-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract, along with 1,500 mg glucosamine and 1,000 mg chondroitin.

—Dayna Dye

* *Advances in Therapy.* 2009 Sep;26(9).

IN THE NEWS



Soy Shows Promise for Colon Cancer Prevention

An article featured in a recent issue of the journal *Cancer Research* reveals the discovery of a team at Children's Hospital & Research Center in Oakland of compounds occurring in soy that could help prevent and possibly treat colon cancer.*

Julie Saba, MD and her associates discovered that natural lipid molecules called sphingadienes may be responsible for some of the cancer-preventive benefits found in soy. They first identified the compounds in fruit flies and found that they had the effect of inducing the death of mutant cells.

"It's very exciting," enthused Dr. Saba. "First, we are encouraged to find a natural molecule that could be consumed through soy products as a strategy to help prevent colon cancer. Second, this information is important because we can build on our understanding of the structure and metabolism of sphingadienes in terms of developing new drugs to treat people who already have colon cancer."

—Dayna Dye

* *Cancer Research*. 2009 Dec. 15.

Antioxidants Could Offer Protection to the Lungs During Flu Season

A report published in a recent issue of the *FASEB Journal* revealed a protective effect for antioxidants against lung damage caused by influenza.

Sadis Matalon and colleagues at the University of Alabama demonstrated that the virus damages the lungs via its M2 protein, which attacks the cells of the lungs' lining by disrupting their ability to remove fluid. In an experiment using human airway cells, those transfected with M2 revealed increased levels of damaging molecules known as reactive oxygen species. Co-administration of M2 with glutathione ester (an antioxidant compound) prevented M2 from causing damage.

"The recent outbreak of H1N1 influenza and the rapid spread of this strain across the world highlight the need to better understand how this virus damages the lungs and to find new treatments," Dr. Matalon remarked. "Additionally, our research shows that antioxidants may prove beneficial in the treatment of flu."

—Dayna Dye

* Available at: <http://www.fasebj.org/cgi/content/abstract/23/11/3829>. Accessed December 1, 2009.



Resveratrol May Prove to be HRT Alternative

A study published in the *Journal of Nutritional Biochemistry* shows that resveratrol may possibly offer a safer hormone replacement therapy (HRT) alternative and chemoprevention of breast cancer because of its estrogenic activity and high antitumor activity.*

Resveratrol is a phytoestrogen, which is a natural plant substance found in foods such as grape skins and red wine that display weak estrogen-like activity toward mammals. The directive of the study was to assess the estrogen-like effects and antitumor effects of individual dietary phytoestrogens by analyzing their effects on tumor cell growth, cell cycle activity and apoptosis (programmed cell death).



"Because it (resveratrol) stimulated the transcription of endogenous estrogen receptor (ER) and proapoptotic effects, this phytoestrogen is the most promising candidate as an HRT alternative and chemopreventive reagent for breast cancer," the researchers concluded.

—Jon Finkel

* *J Nutr Biochem*. 2009 Oct 2.

Achieve Multiple Health Benefits From Highly Absorbable Curcumin

How Much Curcumin Are You Absorbing?

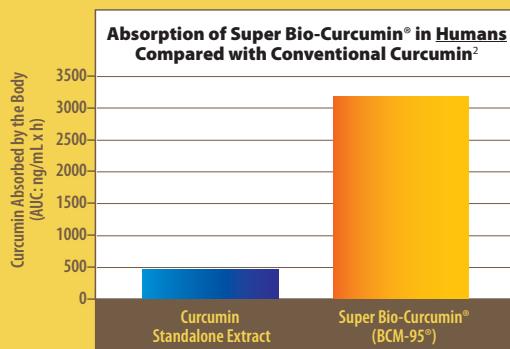


Chart 1. Super Bio-Curcumin® (BCM-95®) showed 6.9 times greater bioavailability (absorption and sustainability over 8 hours) in humans compared with conventional curcumin (as measured by the area under the curve [AUC] in a plot of blood levels against time, that is, the total amount of curcumin absorbed by the body over 8 hours).

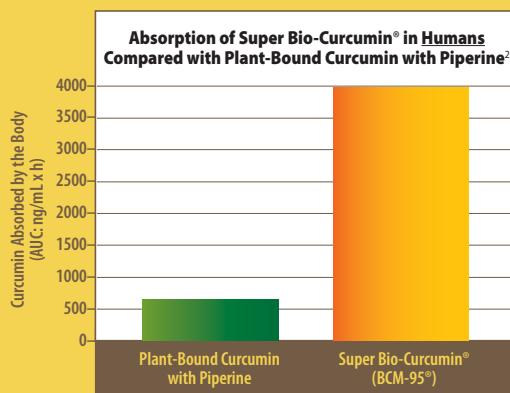


Chart 2. Super Bio-Curcumin® (BCM-95®) showed 6.3 times greater bioavailability (absorption and sustainability over 8 hours) in humans compared with plant-bound curcumin with piperine (as measured by the area under the curve [AUC] in a plot of blood levels against time, that is, the total amount of curcumin absorbed by the body over 8 hours).

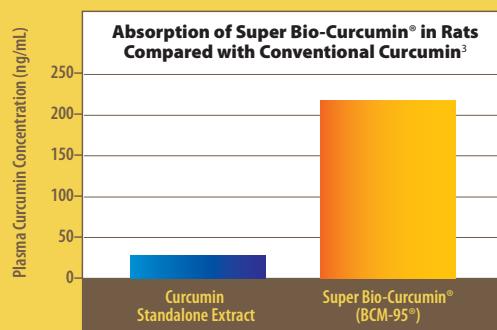


Chart 3. Bioavailability in rats fed with BCM-95® is 7.8 times higher than conventional curcumin.

Curcumin and other *turmeric* constituents are impressing scientists around the world with their remarkable health benefits, such as suppressing inflammatory factors, fighting free radicals, and promoting healthy DNA function. The problem is that curcumin is *poorly* absorbed into the bloodstream.

Super Bio-Curcumin® has been shown to **absorb up to seven times better** than conventional curcumin. This patent-pending formula thus represents the most cost-effective way to supplement with this critical nutrient.^{1,2}

The remarkable absorption studies charted to the left reveal that just one 400 mg capsule a day of this turmeric compound can provide *curcumin* blood levels equivalent to ingesting **2,500–2,800 mg** of commercial curcumin supplements.

Not only does this novel turmeric formulation provide far greater *peak* blood levels, but the curcumin also remains in the bloodstream almost **twice as long** compared with conventional supplements.

What's more, this enhanced absorption delivery complex provides other beneficial turmeric compounds in addition to standardized *curcumin*.

SUPER BIO-CURCUMIN® WITH BCM-95®



ITEM #00407

Life Extension®'s **Super Bio-Curcumin®** contains the patent-pending BCM-95® turmeric compound. A bottle containing **60 Super Bio-Curcumin® with BCM-95®** capsules retails for \$30. If a member buys four bottles during **Super Sale**, the price is reduced to just **\$17.89** per bottle. Each bottle will last most members two months.

References:

1. *Spice India*. 2006 Sept;19(9):11-5.
2. A novel bioenhanced preparation of curcuminoids. Study submitted for publication, 2007.
3. Bioavailability study of BCM-95® in rats. Orcas International Inc. 2006.

To order Super Bio-Curcumin® with BCM-95®, call 1-800-544-4440 or visit www.LifeExtension.com

Bio-Curcumin® and BCM-95® are registered trademarks of Dolcas-Biotech, LLC.

These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

LARGEST SELECTION OF VITAMIN D SUPPLEMENTS

More studies have been published over the past four years supporting the health benefits of **vitamin D** than possibly any other compound.

The good news for consumers is that vitamin D is a very **low cost** supplement.

The objective of taking a vitamin D supplement is to achieve **25-hydroxyvitamin D** blood levels of **50 ng/mL** (and higher).

Life Extension's® exclusive analysis of over 13,000 **vitamin D** blood tests reveals that the minimum intake for most aging people should be around **7,000 IU** a day. Some individuals need **10,000 IU** of vitamin D daily.

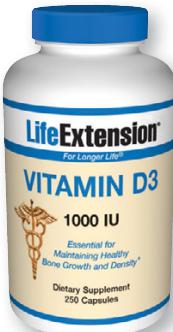
You can choose the right dose of **vitamin D3** for you from the large selection of vitamin D3 supplements below. Remember to factor in that you may be getting **1,000-3,000 IU** of vitamin D in multi-nutrient formulas you already take.

■ Vitamin D3 1,000 IU • 250 capsules

Retail: \$12.50

Four bottle Member/Super Sale Price: \$7.59

Commercial companies offered only **400 IU** vitamin D products when Life Extension long ago introduced this **1,000 IU** version. For most people, this **1000 IU** potency is *insufficient* to attain optimal vitamin D blood levels. For smaller individuals who obtain **2,000-3,000 IU** in their multi-nutrient formulas (and children), this potency of vitamin D may be suitable. **Item# 00251**

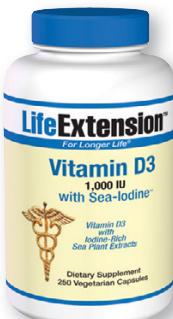


■ Vitamin D3 1,000 IU with 1,000 IU of Sea-Iodine • 250 vegetarian capsules

Retail: \$22

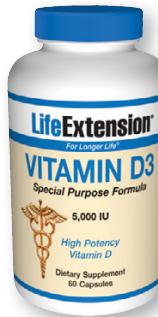
Four bottle Member/Super Sale Price: \$13.50

Most people do not ingest enough vitamin D and iodine, especially those seeking to reduce their salt intake. Combining **vitamin D3** and **iodine** into one capsule makes taking these two nutrients economical and convenient. **Item# 01371**



To order any of these high-potency vitamin D3

supplements at low Super Sale prices, call **1-800-544-4440**
or visit www.LifeExtension.com



■ Vitamin D3 5,000 IU • 60 capsules

Retail: \$11

Four bottle Member/Super Sale Price: \$6.68

For those obtaining **1,000-3,000 IU** of vitamin D in their multi-nutrient formulas, this **5,000 IU** potency is what most need to achieve optimal vitamin D blood levels. **Item# 00713**



■ Vitamin D3 5,000 IU with 1,000 IU of Sea-Iodine • 60 vegetarian capsules

Retail: \$14

Four bottle Member/Super Sale Price: \$8.44

Most people do not ingest enough vitamin D and iodine, especially those seeking to reduce their salt intake. Combining **5,000 IU** of **vitamin D3** and **1,000 IU** of **iodine** into one capsule make taking these two nutrients economical and convenient. **Item# 01372**



■ Vitamin D3 7,000 IU • 60 capsules

Retail: \$14

Four bottle Member/Super Sale Price: \$8.51

Some individuals (such as those weighing more than 180 pounds) may require higher potencies of vitamin D. When combined with **1,000-3,000 IU** obtained from multi-nutrient formulas, this **7,000 IU** **vitamin D3** capsule should enable these individuals to attain **25-hydroxyvitamin D** blood levels above the desired range of **50 ng/mL**. **Item# 01418**



■ Vitamin D3 Liquid Emulsion 2,000 IU • 1 ounce

Retail: \$28

Four bottle Member/Super Sale Price: \$16.88

For those rare individuals who have difficulty absorbing enough vitamin D3 from powdered capsules, this liquid emulsion of vitamin D can be used. **Item# 00864**

CAUTION: Individuals consuming more than 2,000 IU/day of vitamin D (from diet and supplements) should periodically obtain a serum 25-hydroxyvitamin D measurement. Do not exceed 10,000 IU per day unless recommended by your doctor. Vitamin D supplementation is not recommended for individuals with hypercalcemia (high blood calcium levels). People with kidney disease, certain medical conditions (such as hyperparathyroidism or sarcoidosis), and those who use cardiac glycosides (digoxin) or thiazide diuretics should consult a physician before using supplemental vitamin D.

Low levels of testosterone have been implicated in a host of life-threatening health problems. Maintaining normal testosterone levels is one of the most important steps you can take to regain your health and improve your performance.

If you're over 40, odds are you're already starting to feel the debilitating effects of low testosterone. Research shows that by the time they are 60 years old, men typically produce **60% less** testosterone than they did at age 20.¹ With this drastic reduction comes several well-documented problems—from the blues and cognitive impairment to reduced sex drive and abdominal weight gain. Normal testosterone has been associated with maintaining a healthy cardiovascular system.²

RESTORING HEALTH AND SEXUAL VIGOR

Life Extension®'s reformulated **Super MiraForte** contains high potencies of **chrysin** and **nettle root**—plant extracts that naturally reduce the aromatization (conversion) of testosterone to estrogen to enhance free testosterone levels.³ **Bioperine®** is included to facilitate the absorption of **chrysin** (a natural flavonoid) into the bloodstream.

Muira puama is a rainforest herb classified in the Brazilian Pharmacopeia as an aphrodisiac. In a trial of men with decreased libido and other sexual issues, **62%** of those taking **muira puama** reported positive results in regard to libido, while **51%** of those with a common sexual problem felt that the herb was helpful.⁴ A second trial examined men with decreased libido and found that **85%** of the test subjects taking **muira puama** enjoyed an enhanced libido, **90%** had improved sexual function, and **100%** of test subjects experienced an increase in intercourse frequency.⁵

To augment these protective effects, a standardized **lignan extract** from Norwegian spruce has been added to Super MiraForte. These lignans convert to enterolactone in the intestine that is then rapidly absorbed into the bloodstream where it provides significant biological effects.⁶ Enterolactone has demonstrated **anti-estrogen** and **anti-DHT** effects that are of particular importance for the aging prostate gland.⁷⁻⁹

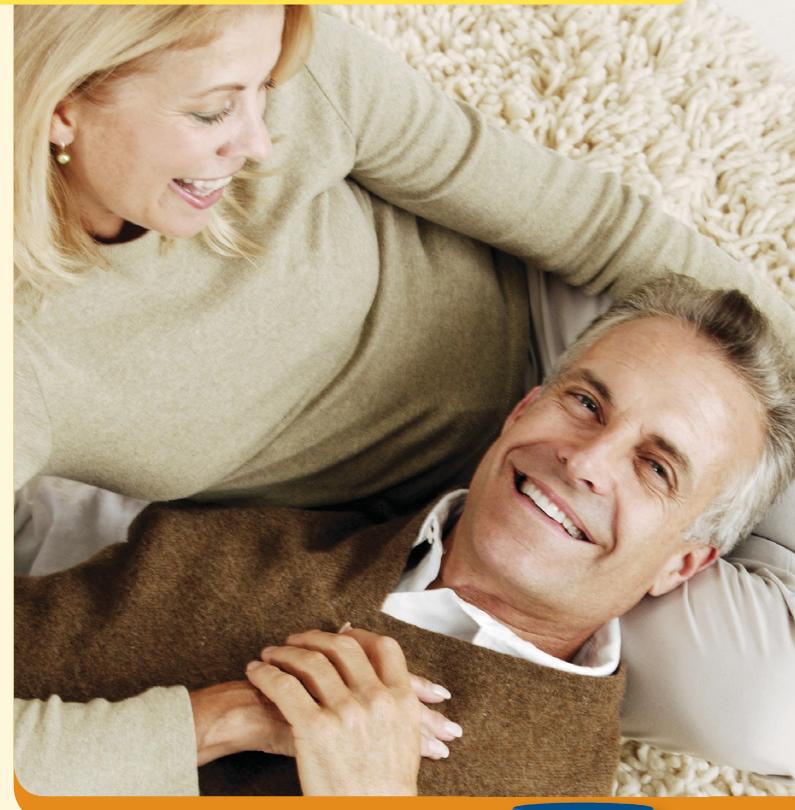
The suggested daily dose of four capsules of Super MiraForte contains potencies of the following nutrients:

Chrysin	1500 mg
Bioperine®	15 mg
Muira puama	850 mg
Nettle root	282 mg
Ginger root	50 mg
Chelated elemental zinc	15 mg
Maca	320 mg
HMRlignan™ Norway	33.4 mg
Spruce lignan extract	

SUPER MIRAFORTE

SUPPORTS HEALTHY TESTOSTERONE LEVELS—NATURALLY

Now With Standardized Lignans



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References

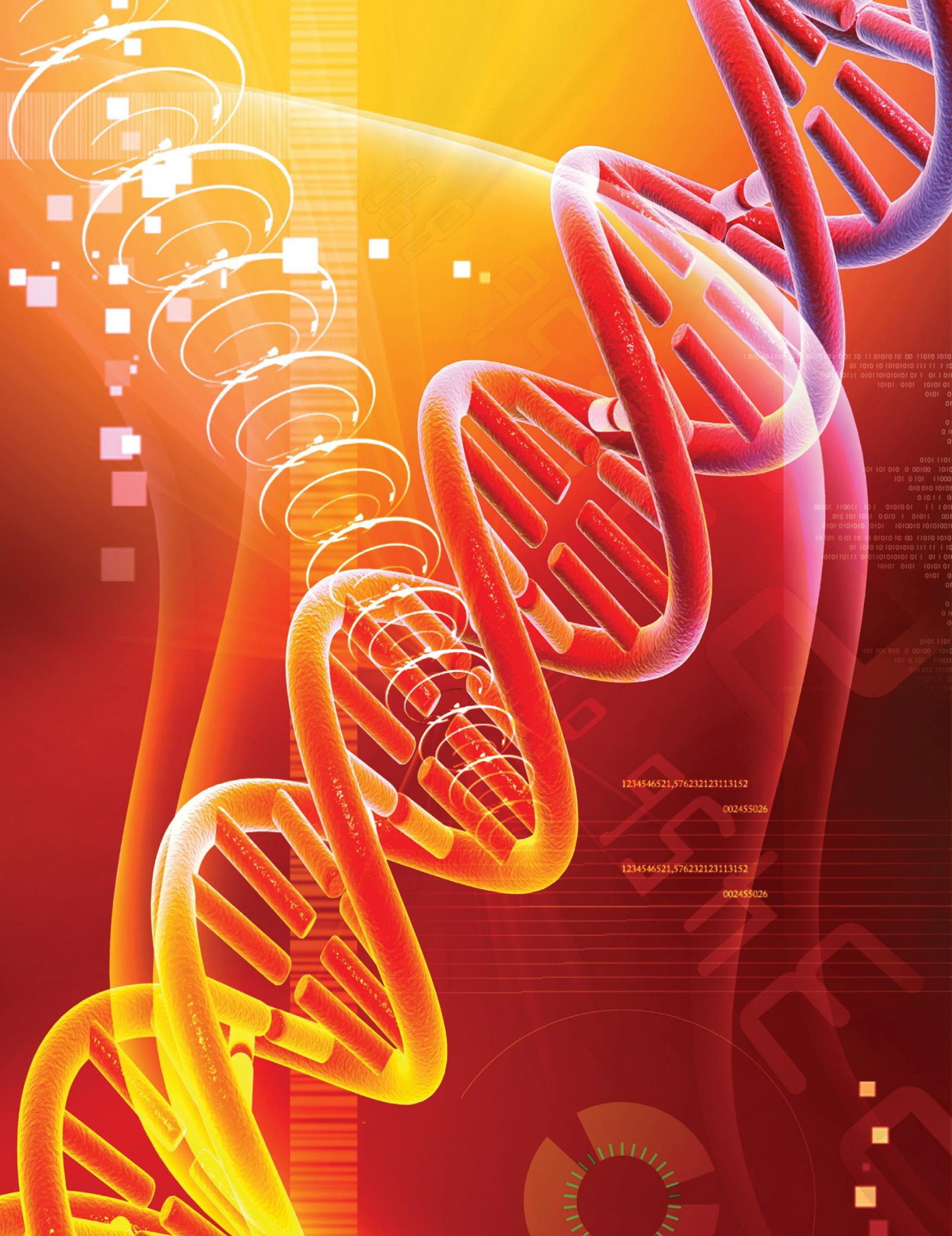
1. *J Clin Endocrinol Metab.* 1991 Nov;73(5):1016-25.
2. *J Clin Endocrinol Metab.* 2002 Aug;87(8):3632-9.
3. *Science.* 1984 Sep 7;225(4666):1032-4.
4. Presented at the First International Congress on Ethnopharmacology, Strasbourg, France. 1990;June 5-9.
5. *Ethnopharmacology.* 1995 Mar.
6. *Exp Biol Med (Maywood).* 2005 Mar;230(3):217-23.
7. *Br J Nutr.* 2007 Aug;98(2):388-96.
8. *Cancer Causes Control.* 2006 Mar;17(2):169-80.
9. *Cancer Epidemiol Biomarkers Prev.* 2005 Jan;14(1):213-20.

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Five Natural Compounds Simulate Caloric Restriction

BY JULIUS GOEPP, MD

The most scientifically validated way to extend life span—from single-celled organisms to mammals—is *caloric restriction*.¹ This technique has been shown to boost life span by nearly 100% in some species.²

Reducing the number of ingested calories—while maintaining healthy intake of essential nutrients—triggers a cascade of anti-aging mechanisms in the body. More than 70 years of research have established its life-extending power.³

Modern science has only recently begun to unravel the mystery behind how *calorie restriction* so radically extends life span. At its core lies favorable alterations in **gene expression**—one of the body's adaptive responses to reduced calorie intake. By activating certain genes and disabling others, caloric restriction dramatically slows aging.

This retardation of aging delays the onset of degenerative disease while improving biomarkers of youth, ranging from **metabolic rate** and **insulin sensitivity** to **cardiac health** and **cognitive function**.⁴

By the same token, the consequences of caloric **overconsumption** are equally profound. Every excess calorie brings you one step closer to age-related illness and death.

This is true at any stage of life, whether you're lean or overweight, regardless of your health or nutritional status.⁵⁻⁷

Eating more than your body needs (**excessive energy intake**) can load the blood with **triglycerides**, **glucose**, **homocysteine**, and **pro-inflammatory chemicals**. The results are accelerated aging processes, prompting deterioration across multiple biomarkers of health.⁸

Most humans find it difficult to submit to a sufficiently *rigorous* dietary regimen, leaving the benefits of caloric restriction tantalizingly out of reach. Until now!

In this article you will discover the most compelling evidence to date on the effects of caloric restriction. In a **milestone 20-year study** of Rhesus monkeys—our close genetic relatives—a modestly restricted diet resulted in a **three-fold reduction in the risk of age-related disease!**⁹

You will also learn about **five** natural compounds that **favorably modulate gene expression** to support a more youthful health profile. These “caloric restriction mimics” *simulate* many of the beneficial effects of caloric restriction through multiple pathways.

Aging individuals may now enjoy some of the youth-promoting, disease-fighting mechanisms of caloric restriction—without strict dietary measures. >>

A Milestone in Longevity Research

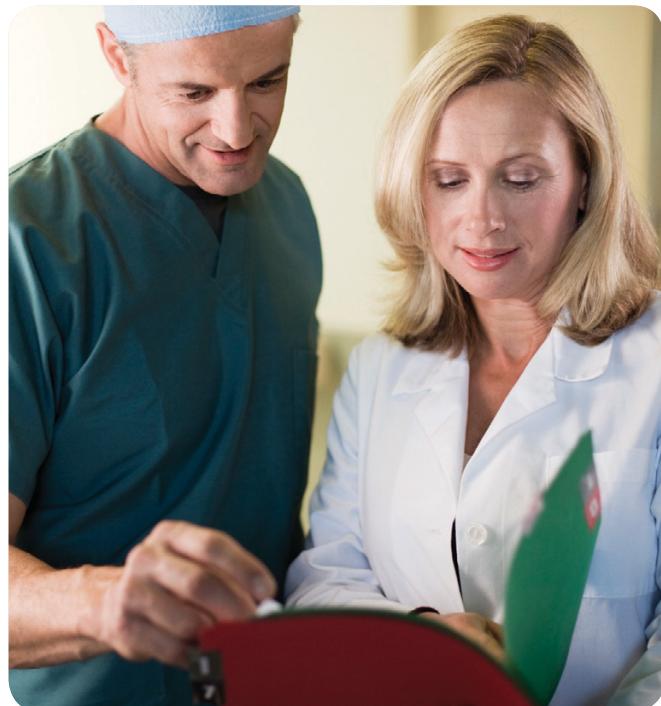
In 1989, a group of anti-aging researchers launched what would prove to be the most comprehensive study of caloric restriction (CR) to date.¹⁰ Rhesus monkeys were chosen as subjects because they exhibit biological and aging characteristics strikingly similar to humans. The results, published in 2009 in the prestigious journal *Science*,⁹ provide compelling evidence of CR's age-delaying, disease-fighting power.

The monkeys were split into two groups. Half were allowed to eat naturally, without restraint; the other half subsisted on a **nutrient-replete diet 30% lower in total calories** than they would normally consume. After **20** years, **37%** of controls had died of age-related causes, as opposed to just **13%** in the calorie-restricted group. In other words, caloric restriction cut degenerative disease risk by **a factor of three**.⁹

This study's findings are even more tantalizing when one sees that it did not require *severe* lowering of calorie intake to produce these striking results. Americans typically consume **100%** more calories than they need. It only required a calorie reduction of **30%** to achieve the remarkable benefits in this study.

The Rhesus monkey study also conclusively demonstrated caloric restriction's protective power. Over the course of **20** years, virtually **all** measured biomarkers of health were superior in the CR group.⁹

These findings have profound implications for humans. Rhesus monkeys are as vulnerable to chronic, age-related disease as we are. They lose their strength



over time. Many become obese, fall prey to metabolic syndrome, and succumb to diabetes. They develop cardiovascular disease and cancer. And like aging humans, their brains shrink as they get older.

Yet among the calorie-restricted group, incidence of cardiovascular disease was **half** the rate of controls. Not **one** member exhibited any symptoms of impaired glucose control or diabetes, whereas **40%** of monkeys who ate as much as they wanted had become diabetic or pre-diabetic.⁹

Calorie-restricted monkeys lost **fat** weight, but did **not** sustain loss of **muscle** mass observed in the control group. CR also inhibited reduction in **brain** volume, especially in areas governing cognitive and motor function.⁹

The robust health of the CR group compared to controls is clearly evident in photographs of the monkeys from the control and CR groups.⁹ (See figure 1.)

Benefits of Calorie Restriction in Humans

All available data indicate that calorie restriction also slows aging and reduces the risk of killer diseases in **humans** as well. Individuals who impose a **20%** reduction in their calorie consumption for **2-6 years** lose **fat** weight and show significant improvement in their markers of aging, including **blood pressure**, **cholesterol** levels, and **glucose** control.¹¹

Even *brief* periods of caloric restriction can temporarily improve core body temperature and insulin

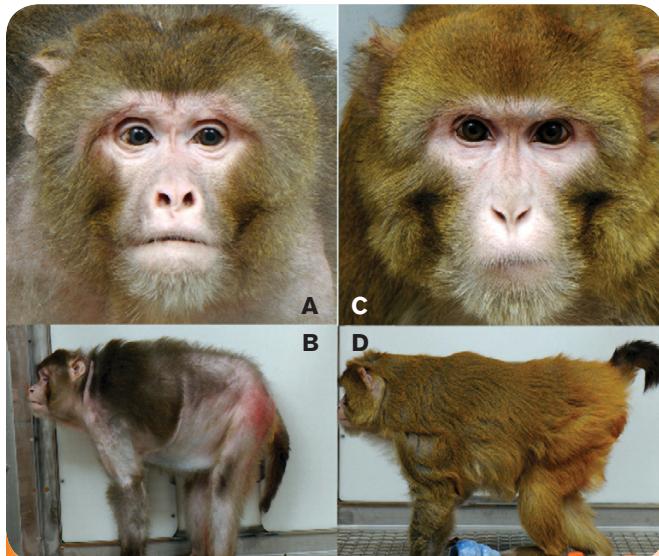


Figure 1. Appearance of Rhesus monkeys in old age (approximately 27.6 years). A and B show a typical control animal. C and D show an age-matched calorically restricted animal.⁹

sensitivity, which happen to be markers of longevity.^{11,12} In clinical studies, short intervals of caloric restriction have reduced *systemic inflammation*—an underlying factor of so many chronic, degenerative diseases.

More youthful heart muscle performance¹³ has also been observed—caloric restriction appears to increase the number of vital energy-producing *mitochondria* in heart and skeletal muscle, reducing the oxidative damage that accelerates aging.¹⁴⁻¹⁶

No other intervention documented in the scientific literature can compete with *caloric restriction* as a candidate for slowing aging and supporting youthful health in humans.

It is regrettable that most aging individuals don't lower calorie intake by the **20-30% reduction** required to reap the benefits of caloric restriction, as it can dramatically lower disease risk and add *years* to their lives.

Caloric Restriction “Mimics” and Gene Expression

The problem many people have in sufficiently reducing their calorie intake has led to a novel scientific solution.

Researchers have identified a select group of nutrients that trigger many of the same underlying mechanisms of action as caloric restriction. Among the most promising of these **caloric restriction mimics** and enhancers are **resveratrol**, **pterostilbene**, **quercetin**, and **grape seed extract**, along with **black tea extract**. These nutraceuticals have been shown to generate many of the same effects in the body as caloric restriction, without significant dietary modification. In particular, they “mimic” caloric restriction's favorable impact on **genes** that influence the aging process.

Genes have the capacity to directly affect life span by regulating a broad spectrum of aging factors, from inflammation and metabolic function to immune response. Calorie restriction exerts a beneficial effect on the *activity of gene expression*, supporting healthy cellular function through numerous physiological pathways. This includes:

- Blocking inflammatory factors
- Optimizing fat and carbohydrate metabolism
- Lowering serum glucose
- Supporting endothelial function
- Inhibiting cancer development and proliferation

Compounds that mimic caloric restriction bring about favorable changes in gene expression and improve the primary biomarkers of aging.

Caloric Restriction

- Excessive caloric intake is strongly associated with the onset of degenerative disease and shorter life span.
- Research shows that calorie restriction (CR) can extend life span and afford significant protection against age-related disease in many animal species.
- A landmark 20-year study demonstrated that caloric restriction powerfully counters the effects of aging in primates closely related to humans. CR produced a **three-fold reduction** in mortality from age-related conditions.
- A caloric restriction regimen strict enough to yield these benefits is difficult or impractical for most people.
- Nutrients known as “caloric restriction mimics” (or mimetics) afford a novel scientific solution. Working in tandem, their biomolecular action exerts similar effects on gene expression, providing the benefits of CR without severe dietary modification.
- Resveratrol, pterostilbene, quercetin, and extracts of grape seed rank among the most promising of CR mimics, while the polyphenols in black tea support these metabolic effects. Research shows that they powerfully inhibit systemic inflammation, enhance mitochondrial health, prevent cancer, and protect brain and heart tissue from age-related deterioration.



Controlling Nuclear Factor-Kappa B

Calorie restriction sharply limits expression of **nuclear factor-kappa B** (NF-**kB**). In the cells, **NF-**kB**** is a critical **gene regulator** that governs response to pro-inflammatory cytokines, free radicals, cholesterol levels, immune function, and cancer prevention.¹⁷⁻²⁴

The nutrients that mimic caloric restriction also act to *stabilize* NF-**kB** regulation and help combat the age-related conditions associated with unfavorable NF-**kB** activity.

Resveratrol activates **sirtuins**,²⁵ a powerful family of “information regulator” proteins that inhibit NF-**kB**, reducing inflammation throughout the body, such as that caused by second-hand cigarette smoke.²⁴ Resveratrol also prevents inflammatory mast cells from releasing the histamines that trigger asthma and allergic reactions.²⁶

Resveratrol radically decreases production of the **adhesion molecules** that attract inflammatory cells to vascular walls, one of the principal mechanisms of atherosclerosis.^{27,28} Adhesion molecules also permit cancer cells to invade tissue and metastasize. Resveratrol’s influence over NF-**kB** has also enhanced beneficial UV-induced programmed cell death (*apoptosis*) in skin cancer cells.²⁹

Found in blueberries, **pterostilbene** is a polyphenol closely related to resveratrol.³⁰ It limits NF-**kB** activity through multiple complementary mechanisms.³¹ In vitro, pterostilbene suppresses invasive tumor activity and enhances therapeutic destruction of cancer cells.^{32,33}

Quercetin’s ability to protect against chronic inflammatory conditions such as asthma, inflammatory bowel disease, and arthritis is due in part to its capacity for NF-**kB** inhibition.^{34,35}

Grape seed extract also disrupts cellular inflammation signaling by blocking NF-**kB**.³⁶ Its effect on pro-inflammatory cytokine production in fat cells may even help combat obesity and type 2 diabetes.³⁷

Black tea extract’s influence on NF-**kB** activity has been shown to specifically protect cells from damage associated with inflammation,³⁸ cancer,^{39,40} Parkinson’s disease,⁴¹ and stroke.⁴²

Suppressing Inflammatory Cytokines

Caloric restriction inhibits expression of genes that produce inflammatory cytokines—such as interleukins and tumor necrosis factor (TNF), as well as cyclooxygenase-2 (COX-2)—that are powerfully implicated in the onset of cancer, atherosclerosis, and chronic inflammation.⁴³⁻⁴⁶

Resveratrol and **pterostilbene** block the release of a host of inflammatory cytokines and enzymes found throughout the body—particularly tissues and organs stressed by environmental factors, infection, or trauma.^{22,47,48} Pterostilbene powerfully suppresses the expression of inflammatory COX-2,^{31,49} helping lower the risk of cancer as well as a host of inflammatory diseases.^{50,51}

Quercetin inhibits the COX-2 enzyme involved in early-stage colon cancer, and suppresses numerous cytokines involved in allergies and autoimmune disorders.^{34,35,53}

Grape seed extract specifically prevents fatty tissue from releasing inflammatory cytokines and adipokines that would otherwise provoke insulin resistance and atherosclerosis associated with metabolic syndrome.³⁷ By downregulating reactive cytokines, grape seed extract has been shown in animals to protect organs from ischemia-reperfusion damage (injury caused by the return of blood flow after a period of restriction).⁵⁴

Black tea extract reduces levels of **STAT-1**, a protein which “tells” the cell nucleus to activate genes that produce inflammatory cytokines.⁵⁵

Cancer Prevention

Calorie restriction (CR) upregulates genes that suppress cancer and downregulates genes that permit cancers to form or spread. CR prevents cancer cell reproduction and proliferation, while inhibiting the blood vessel growth cancer cells require to develop and metastasize.^{46,56-62}

Resveratrol and **pterostilbene** mirror these effects. They combat cancer at every stage of development, inducing apoptosis (programmed cell death) in a variety of human cancer types, while preserving healthy cells.^{32,63-66} Resveratrol also suppresses cancer proliferation by modulating expression of proteins involved in the reproductive cycle of abnormal cells.^{67,68}

Quercetin activates “executioner” proteins while inhibiting survival proteins in human cancer cells, blocking their reproduction.⁶⁹⁻⁷¹ Quercetin and resveratrol have also been shown to block the expression of *vascular endothelial growth factor* (VEGF), an effect that may help starve tumors of their blood supply.⁷²

Grape seed extract induces expression of a protein that arrests cancer cells early in their reproductive cycle, preventing further development and destroying them.⁷³ Similar to quercetin, grape seed extract fights angiogenesis by suppressing the VEGF signaling pathway.⁷⁴

Black tea extract reduces expression of genes that cancer cells use to proliferate, survive, infiltrate healthy tissue, supply themselves with blood, and metastasize to other organs.⁷⁵ It has also been shown to upregulate expression of proteins that arrest the cell reproductive cycle and induce cellular death specifically in cancers.⁷⁶

Enhanced Glucose Control

Caloric restriction enhances glucose control.^{77,78} Recall that not one of the Rhesus monkeys in the study discussed earlier developed diabetes or exhibited symptoms of impaired glucose control.⁹ Caloric restriction triggers gene regulators called *peroxisome proliferator-activated receptors* (PPARs), a class of proteins responsible for healthy fat and carbohydrate metabolism. They also play key roles in optimizing mitochondrial health^{57,59} and thwarting the onset of metabolic syndrome and diabetes.⁷⁹

Resveratrol⁸⁰ and **pterostilbene**^{81,82} upregulate the production and activity of PPAR, launching a set of cellular processes that support a youthful metabolic profile. The PPAR activator resveratrol has been shown to:

- Prevent fat cells from absorbing sugar and converting it to fat⁸³
- Reduce inflammation and insulin resistance in fat cells⁸⁴
- Boost mitochondrial function⁸⁵



Grape seed extract modulates a *different* set of PPARs that regulate fat storage. Grape seed extract induces **fat** metabolism while inhibiting the development of new fat cells.^{86,87} It also protects endothelial cells by preventing the inflammatory response to proteins damaged by glucose (the age-accelerating process known as **glycation**).⁸⁸

Resveratrol further exerts a favorable influence on blood sugar metabolism at the cellular level, reducing glucose production in liver cells in a way that mimics prolonged calorie restriction.⁸⁹ In diabetic animals, resveratrol has been shown to help restore blood sugar to normal by modulating the activity of several enzymes involved in sugar metabolism.⁹⁰

Pterostilbene and **grape seed extract** generate similar beneficial changes that help promote healthy blood sugar levels.^{91,92} Grape seed extract activates genes that trigger **glucose uptake**. This assists cells in the *absorption* and removal of glucose from circulation.⁹²

Quercetin has been shown to stimulate the proliferation of pancreatic cells that help modulate blood glucose levels in both diabetic and non-diabetic animal models.⁹³ It also markedly reduces expression of the enzyme that produces **sorbitol**, a sugar alcohol known to cause cataracts and blindness.⁹⁴

Black tea extract polyphenols inhibit **lipase**,⁹⁵ an enzyme that breaks down fat in the stomach and small intestines. This helps block *absorption* of fat into the bloodstream.⁹⁶

In animal models, the *theaflavins* in black tea extract prevent after-meal elevations in blood glucose and may protect against the *metabolic syndrome*.^{97,98} This effect may help increase signaling for a powerful longevity factor called **FOXO1a**.⁹⁹

Summary

Caloric restriction (CR) is the most scientifically validated method shown to reliably extend life span in multiple species, from microorganisms to mammals.

A milestone 20-year study provides the most conclusive evidence to date of its efficacy in Rhesus monkeys, our close genetic relatives. The discovery of calorie restriction-mimicking nutrients makes it possible for aging humans to *emulate* some of CR's beneficial mechanisms of action, especially as an adjunct to modestly reducing one's overall calorie intake.¹⁰⁰ The unique ability of these nutrients to modulate **gene expression** exerts system-wide effects that, in addition to influencing many of the same pathways activated by calorie restriction, can also significantly reduce degenerative disease risk. ●

If you have any questions on the scientific content of this article, please call a Life Extension® Health Advisor at 1-866-864-3027.

References

1. Genaro Pde S, Sarkis KS, Martini LA. Effect of caloric restriction on longevity. *Arq Bras Endocrinol Metabol*. 2009 Jul;153(5):667-72.
2. McCay CM, Crowell MF. *The Scientific Monthly*. 1934 Nov;39(5):405-14.
3. McCay CM, Crowell MF, Maynard LA. The effect of retarded growth upon the length of life span and upon the ultimate body size. *Nutrition*. 1935;5:155-71.
4. Heilbronn L, Ravussin E. Calorie restriction and aging: review of the literature and implications for studies in humans. *Am J Clin Nutr*. 2003 Sept;78(3):361-9.
5. Fontana L, Villareal DT, Weiss EP, et al. Calorie restriction or exercise: effects on coronary heart disease risk factors. A randomized, controlled trial. *Am J Physiol Endocrinol Metab*. 2007 Jul;293(1):E197-202.
6. Tan CY, Vidal-Puig A. Adipose tissue expandability: the metabolic problems of obesity may arise from the inability to become more obese. *Biochem Soc Trans*. 2008 Oct;36(Pt 5):935-40.
7. Korner J, Woods SC, Woodworth KA. Regulation of energy homeostasis and health consequences in obesity. *Am J Med*. 2009 Apr;122(4 Suppl 1):S12-8.
8. Fontana L. Nutrition, adiposity and health. *Epidemiol Prev*. 2007 Sep-Oct;31(5):290-4.
9. Colman RJ, Anderson RM, Johnson SC, et al. Caloric restriction delays disease onset and mortality in rhesus monkeys. *Science*. 2009 Jul 10;325(5937):201-4.
10. Colman RJ, Roecker EB, Ramsey JJ, Kemnitz JW. The effect of dietary restriction on body composition in adult male and female rhesus macaques. *Aging (Milano)*. 1998 Apr;10(2):83-92.
11. Everitt AV, Le Couteur DG. Life extension by calorie restriction in humans. *Ann N Y Acad Sci*. 2007 Oct;1114:428-33.
12. Heilbronn LK, de Jonge L, Frisard MI, et al. Effect of 6-month calorie restriction on biomarkers of longevity, metabolic adaptation, and oxidative stress in overweight individuals: a randomized controlled trial. *JAMA*. 2006 Apr 5;295(13):1539-48.
13. Holloszy JO, Fontana L. Caloric restriction in humans. *Exp Gerontol*. Aug 2007;42(8):709-12.
14. Menshikova EV, Ritov VB, Toledo FG, Ferrell RE, Goodpaster BH, Kelley DE. Effects of weight loss and physical activity on skeletal muscle mitochondrial function in obesity. *Am J Physiol Endocrinol Metab*. 2005 Apr;288(4):E818-25.
15. Civitarese AE, Carling S, Heilbronn LK, et al. Calorie restriction increases muscle mitochondrial biogenesis in healthy humans. *PLoS Med*. 2007 Mar;4(3):e76.
16. Heilbronn LK, Gan SK, Turner N, Campbell LV, Chisholm DJ. Markers of mitochondrial biogenesis and metabolism are lower in overweight and obese insulin-resistant subjects. *J Clin Endocrinol Metab*. 2007 Apr;92(4):1467-73.
17. Zou Y, Yoon S, Jung KJ, et al. Upregulation of aortic adhesion molecules during aging. *J Gerontol A Biol Sci Med Sci*. 2006 Mar;61(3):232-44.
18. Jung KJ, Maruyama N, Ishigami A, Yu BP, Chung HY. The redox-sensitive DNA binding sites responsible for age-related downregulation of SMP30 by ERK pathway and reversal by calorie restriction. *Antioxid Redox Signal*. 2006 Mar-Apr;8(3-4):671-80.
19. Kim HJ, Yu BP, Chung HY. Molecular exploration of age-related NF- κ B/IKK downregulation by calorie restriction in rat kidney. *Free Radic Biol Med*. 2002 May 15;32(10):991-1005.
20. Chandrasekar B, Nelson JF, Colston JT, Freeman GL. Calorie restriction attenuates inflammatory responses to myocardial ischemia-reperfusion injury. *Am J Physiol Heart Circ Physiol*. 2001 May;280(5):H2094-102.
21. Chung HY, Kim HJ, Kim KW, Choi JS, Yu BP. Molecular inflammation hypothesis of aging based on the anti-aging mechanism of calorie restriction. *Microsc Res Tech*. 2002 Nov 15;59(4):264-72.
22. Chung HY, Kim HJ, Kim JW, Yu BP. The inflammation hypothesis of aging: molecular modulation by calorie restriction. *Ann N Y Acad Sci*. 2001 Apr;928:327-35.
23. Kim DH, Kim JY, Yu BP, Chung HY. The activation of NF- κ B through Akt-induced FOXO1 phosphorylation during aging and its modulation by calorie restriction. *Biogerontology*. 2008 Feb;9(1):33-47.
24. Yang SR, Wright J, Bauter M, Seweryniak K, Kode A, Rahman I. Sirtuin regulates cigarette smoke-induced proinflammatory mediator release via RelA/p65 NF- κ B in macrophages in vitro and in rat lungs in vivo: implications for chronic inflammation and aging. *Am J Physiol Lung Cell Mol Physiol*. 2007 Feb;292(2):L567-76.
25. Pallas M, Casadesus G, Smith MA, et al. Resveratrol and neurodegenerative diseases: activation of SIRT1 as the potential pathway towards neuroprotection. *Curr Neurovasc Res*. 2009 Feb;6(1):70-81.
26. Kang OH, Jang HJ, Chae HS, et al. Anti-inflammatory mechanisms of resveratrol in activated HMC-1 cells: pivotal roles of NF- κ B and MAPK. *Pharmacol Res*. 2009 May;59(5):330-7.
27. Park HJ, Jeong SK, Kim SR, et al. Resveratrol inhibits *Porphyromonas gingivalis* lipopolysaccharide-induced endothelial adhesion molecule expression by suppressing NF- κ B activation. *Arch Pharm Res*. 2009 Apr;32(4):583-91.
28. Park JS, Kim KM, Kim MH, et al. Resveratrol inhibits tumor cell adhesion to endothelial cells by blocking ICAM-1 expression. *Anticancer Res*. 2009 Jan;29(1):355-62.
29. Roy P, Kalra N, Nigam N, et al. Resveratrol enhances ultraviolet B-induced cell death through nuclear factor- κ B pathway in human epidermoid carcinoma A431 cells. *Biochem Biophys Res Commun*. 2009 Jun 26;384(2):215-20.
30. Rimando AM, Kalt W, Magee JB, Dewey J, Ballington JR. Resveratrol, pterostilbene, and piceatannol in Vaccinium berries. *J Agric Food Chem*. 2004 Jul 28;52(15):4713-9.



31. Cichocki M, Paluszczak J, Szaefer H, Piechowiak A, Rimando AM, Baer-Dubowska W. Pterostilbene is equally potent as resveratrol in inhibiting 12-O-tetradecanoylphorbol-13-acetate activated NFκB, AP-1, COX-2, and iNOS in mouse epidermis. *Mol Nutr Food Res*. 2008 Jun;52 Suppl 1:S62-70.

32. Pan MH, Chiou YS, Chen WJ, Wang JM, Badmaev V, Ho CT. Pterostilbene inhibited tumor invasion via suppressing multiple signal transduction pathways in human hepatocellular carcinoma cells. *Carcinogenesis*. 2009 Jul;30(7):1234-42.

33. Priego S, Feddi F, Ferrer P, et al. Natural polyphenols facilitate elimination of HT-29 colorectal cancer xenografts by chemoradiotherapy: a Bcl-2- and superoxide dismutase 2-dependent mechanism. *Mol Cancer Ther*. 2008 Oct;7(10):3330-42.

34. Ruiz PA, Braune A, Holzswimmer G, Quintanilla-Fend L, Haller D. Quercetin inhibits TNF-induced NF-κB transcription factor recruitment to proinflammatory gene promoters in murine intestinal epithelial cells. *J Nutr*. 2007 May;137(5):1208-15.

35. Min YD, Choi CH, Bark H, et al. Quercetin inhibits expression of inflammatory cytokines through attenuation of NF-κB and p38 MAPK in HMC-1 human mast cell line. *Inflamm Res*. 2007 May;56(5):210-5.

36. Terra X, Valls J, Vitrac X, et al. Grape-seed procyanidins act as antiinflammatory agents in endotoxin-stimulated RAW 264.7 macrophages by inhibiting NF-κB signaling pathway. *J Agric Food Chem*. 2007 May 30;55(11):4357-65.

37. Chacon MR, Ceperuelo-Mallafré V, Maymo-Masip E, et al. Grape-seed procyanidins modulate inflammation on human differentiated adipocytes in vitro. *Cytokine*. 2009 Aug;47(2):137-42.

38. Ukil A, Maity S, Das PK. Protection from experimental colitis by theaflavin-3,3'-digallate correlates with inhibition of IKK and NF-κB activation. *Br J Pharmacol*. 2006 Sep;149(1):121-31.

39. Roy P, Nigam N, Singh M, et al. Tea polyphenols inhibit cyclooxygenase-2 expression and block activation of nuclear factor-κB and Akt in diethylnitrosoamine induced lung tumors in Swiss mice. *Invest New Drugs*. 2009 Jun 11.

40. Patel R, Krishnan R, Ramchandani A, Maru G. Polymeric black tea polyphenols inhibit mouse skin chemical carcinogenesis by decreasing cell proliferation. *Cell Prolif*. 2008 Jun;41(3):532-53.

41. Levites Y, Youdim MB, Maor G, Mandel S. Attenuation of 6-hydroxydopamine (6-OHDA)-induced nuclear factor-κB (NF-κB) activation and cell death by tea extracts in neuronal cultures. *Biochem Pharmacol*. 2002 Jan 1;63(1):21-9.

42. Cai F, Li C, Wu J, et al. Modulation of the oxidative stress and nuclear factor κB activation by theaflavin 3,3'-gallate in the rats exposed to cerebral ischemia-reperfusion. *Folia Biol (Praha)*. 2007;53(5):164-72.

43. Chung HY, Cesari M, Anton S, et al. Molecular inflammation: underpinnings of aging and age-related diseases. *Ageing Res Rev*. 2009 Jan;8(1):18-30.

44. Kim YJ, Kim HJ, No JK, Chung HY, Fernandes G. Anti-inflammatory action of dietary fish oil and calorie restriction. *Life Sci*. 2006 Apr 18;78(21):2523-32.

45. Miliaras S, Anogeianaki A, Meditskou S, et al. Effects of rich-in-fat diets and highly selective COX-2 inhibitors on 7,12-dimethylbenz(A)-anthracene-induced tumor growth. *Int J Immunopathol Pharmacol*. 2009 Apr-Jun;22(2):323-32.

46. Jung KJ, Lee EK, Kim JY, et al. Effect of short term calorie restriction on pro-inflammatory NF-κB and AP-1 in aged rat kidney. *Inflamm Res*. 2009 Mar;58(3):143-50.

47. Culpitt SV, Rogers DF, Fenwick PS, et al. Inhibition by red wine extract, resveratrol, of cytokine release by alveolar macrophages in COPD. *Thorax*. 2003 Nov;58(11):942-6.

48. Wu CT, Yu HP, Chung CY, Lau YT, Liao SK. Attenuation of lung inflammation and pro-inflammatory cytokine production by resveratrol following trauma-hemorrhage. *Chin J Physiol*. 2008 Dec 31;51(6):363-8.

49. Pan MH, Chang YH, Tsai ML, et al. Pterostilbene suppressed lipopolysaccharide-induced up-expression of iNOS and COX-2 in murine macrophages. *J Agric Food Chem*. 2008 Aug 27;56(16):7502-9.

50. de Souza Pereira R. Selective cyclooxygenase-2 (COX-2) inhibitors used for preventing or regressing cancer. *Recent Pat Anticancer Drug Discov*. 2009 Jun;4(2):157-63.

51. Tunon MJ, García-Mediavilla MV, Sánchez-Campos S, González-Gallego J. Potential of flavonoids as anti-inflammatory agents: modulation of pro-inflammatory gene expression and signal transduction pathways. *Curr Drug Metab*. 2009 Mar;10(3):256-71.

52. Cruz-Correia M, Shoskes DA, Sanchez P, et al. Combination treatment with curcumin and quercetin of adenomas in familial adenomatous polyposis. *Clin Gastroenterol Hepatol*. 2006 Aug;4(8):1035-8.

53. García-Mediavilla V, Crespo I, Collado PS, et al. The anti-inflammatory flavones quercetin and kaempferol cause inhibition of inducible nitric oxide synthase, cyclooxygenase-2 and reactive C-protein, and down-regulation of the nuclear factor κB pathway in Chang Liver cells. *Eur J Pharmacol*. 2007 Feb 28;557(2-3):221-9.

54. Sehirli O, Ozel Y, Dulundu E, Topaloglu U, Ercan F, Sener G. Grape seed extract treatment reduces hepatic ischemia-reperfusion injury in rats. *Phytother Res*. 2008 Jan;22(1):43-8.

55. Cai F, Li CR, Wu JL, et al. Theaflavin ameliorates cerebral ischemia-reperfusion injury in rats through its anti-inflammatory effect and modulation of STAT-1. *Mediators Inflamm*. 2006;2006(5):30490.

56. Alcain FJ, Villalba JM. Sirtuin inhibitors. *Expert Opin Ther Pat*. 2009 Mar;19(3):283-94.

57. Alcain FJ, Villalba JM. Sirtuin activators. *Expert Opin Ther Pat*. 2009 Apr;19(4):403-14.

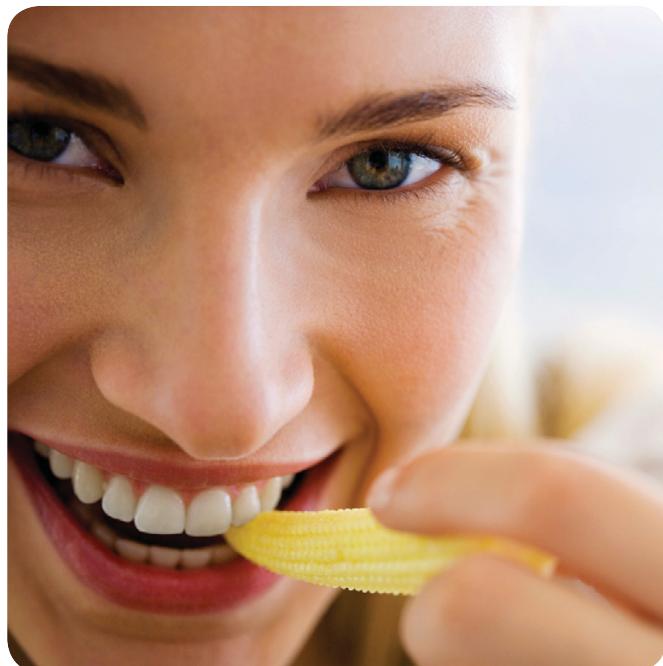
58. Kanfi Y, Peshti V, Gozlan YM, Rathaus M, Gil R, Cohen HY. Regulation of SIRT1 protein levels by nutrient availability. *FEBS Lett*. 2008 Jul 9;582(16):2417-23.

59. Al-Regaiey KA, Masternak MM, Bonkowski MS, Panici JA, Kopchick JJ, Bartke A. Effects of caloric restriction and growth hormone resistance on insulin-related intermediates in the skeletal muscle. *J Gerontol A Biol Sci Med Sci*. 2007 Jan;62(1):18-26.

60. Fay JR, Steele V, Crowell JA. Energy homeostasis and cancer prevention: the AMP-activated protein kinase. *Cancer Prev Res (Phila Pa)*. 2009 Apr;2(4):301-9.

61. Potente M, Dimmeler S. Emerging roles of SIRT1 in vascular endothelial homeostasis. *Cell Cycle*. 2008 Jul 15;7(14):2117-22.

62. Kritchevsky D. Caloric restriction and cancer. *J Nutr Sci Vitaminol (Tokyo)*. 2001 Feb;47(1):13-9.



63. Pan MH, Chang YH, Tsai ML, et al. Pterostilbene suppressed lipopolysaccharide-induced up-expression of iNOS and COX-2 in murine macrophages. *J Agric Food Chem.* 2008 Aug 27;56(16):7502-9.

64. Ferrer P, Asensi M, Priego S, et al. Nitric oxide mediates natural polyphenol-induced Bcl-2 down-regulation and activation of cell death in metastatic B16 melanoma. *J Biol Chem.* 2007 Feb 2;282(5):2880-90.

65. Pan MH, Chang YH, Badmaev V, Nagabhushanam K, Ho CT. Pterostilbene induces apoptosis and cell cycle arrest in human gastric carcinoma cells. *J Agric Food Chem.* 2007 Sep 19;55(19):7777-85.

66. Pan MH, Lin YT, Lin CL, Wei CS, Ho CT, Chen WJ. Suppression of Heregulin-beta1/HER2-Modulated Invasive and Aggressive Phenotype of Breast Carcinoma by Pterostilbene via Inhibition of Matrix Metalloproteinase-9, p38 Kinase Cascade and Akt Activation. *Evid Based Complement Alternat Med.* 2009 Jul 16.

67. Hsieh TC. Uptake of resveratrol and role of resveratrol-targeting protein, quinone reductase 2, in normally cultured human prostate cells. *Asian J Androl.* 2009 Sep 21.

68. Gescher AJ. Resveratrol from red grapes - pedestrian polyphenol or useful anticancer agent? *Planta Med.* 2008 Oct;74(13):1651-5.

69. Chien SY, Wu YC, Chung JG, et al. Quercetin-induced apoptosis acts through mitochondrial- and caspase-3-dependent pathways in human breast cancer MDA-MB-231 cells. *Hum Exp Toxicol.* 2009 Aug;28(8):493-503.

70. Kang JW, Kim JH, Song K, Kim SH, Yoon JH, Kim KS. Kaempferol and quercetin, components of Ginkgo biloba extract (EGb 761), induce caspase-3-dependent apoptosis in oral cavity cancer cells. *Phytother Res.* 2009 Jul 7.

71. Tan J, Wang B, Zhu L. Regulation of survivin and Bcl-2 in HepG2 cell apoptosis induced by quercetin. *Chem Biodivers.* 2009 Jul;6(7):1101-10.

72. Kaindl U, Eyberg I, Rohr-Udilova N, Heinzle C, Marian B. The dietary antioxidants resveratrol and quercetin protect cells from exogenous pro-oxidative damage. *Food Chem Toxicol.* 2008 Apr;46(4):1320-6.

73. Kaur M, Singh RP, Gu M, Agarwal R, Agarwal C. Grape seed extract inhibits in vitro and in vivo growth of human colorectal carcinoma cells. *Clin Cancer Res.* 2006 Oct 15;12(20 Pt 1):6194-202.

74. Wen W, Lu J, Zhang K, Chen S. Grape seed extract inhibits angiogenesis via suppression of the vascular endothelial growth factor receptor signaling pathway. *Cancer Prev Res (Phila Pa).* 2008 Dec;1(7):554-61.

75. Letchoumy PV, Mohan KV, Prathiba D, Hara Y, Nagini S. Comparative evaluation of antiproliferative, antiangiogenic and apoptosis inducing potential of black tea polyphenols in the hamster buccal pouch carcinogenesis model. *J Carcinog.* 2007;6:19.

76. Prasad S, Kaur J, Roy P, Kalra N, Shukla Y. Theaflavins induce G2/M arrest by modulating expression of p21waf1/cip1, cdc25C and cyclin B in human prostate carcinoma PC-3 cells. *Life Sci.* 2007 Oct 13;81(17-18):1323-31.

77. Ugochukwu NH, Figgers CL. Modulation of the flux patterns in carbohydrate metabolism in the livers of streptozotocin-induced diabetic rats by dietary caloric restriction. *Pharmacol Res.* 2006 Sep;54(3):172-80.

78. Liang F, Kume S, Koya D. SIRT1 and insulin resistance. *Nat Rev Endocrinol.* 2009 Jul;5(7):367-73.

79. Carvajal K, Hernández-Esquível Mde L, Moreno-Sánchez R. PPARs, metabolic syndrome and cardiac diseases. *Arch Cardiol Mex.* 2007 Oct-Dec;77 Suppl 4:S4-66-76.

80. Cheng G, Zhang X, Gao D, Jiang X, Dong W. Resveratrol inhibits MMP-9 expression by up-regulating PPAR alpha expression in an oxygen glucose deprivation-exposed neuron model. *Neurosci Lett.* 2009 Feb 20;451(2):105-8.

81. Mizuno CS, Ma G, Khan S, Patny A, Avery MA, Rimando AM. Design, synthesis, biological evaluation and docking studies of pterostilbene analogs inside PPARalpha. *Bioorg Med Chem.* 2008 Apr 1;16(7):3800-8.

82. Rimando AM, Nagmani R, Feller DR, Yokoyama W. Pterostilbene, a new agonist for the peroxisome proliferator-activated receptor alpha-isoform, lowers plasma lipoproteins and cholesterol in hypercholesterolemic hamsters. *J Agric Food Chem.* 2005 May 4;53(9):3403-7.

83. Floyd ZE, Wang ZQ, Kilroy G, Cefalu WT. Modulation of peroxisome proliferator-activated receptor gamma stability and transcriptional activity in adipocytes by resveratrol. *Metabolism.* 2008 Jul;57(7 Suppl 1):S32-8.

84. Kennedy A, Overman A, Lapoint K, et al. Conjugated linoleic acid-mediated inflammation and insulin resistance in human adipocytes are attenuated by resveratrol. *J Lipid Res.* 2009 Feb;50(2):225-32.

85. Lopez-Lluch G, Irusta PM, Navas P, de Cabo R. Mitochondrial biogenesis and healthy aging. *Exp Gerontol.* 2008 Sep;43(9):813-9.

86. Pinent M, Blade MC, Salvado MJ, Arola L, Ardevol A. Intracellular mediators of procyanidin-induced lipolysis in 3T3-L1 adipocytes. *J Agric Food Chem.* 2005 Jan 26;53(2):262-6.

87. Pinent M, Blade MC, Salvado MJ, et al. Grape-seed derived procyanidins interfere with adipogenesis of 3T3-L1 cells at the onset of differentiation. *Int J Obes (Lond).* 2005 Aug;29(8):934-41.

88. Ma L, Gao HQ, Li BY, Ma YB, You BA, Zhang FL. Grape seed proanthocyanidin extracts inhibit vascular cell adhesion molecule expression induced by advanced glycation end products through activation of peroxisome proliferators-activated receptor gamma. *J Cardiovasc Pharmacol.* 2007 May;49(5):293-8.

89. Liu Y, Dentin R, Chen D, et al. A fasting inducible switch modulates gluconeogenesis via activator/coactivator exchange. *Nature.* 2008 Nov 13;456(7219):269-73.

90. Palsamy P, Subramanian S. Modulatory effects of resveratrol on attenuating the key enzymes activities of carbohydrate metabolism in streptozotocin-nicotinamide-induced diabetic rats. *Chem Biol Interact.* 2009 May 15;179(2-3):356-62.

91. Pari L, Satheesh MA. Effect of pterostilbene on hepatic key enzymes of glucose metabolism in streptozotocin- and nicotinamide-induced diabetic rats. *Life Sci.* 2006 Jul 10;79(7):641-5.

92. Pinent M, Blay M, Blade MC, Salvado MJ, Arola L, Ardevol A. Grape seed-derived procyanidins have an antihyperglycemic effect in streptozotocin-induced diabetic rats and insulinomimetic activity in insulin-sensitive cell lines. *Endocrinology.* 2004 Nov;145(11):4985-90.

93. Vessal M, Hemmati M, Vasei M. Antidiabetic effects of quercetin in streptozocin-induced diabetic rats. *Comp Biochem Physiol C Toxicol Pharmacol.* 2003 Jul;135C(3):357-64.

94. Ramana BV, Raju TN, Kumar VV, Reddy PU. Defensive role of quercetin against imbalances of calcium, sodium, and potassium in galactosemic cataract. *Biol Trace Elem Res.* 2007 Oct;119(1):35-41.

95. Kusano R, Andou H, Fujieda M, Tanaka T, Matsuo Y, Kouno I. Polymer-like polyphenols of black tea and their lipase and amylase inhibitory activities. *Chem Pharm Bull (Tokyo).* 2008 Mar;56(3):266-72.

96. Kobayashi M, Ichitani M, Suzuki Y, et al. Black-tea polyphenols suppress postprandial hypertriacylglycerolemia by suppressing lymphatic transport of dietary fat in rats. *J Agric Food Chem.* 2009 Aug 12;57(15):7131-6.

97. Shoji Y, Nakashima H. Glucose-lowering effect of powder formulation of African black tea extract in KK-A(y)TaJcl diabetic mouse. *Arch Pharm Res.* 2006 Sep;29(9):786-94.

98. Ramadan G, El-Beih NM, Abd El-Ghaffar EA. Modulatory effects of black v. green tea aqueous extract on hyperglycaemia, hyperlipidaemia and liver dysfunction in diabetic and obese rat models. *Br J Nutr.* 2009 Oct 13:1-9.

99. Cameron AR, Anton S, Melville L, et al. Black tea polyphenols mimic insulin/insulin-like growth factor-1 signalling to the longevity factor FOXO1a. *Aging Cell.* 2008 Jan;7(1):69-77.

100. Morris BJ. A forkhead in the road to longevity: the molecular basis of lifespan becomes clearer. *J Hypertens.* 2005 Jul;23(7):1285-3.

ARE YOU TAKING THE OPTIMAL FORMS OF VITAMIN E?

According to the *Proceedings of the National Academy of Sciences*, **alpha** tocopherol (regular vitamin E) displaces critically important **gamma** tocopherol in the cells.¹ While **alpha** tocopherol inhibits free-radical production, **gamma** tocopherol is required to trap and neutralize existing free radicals.²

Four prestigious scientific journals have highlighted **gamma tocopherol** as one of the most critically important forms of **vitamin E** for those seeking optimal health benefits.

Most commercial vitamin E supplements contain little, if any, **gamma tocopherol**. They instead rely on **alpha tocopherol** as the primary ingredient. However, it is **gamma tocopherol** (not the **alpha** form) that quenches **peroxynitrite**, the free radical that plays a major role in the development of **age-related decline**.^{2,3}

SESAME LIGNANS: The Natural Vitamin E Booster

Life Extension® has uncovered research suggesting that adding **sesame lignans** to **gamma tocopherol** may significantly enhance its beneficial effects. Sesame and its lignans have been shown to boost antioxidant levels and help maintain already-normal blood pressure.*

In a human study that combined **gamma tocopherol** with **sesame lignans**, **gamma tocopherol/sesame** was **25% more effective** than **gamma tocopherol/tocotrienols** in suppressing tissue measurements for free-radical and inflammatory damage.^{4,5} Since tocotrienols are considered nature's most potent antioxidants, the fact that low-cost **gamma tocopherol with sesame** is more effective is a remarkable finding.

Life Extension has fortified the popular **Gamma E Tocopherol** supplement with standardized **sesame lignans**. Consumers obtain superior benefits at a much **lower cost**.

WORLD'S MOST COMPREHENSIVE VITAMIN E FORMULA!

Each softgel of **Gamma E Tocopherol with Sesame Lignans** provides:

Gamma tocopherol	237 mg
Sesame lignan extract	20 mg
Delta tocopherol	79 mg
Alpha tocopherol	36 mg
Beta tocopherol	3.6 mg

This formula provides potent doses of **gamma tocopherol** along with **sesame lignan extract** to augment the **antioxidant** effects of **gamma tocopherol**.

The retail price for 60 softgels of **Gamma E Tocopherol with Sesame Lignans** is **\$32**. If a member buys four bottles during **Super Sale**, the price is reduced to only **\$19.58 per bottle**.

To order **Gamma E Tocopherol with Sesame Lignans**,
call **1-800-544-4440**
or visit www.lifeextension.com

References

1. *J Natl Cancer Inst.* 2000 Dec;20;92(24):1966-7.
2. *Proc Natl Acad Sci USA.* 1997 Apr 1; 94(7):3217-22.
3. *Atherosclerosis.* 1999 May;144(1):117-22.
4. *J Nutr.* 1992 Dec;122(12):2440-6.
5. *Lipids.* 1995 Nov;30(11):1019-28.



Item # 00759

Antioxidant Vitamins & Cancer: Some scientific evidence suggests that consumption of antioxidant vitamins may reduce the risk of certain forms of cancer. However, the FDA does not endorse this claim because this evidence is limited and not conclusive.



Vital Greens Plant Foods In a Convenient Powder Blend

We try to eat healthfully, but getting enough **plant foods** each day can be a challenge for many people.

The new **Vital Greens Mix** is a blend of *organic* whole plant foods that provide diverse benefits ranging from protecting DNA integrity and maintaining immune function to suppressing vascular risk factors and restoring healthy intestinal flora.

Some of the ingredients in the new **Vital Greens Mix** superfood include:

- **Chia**, a rich source of protein, fiber, vitamins, minerals, and alpha-linolenic acid (an omega-3 precursor). Recent research at the University of Toronto has shown that regular **chia** intake may help maintain healthy **blood pressure** and **C-reactive protein** levels, both of which are well-known markers of cardiovascular disease.¹
- **Green Grasses and Sprouts** to provide a full array of **chlorophyll-rich**, nutrient-dense natural “green” foods that help **detoxify** and maintain **youthful DNA integrity**. In addition, clinical trials with *chlorella* supplements demonstrate that daily use of this green food may *support healthy immunity and wound healing*.^{2,3}
- **Probiotics** to sustain **healthy bacterial balance** in the gut, which may support overall *immune* and *digestive health*.
- **A mushroom blend** combining *maitake*, *shiitake* and *cordyceps* extracts to support **immune health** and provide adaptogenic benefits for balance and resistance.



Item #01098

Each serving provides:

Chia Seed (organic)	3000 mg
Chlorella (organic)	1000 mg
Spirulina (organic)	1000 mg
Acerola (organic)	750 mg
Wheat Grass (organic)	430 mg
Maitake Mushroom Extract (organic)	400 mg
Wheat Sprout (organic)	300 mg
Barley Malt (organic)	300 mg
Probiotic Blend (<i>Lactobacillus acidophilus</i> , <i>Lactobacillus casei</i> , <i>Lactobacillus plantarum</i> , <i>Lactobacillus rhamnosus</i> , <i>Bifidobacterium breve</i> , <i>Bifidobacterium longum</i>)	300 mg
Shiitake Mushroom (organic)	300 mg
Cordyceps Mushroom Extract (organic)	300 mg
Dulse (organic)	250 mg
Beet Juice (organic)	250 mg
Barley Grass (organic)	200 mg
Brown Rice Bran (organic)	200 mg
Cilantro (organic)	200 mg
Suma (organic)	150 mg
Flower Pollen Extract (organic)	120 mg

Life Extension®'s Vital Greens Mix is a great way to start off a meal. It induces some immediate satiety while providing beneficial plant foods that help neutralize mutagenic components of modern diets.

Each scoop of **Life Extension's Vital Greens Mix** provides a daily dose of nutrient-dense foods and herbal extracts. The retail price of a 30-serving jar of this organic superfood is \$48. When a member buys four jars during **Super Sale**, the price is reduced to only **\$29.70 per bottle**.

To order the new **Vital Greens Mix**,
call **1-800-544-4440** or visit
www.LifeExtension.com

REFERENCES:

1. *Diabetes Care*. 2007 Nov;30(11):2804-10.
2. *CMAJ*. 2003 Jul 22;169(2):111-7.
3. *Altern Ther Health Med*. 2001 May-Jun;7(3):79-91.

Activate Your Longevity Genes with New CALORIE RESTRICTION MIMETIC Formula

Scientists have *intensely* focused on finding out **how calorie restriction** so radically extends life span.

In response to **reduced calorie intake**, beneficial "youth" genes are **activated** while detrimental "senescence" genes are **disabled**. These favorable **gene expression** changes play a critical role in the ability of calorie restriction to **slow aging**.

The incredible news is that a select group of nutrients has been discovered that trigger many of the same favorable mechanisms (including more youthful **gene expression** patterns) as **caloric restriction**.

Among the most promising **caloric restriction mimics** and enhancers are **resveratrol**, **pterostilbene**, **quercetin**, and **grape seed polyphenols**, along with **black tea extract**. These nutrients have been shown to generate many of the same effects in the body as caloric restriction, without significant dietary modification.¹⁻⁸ In particular, they help "mimic" caloric restriction's favorable impact on **genes** that influence the aging process.¹⁻⁸

A new **Calorie Restriction Mimetic Formula** includes **resveratrol**, **pterostilbene**, **quercetin**, **grape seed polyphenols**, and **black tea extract** to provide even broader-spectrum gene expression support in one nutritional compound. The suggested dose of two capsules each day of the new **Calorie Restriction Mimetic Formula** provides:

Trans-Resveratrol	250 mg
Trans-Pterostilbene	3 mg
Quercetin	150 mg
Black tea extract	300 mg
Grape seed polyphenols	50 mg

A bottle containing 60 vegetarian capsules of the **new Calorie Restriction Mimetic Formula** retails for \$36. If a member buys four bottles during the **Super Sale**, the cost is reduced to **\$22.28** per bottle.



To order the new **Calorie Restriction Mimetic Formula** call **1-800-544-4440** or visit to www.LifeExtension.com

Note: Those taking other **resveratrol** products may consider switching to this new **Calorie Restriction Mimetic Formula**.

REFERENCES

1. *Cell*. 2006 Dec 15;127(6):1109-22.
2. *Endocrinology*. 2008 Jan;149(1):84-92.
3. *Crit Care Med*. 2004 Oct;32(10):2097-103.
4. *J Agric Food Chem*. 1999 Apr;47(4):1416-21.
5. *Arch Pharm Res*. 2002 Oct;25(5):561-71.
6. *Nutr Cancer*. 1999;35(1):80-6.
7. *Anticancer Agents Med Chem*. 2006 Sep;6(5):389-406.
8. *Nature*. 2006 Nov 16;444(7117):337-42.



Block *Absorption* of Killer Carbohydrates

BY JULIUS GOEPP, MD

If you're an American over 20 years old, you may already suffer from some form of **impaired glucose control**.

According to 2007 data from the National Institutes of Health, 25% of Americans 20 years and older had abnormally high levels of **glucose** in the blood—a **pre-diabetic** state. That number leapt to 35% in individuals 60 and older. Extrapolating from the total US population, nearly **60 million American adults** may now be pre-diabetic.¹

These statistics make it clear that the glut of excess **carbohydrate** calories lurking in the Western diet undermines our best efforts to maintain healthy metabolic function and body weight.

Far too many people have lost their *metabolic capacity* to properly **process** carbohydrates. The result is a looming public health disaster, as **impaired glucose control** initiates a series of **metabolic disorders** that sharply increases disease risk.

To reverse this lethal epidemic of impaired glucose control, aging individuals often require more **aggressive** measures than merely *limiting* carbohydrate intake.

Fortunately, researchers have discovered proven methods to limit one's exposure to carbohydrate calories without dramatic changes in diet. It centers on a group of natural compounds called **enzyme inhibitors**.

These interventions *block* the action of **sucrase**, **amylase**, and **glucosidase**, digestive enzymes responsible for the breakdown of various *forms* of carbohydrate. Instead of absorbing and converting these sugars into higher than desired blood glucose levels, you may be able to safely eliminate many of these excess carbohydrate calories.

These natural “carb-blockers” provide a *practical* approach to help lower glucose levels, improve age-related markers of health, and regain glycemic control.² > >

Combating the Scourge of Sucrose

Starting in early childhood, most Americans are exposed to far too much *refined* carbohydrates in the form of sucrose (table sugar). It is virtually omnipresent in processed foods. It also happens to be an enemy to human health and an age-accelerator.

Many individuals do not even knowingly ingest it. According to a 2009 study in the *Journal of Nutrition*, only **3%** of dietary sucrose is deliberately *added* by consumers; **82%** is added by manufacturers.³ And while the World Health Organization (WHO) recommends that sucrose comprise only **10%** of total energy intake, many people substantially exceed this level from the earliest years of life—usually at the expense of key nutrients.^{3,4}

Manufacturers know that most people will consistently choose sweetened over unsweetened foods and beverages.⁵ People will often *continue* to do so at the expense of their own health. After years of chronic exposure to sucrose, physiological processes not unlike addiction take hold. Rapid absorption of sugars from a high-sucrose meal triggers a dangerous sequence of unfavorable hormonal and metabolic alterations that promote still greater consumption, especially in overweight individuals.^{6,7} The result is the dangerously high incidence of metabolic disease we see today—obesity, type 2 diabetes, and metabolic syndrome.⁸⁻¹⁰

This alarming trend has led to a set of novel, evidence-based solutions. Aging individuals now have practical means at their disposal to slow or reverse

the long-term consequences of chronic sucrose overexposure.

Sucrose is composed of two simple sugar molecules, glucose and fructose. It is poorly *absorbed* in the intestine in this form. In order to be utilized, it must first be broken down by the digestive enzyme **sucrase**. Blocking the enzymatic action of *sucrase* therefore limits uptake of *sucrose*.

L-arabinose

Researchers have identified a potent *sucrase* inhibitor called **L-arabinose**. Although it is a simple plant sugar, L-arabinose is indigestible and cannot be absorbed into the blood. Instead it remains in the digestive tract and is eventually excreted.^{11,12} By blocking metabolism of sucrose, L-arabinose *inhibits* the spike in blood sugar and fat synthesis that would otherwise follow a sugar-rich meal.¹² In animal models, L-arabinose virtually eliminates the rise in blood sugar following administration of sucrose, with blood glucose levels rising only **2%** higher than in control animals that did not receive sucrose. L-arabinose did not exert *any* effect on serum glucose levels in control animals that did not receive sucrose.¹³

L-arabinose has been proven safe in both short- and long-term studies, and may contribute to lowered levels of glycosylated hemoglobin (hemoglobin A1C),¹⁴ a measure of chronic exposure to sugar in the blood. A study combining L-arabinose and **white bean extract** (see next page) not only smoothed out postprandial glucose spikes and reduced insulin levels—it *lowered* systolic blood pressure.¹⁴



Chromium

The trace element **chromium** is another potent aid in the management of healthy blood sugar levels. Scientists have known for years that chromium aids in the management of type 2 diabetes, where it not only helps control blood glucose but also reduces **total cholesterol levels**.¹⁵⁻¹⁷ The combination of chromium with L-arabinose attacks multiple targets to prevent sucrose-induced blood sugar elevation and reduce sugar calorie exposure.

Complex Carbohydrate Enzyme Inhibitors

White Bean Extract

Extracts from the common white kidney bean, *Phaseolus vulgaris*, are powerful blockers of the enzyme *alpha-amylase*, which is secreted by the pancreas.^{18,19} Alpha-amylase breaks down long-chain, complex starch molecules into simple sugars and short-chain *oligosaccharides* for absorption in the small intestine. Blocking *alpha-amylase* inhibits the metabolism of starches and *slows* the rate at which free sugars are absorbed.

White bean extract shows enormous potential for preventing the blood sugar and insulin spikes that are associated with many chronic health disorders.¹³ Slowing starch digestion also prolongs the amount of time it takes for the stomach to empty its contents, further reducing the amount of carbohydrate calories released at any one time into the intestine.²⁰

White bean extracts operate along numerous overlapping pathways in multiple, related physiological systems. Laboratory research shows that supplementation with *white bean extract* promotes weight loss in obese animals, with dramatic reduction in fat accumulation without loss of muscle mass.^{21,22} Plasma *insulin* levels also dropped substantially following a high-carbohydrate meal including white bean extract in pre-clinical studies, reflecting a much more gradual rise in blood sugar levels.²²

Amylase inhibition with *white bean extract* has proven particularly effective in reducing glycemia (sugar load in the blood) in studies on diabetic animals. Supplementation in diabetic rats not only substantially lowered mean blood sugar levels; it also reduced the animals' total food and water intake (water intake is increased in untreated diabetes because of the amount lost in sugar-laden urine).²³ White bean extract has also been shown to boost levels of an intestinal hormone called *cholecystokinin* (CCK), which produces the sensation of satiety following a meal and reduces the urge to continue eating.²⁴ Research further indicates it can restore overactive intestinal sugar-digesting enzymes



Irvingia

- Roughly 1 in 5 Americans are pre-diabetic, a result of excess carbohydrate consumption.
- The digestive enzymes *sucrase*, *amylase*, and *glucosidase* are primarily responsible for enabling carbohydrate absorption into the blood.
- L-arabinose—a natural but largely indigestible sugar—blocks *sucrase* activity, preventing the sugar *sucrose* from entering the bloodstream.
- Extracts of white bean, seaweed, and Irvingia block the breakdown of starch in the intestine, preventing their calories from being absorbed.
- Green tea extract helps your body burn additional calories at rest.
- Collectively, these natural compounds may dramatically inhibit total carbohydrate intake, limit postprandial blood sugar and insulin spikes, and generate weight loss.



to nearly normal levels, further reducing the sugar load.²³

White bean extract has yielded equally compelling results in human studies. It has been shown to diminish the effects of high-glycemic index foods (like white bread) that are notorious for producing sharp, potentially dangerous postprandial blood sugar spikes, helping to alleviate metabolic burden throughout the body.²⁵ In obese but otherwise healthy patients, supplementation with white bean extract produces weight loss *more than twice as fast* as in controls.²⁶ Even in people who are only slightly overweight, *white bean extract* reduces body weight, body mass index (BMI), fat mass, fat tissue thickness, and waist circumference, while maintaining lean body mass²⁷—key elements that are linked with metabolic syndrome and cardiovascular disease risk.

White bean extract is highly effective even in aging individuals whose diets contain the highest amounts of carbohydrates.²⁸ By blocking *amylase* and slowing starch digestion, it increases the amount of intact carbohydrates reaching the lower bowel, where they are consumed by **beneficial intestinal bacteria**.¹⁹ It has also been shown to induce a **three-fold reduction** in serum triglycerides compared to controls.²⁶

White bean extract has an excellent safety profile, even at high doses.²⁹ As with all the carbohydrate inhibitors reviewed here, it does not reduce blood sugar to *below* normal levels—an important

feature for diabetics in whom large swings in glucose levels are especially dangerous.¹³

It should be noted that ingestion of white bean extract may result in minor intestinal protein losses, requiring additional protein intake.³⁰

Irvingia

Extract of the wild African mango *Irvingia gabonensis* can produce weight loss by inhibiting calorie absorption and storage through multiple mechanisms. It also exerts potent anti-diabetic and anti-obesity effects.³¹ Studies from Africa in the mid-1980's showed that Irvingia extract normalized blood sugar in diabetic patients while increasing the activity of enzymes involved in cellular energy metabolism.³² Irvingia extract also lowers blood levels of dangerous LDL and VLDL in diabetics, while increasing beneficial HDL levels.³³ It mobilizes liver enzyme activity in favor of glucose storage and *away* from its release into the bloodstream, thereby improving glucose control.³¹

Irvingia works in part by inhibiting the amylase enzyme, and thus provides support for those attempting to restrict their total carbohydrate exposure and lose weight.³¹ In one double-blind, placebo-controlled

White bean extract... has been shown to diminish the effects of **high-glycemic index foods** that produce sharp, potentially dangerous postprandial blood sugar spikes.

study of obese but otherwise healthy adults, one month of supplementation produced a **5.3%** body weight loss in supplemented patients, compared with only a **1.3%** loss in the control group.³⁴ These individuals also saw significant improvement in their lipid profiles. Additional studies confirm these findings, demonstrating significant reductions in body fat content, waist circumference, blood sugar levels, and markers of fat tissue regulation.^{35,36}

In addition to blocking amylase activity, *Irvingia* extract acts directly on fat cells to reduce lipid formation and storage.^{31,37} Its operation is multi-modal—down-regulating genes that promote fat production—while upregulating factors that suppress it, including *adiponectin*,^{36,37} a glucose-regulating hormone that lowers fat-mediated inflammation and enhances insulin sensitivity.³⁸ Adiponectin levels correlate with a reduction in risk for metabolic syndrome.^{39,40}

Seaweed Extracts

Extracts from several seaweed species (actually the complex algae known as kelp) are potent inhibitors of *amylase* and another digestive enzyme called *alpha-glucosidase*. They have proven to be cost-effective means of preventing the progression of diabetes in pre-clinical models.⁴¹ In particular, extracts of *Fucus vesiculosus* and *Ascophyllum nodosum* are known to help lower blood glucose in normal and diabetic animals.^{42,43}

A recent detailed study of a seaweed compound containing extracts from both *Fucus* and *Ascophyllum* offers insights into their mechanisms and benefits. The combination was given to laboratory rats prior to a meal with a high glycemic index.⁴⁴ Such meals typically produce a rapid rise in both glucose and insulin levels in the blood, with an equally rapid drop within 60 minutes. The *glucose* spike contributes to excessive formation of dangerous *advanced glycation end-products* (AGEs). The *insulin* spike contributes to paradoxically low blood sugar at about 90 minutes, which can create a sensation of hunger, prompting excess food consumption.

The seaweed combination cut postprandial sugar spikes by **90%** compared with untreated animals.⁴⁴ That in turn reduced the insulin spike by **40%**, which completely eliminated the period of low blood sugar that followed the meal in untreated rats.

In response to this seaweed combination, postprandial glucose and insulin profiles were modified to levels resembling meals with a much lower glycemic load. Blood sugar rose more slowly, achieved a shorter and more modest peak level, and then declined more gradually without ever “bottoming out” at an abnormally low level.

Lethal Dangers of Excess Insulin

As humans age and/or ingest excess calories, a phenomenon known as “insulin resistance” develops whereby cell membranes become less sensitive to the effects of insulin.

As glucose levels increase in the blood (instead of being taken up by cells), the pancreas tries to compensate by secreting higher levels of insulin. Too often the end result is a vicious cycle of **hyperinsulinemia** (excess blood insulin) and **hyperglycemia** (excess blood glucose) as aging cells become less efficient in removing glucose from the blood.

Too much **insulin** damages the body in the following ways:

- Converts excess glucose into fat stores⁴⁶
- Floods the blood with triglycerides and C-reactive protein^{46,47}
- Drives down levels of healthy HDL⁴⁶
- Destabilizes the body's sodium balance, increasing blood pressure^{48,49}
- May contribute to kidney damage⁵⁰
- Ravages the vascular system^{51,52}
- Has been associated with increased risk of certain cancers^{53,54}

Glucose and insulin levels drop in response to reduced calorie intake, exercise, and/or use of compounds before meals that inhibit glucosidase, amylase, and sucrase enzymes in the digestive tract.

A recently completed human clinical trial produced similarly promising results. In this randomized, cross-over, placebo-controlled, double-blind study conducted at Laval University, 23 healthy volunteers consumed **500 mg** of a *Fucus-Ascophyllum* combination along with a high-glycemic index meal of white bread. The seaweed combination produced a **44%** reduction in the glycemic response that normally follows ingestion of such a meal. The seaweed combination also produced a **22%** reduction in the initial insulin production following the meal, and an overall **5.9%** reduction in the area under the curve of the insulin response. The study results will be presented at the *Experimental Biology* meeting in Anaheim in April, 2010.⁴⁵

The findings suggest that in humans, the *Fucus-Ascophyllum* compound may lead to earlier satiety, longer intervals between meals, fewer urges to snack, and lower total calorie intake.



Green Tea Extract

Raising basal metabolic rate is another effective mechanism for offsetting excess carbohydrate intake. Green tea extract boosts the “resting” metabolism by inhibiting an enzyme called catechol-O-methyl transferase or *COMT* that breaks down *noradrenaline*, an adrenaline-like hormone that sustains energy production.⁵⁵ The resulting higher levels of metabolic activity help to burn off excess calories.⁵⁶

In a large clinical trial, a patented green tea phyto-some extract produced exceptional weight loss in obese individuals. Supplemented subjects lost almost **31 lbs** over 3 months, while controls lost just 11 pounds!⁵⁷ Both groups followed a low calorie diet. Multiple studies of overweight and obese adults indicate that green tea extracts can reduce abdominal fat as well as total cholesterol, LDL, and fasting triglyceride levels.^{58,59}

Green tea provides another benefit in helping to alleviate the metabolic burden imposed by excess calorie ingestion. It has been shown to inhibit the *lipase* digestive enzyme that breaks down dietary fats for absorption into the blood.⁶⁰

Summary

Roughly 1 in 5 Americans are pre-diabetic, a result of excess calorie consumption and normal aging. *Sucrase*, *amylase*, and *glucosidase* are digestive enzymes that break down carbohydrates, and *lipase* is a digestive enzyme that breaks down fat, facilitating absorption of excess calories into the blood.

As humans age, the impact of **chronic caloric overload** enabled by these digestive enzymes can lead to an array of life-threatening conditions ranging from high blood sugar and insulin to type 2 diabetes, obesity, and metabolic syndrome.

Natural compounds have been shown to effectively *inhibit* these digestive enzymes and *impede* the *absorption* of excess carbohydrate. **L-arabinose** neutralizes **sucrase**, reducing uptake of sugar (as sucrose) into the blood. Extracts of **white bean**, *Irvingia gabonensis*, and certain **seaweeds** block **amylase** and **glucosidase** activity, further reducing the number of ingested carbohydrate calories that are absorbed. Green or black **tea** extract can help reduce the activity of **lipase**, a digestive enzyme that helps break down fat in the gastrointestinal tract.

These natural compounds have been shown to effectively lower excess calorie **absorption** while reducing postprandial (post-meal) blood sugar and blood fat (triglyceride) spikes. •

If you have any questions on the scientific content of this article, please call a Life Extension® Health Advisor at 1-866-864-3027.

References

1. Available at: <http://diabetes.niddk.nih.gov/DM/PUBS/statistics/#youngpeople>. Accessed November 30, 2009.
2. Bischoff H. Pharmacology of alpha-glucosidase inhibition. *Eur J Clin Invest.* 1994 Aug;24 Suppl 3:3-10.

3. Erkkola M, Kronberg-Kippila C, Kyttala P, et al. Sucrose in the diet of 3-year-old Finnish children: sources, determinants and impact on food and nutrient intake. *Br J Nutr.* 2009 Apr;101(8):1209-17.
4. Ruottinen S, Niinikoski H, Lagstrom H, et al. High sucrose intake is associated with poor quality of diet and growth between 13 months and 9 years of age: the special Turku Coronary Risk Factor Intervention Project. *Pediatrics.* 2008 Jun;121(6):e1676-85.
5. Vorster HH, van Tonder E, Kotze JP, Walker AR. Effects of graded sucrose additions on taste preference, acceptability, glycemic index, and insulin response to butter beans. *Am J Clin Nutr.* 1987 Mar;45(3):575-9.
6. Ludwig DS, Majzoub JA, Al-Zahrani A, Dallal GE, Blanco I, Roberts SB. High glycemic index foods, overeating, and obesity. *Pediatrics.* 1999 Mar;103(3):E26.
7. Van Wymelbeke V, Beridot-Therond ME, de La Gueronniere V, Fantino M. Influence of repeated consumption of beverages containing sucrose or intense sweeteners on food intake. *Eur J Clin Nutr.* 2004 Jan;58(1):154-61.
8. Striegel-Moore RH, Thompson D, Affenito SG, et al. Correlates of beverage intake in adolescent girls: the National Heart, Lung, and Blood Institute Growth and Health Study. *J Pediatr.* 2006 Feb;148(2):183-7.
9. Palmer JR, Boggs DA, Krishnan S, Hu FB, Singer M, Rosenberg L. Sugar-sweetened beverages and incidence of type 2 diabetes mellitus in African American women. *Arch Intern Med.* 2008 Jul 28;168(14):1487-92.
10. Culling KS, Neil HA, Gilbert M, Frayn KN. Effects of short-term low- and high-carbohydrate diets on postprandial metabolism in non-diabetic and diabetic subjects. *Nutr Metab Cardiovasc Dis.* 2009 Jun;19(5):345-51.
11. Seri K, Sanai K, Matsuo N, Kawakubo K, Xue C, Inoue S. L-arabinose selectively inhibits intestinal sucrase in an uncompetitive manner and suppresses glycemic response after sucrose ingestion in animals. *Metabolism.* 1996 Nov;45(11):1368-74.
12. Osaki S, Kimura T, Sugimoto T, Hizukuri S, Iritani N. L-arabinose feeding prevents increases due to dietary sucrose in lipogenic enzymes and triacylglycerol levels in rats. *J Nutr.* 2001 Mar;131(3):796-9.
13. Preuss HG, Echard B, Bagchi D, Stohs S. Inhibition by natural dietary substances of gastrointestinal absorption of starch and sucrose in rats and pigs: 1. Acute studies. *Int J Med Sci.* 2007;4(4):196-202.
14. Preuss HG, Echard B, Bagchi D, Stohs S. Inhibition by natural dietary substances of gastrointestinal absorption of starch and sucrose in rats 2. Subchronic studies. *Int J Med Sci.* 2007;4(4):209-15.
15. Terpilowska S, Zaporowska H. The role of chromium in cell biology and medicine. *Przegl Lek.* 2004;61 Suppl 3:51-4.
16. Broadhurst CL, Domenico P. Clinical studies on chromium picolinate supplementation in diabetes mellitus—a review. *Diabetes Technol Ther.* Dec 2006 Dec;8(6):677-87.
17. Balk EM, Tatsioni A, Lichtenstein AH, Lau J, Pittas AG. Effect of chromium supplementation on glucose metabolism and lipids: a systematic review of randomized controlled trials. *Diabetes Care.* 2007 Aug;30(8):2154-63.
18. Mosca M, Boniglia C, Carratu B, Giammarioli S, Nera V, Sanzini E. Determination of alpha-amylase inhibitor activity of phaseolamin from kidney bean (*Phaseolus vulgaris*) in dietary supplements by HPAEC-PAD. *Anal Chim Acta.* 2008 Jun 9;617(1-2):192-5.
19. Obiro WC, Zhang T, Jiang B. The nutraceutical role of the *Phaseolus vulgaris* alpha-amylase inhibitor. *Br J Nutr.* 2008 Jul;100(1):1-12.
20. Layer P, Zinsmeister AR, DiMagno EP. Effects of decreasing intraluminal amylase activity on starch digestion and postprandial gastrointestinal function in humans. *Gastroenterology.* 1986 Jul;91(1):41-8.
21. Santoro LG, Grant G, Pusztai A. Effects of short-term feeding of rats with a highly purified phaseolin preparation. *Plant Foods Hum Nutr.* 1997;51(1):61-70.
22. Pusztai A, Grant G, Buchan WC, Bardocz S, de Carvalho AF, Ewen SW. Lipid accumulation in obese Zucker rats is reduced by inclusion of raw kidney bean (*Phaseolus vulgaris*) in the diet. *Br J Nutr.* 1998 Feb;79(2):213-21.
23. Tormo MA, Gil-Exijo I, Romero de Tejada A, Campillo JE. White bean amylase inhibitor administered orally reduces glycaemia in type 2 diabetic rats. *Br J Nutr.* 2006 Sep;96(3):539-44.
24. Fantini N, Cabras C, Lobina C, et al. Reducing effect of a *Phaseolus vulgaris* dry extract on food intake, body weight, and glycemia in rats. *J Agric Food Chem.* 2009 Oct 14;57(19):9316-23.
25. Udani JK, Singh BB, Barrett ML, Preuss HG. Lowering the glycemic index of white bread using a white bean extract. *Nutr J.* 2009;8:52.
26. Udani J, Hardy M, Madsen DC. Blocking carbohydrate absorption and weight loss: a clinical trial using Phase 2 brand proprietary fractionated white bean extract. *Altern Med Rev.* 2004 Mar;9(1):63-9.
27. Celleno L, Tolaini MV, D'Amore A, Perricone NV, Preuss HG. A Dietary supplement containing standardized *Phaseolus vulgaris* extract influences body composition of overweight men and women. *Int J Med Sci.* 2007;4(1):45-52.
28. Udani J, Singh BB. Blocking carbohydrate absorption and weight loss: a clinical trial using a proprietary fractionated white bean extract. *Altern Ther Health Med.* 2007 Jul-Aug;13(4):32-7.
29. Chokshi D. Toxicity studies of Blockal, a dietary supplement containing Phase 2 Starch Neutralizer (Phase 2), a standardized extract of the common white kidney bean (*Phaseolus vulgaris*). *Int J Toxicol.* 2006 Sep-Oct;25(5):361-71.
30. Deglaire A, Moughan PJ, Bos C, Tome D. Commercial *Phaseolus vulgaris* extract (starch stopper) increases ileal endogenous amino acid and crude protein losses in the growing rat. *J Agric Food Chem.* 2006 Jul 12;54(14):5197-202.
31. Omoruyi F, Adamson I. Digestive and hepatic enzymes in streptozotocin-induced diabetic rats fed supplements of dikanut (*Irvingia gabonensis*) and cellulose. *Ann Nutr Metab.* 1993;37(1):14-23.
32. Adamson I, Okafor C, Abu-Bakare A. Erythrocyte membrane ATPases in diabetes: effect of dikanut (*Irvingia gabonensis*). *Enzyme.* 1986;36(3):212-5.
33. Adamson I, Okafor C, Abu-Bakare A. A supplement of Dikanut (*Irvingia gabonensis*) improves treatment of type II diabetics. *West Afr J Med.* 1990 Apr-Jun;9(2):108-15.
34. Ngondi JL, Oben JE, Minka SR. The effect of *Irvingia gabonensis* seeds on body weight and blood lipids of obese subjects in Cameroon. *Lipids Health Dis.* 2005;4:12.



35. Oben JE, Ngondi JL, Momo CN, Agbor GA, Sobgui CS. The use of a *Cissus quadrangularis*/*Irvingia gabonensis* combination in the management of weight loss: a double-blind placebo-controlled study. *Lipids Health Dis.* 2008;7:12.

36. Ngondi JL, Etoundi BC, Nyangono CB, Mboufong CM, Oben JE. IGOB131, a novel seed extract of the West African plant *Irvingia gabonensis*, significantly reduces body weight and improves metabolic parameters in overweight humans in a randomized double-blind placebo controlled investigation. *Lipids Health Dis.* 2009;8:7.

37. Oben JE, Ngondi JL, Blum K. Inhibition of *Irvingia gabonensis* seed extract (OB131) on adipogenesis as mediated via down regulation of the PPARgamma and leptin genes and up-regulation of the adiponectin gene. *Lipids Health Dis.* 2008;7:44.

38. Cetinalp-Demircan P, Bekpinar S, Gurdol F, Orhan Y. Adiponectin is a link among inflammation, insulin resistance, and high-density lipoprotein cholesterol but is not associated with paraoxonase activity in premenopausal women. *J Clin Hypertens (Greenwich).* 2009 Nov;11(11):672-7.

39. Fontana L, Klein S, Holloszy JO. Effects of long-term calorie restriction and endurance exercise on glucose tolerance, insulin action, and adipokine production. *Age (Dordr).* 2009 Nov 11.

40. Stenholm S, Koster A, Alley DE, et al. Adipocytokines and the metabolic syndrome among older persons with and without obesity - the InCHIANTI Study. *Clin Endocrinol (Oxf).* 2009 Oct 31.

41. Iwai K. Antidiabetic and antioxidant effects of polyphenols in brown alga *Ecklonia stolonifera* in genetically diabetic KK-A(y) mice. *Plant Foods Hum Nutr.* 2008 Dec;63(4):163-9.

42. Lamela M, Anca J, Villar R, Otero J, Calleja JM. Hypoglycemic activity of several seaweed extracts. *J Ethnopharmacol.* 1989 Nov;27(1-2):35-43.

43. Zhang J, Tiller C, Shen J, et al. Antidiabetic properties of polysaccharide- and polyphenolic-enriched fractions from the brown seaweed *Ascophyllum nodosum*. *Can J Physiol Pharmacol.* 2007 Nov;85(11):1116-23.

44. InnoVactiv, Inc. Data on file.

45. Available at: <http://www.naturalproductsinsider.com/news/2009/12/insea2-reduces-glycemic-response.aspx#>. Accessed December 2, 2009.

46. Stulc T, Cecka R. Lipid-lowering treatment in metabolic syndrome. *Vnitr Lek.* 2009 Jul-Aug;55(7-8):626-30.

47. Andel M, Polák J, Kraml P, Dlouhý P, Stich V. Chronic mild inflammation links obesity, metabolic syndrome, atherosclerosis and diabetes. *Vnitr Lek.* 2009 Jul-Aug;55(7-8):659-65.

48. Hano T, Nishio I. Treatment of hypertension in the patients with obesity. *Nippon Rinsho.* 2001 May;59(5):973-7.

49. Kopf D, Muhlen I, Kroning G, Sendzik I, Huschke B, Lehnert H. Insulin sensitivity and sodium excretion in normotensive off-spring and hypertensive patients. *Metabolism.* 2001 Aug;50(8):929-35.

50. Noda M, Matsuo T, Nagano-Tsuge H, et al. Involvement of angiotensin II in progression of renal injury in rats with genetic non-insulin-dependent diabetes mellitus (Wistar fatty rats). *Jpn J Pharmacol.* 2001 Apr;85(4):416-22.

51. Hegele RA. Premature atherosclerosis associated with monogenic insulin resistance. *Circulation.* 2001 May 8;103(18):2225-9.

52. Katz AS, Goff DC, Feldman SR. Acanthosis nigricans in obese patients: Presentations and implications for prevention of atherosclerotic vascular disease. *Dermatol Online J.* 2000 Sep;6(1):1.

53. Kaaks R. Plasma insulin, IGF-I and breast cancer. *Gynecol Obstet Fertil.* 2001 Mar;29(3):185-91.

54. Nilsen TI, Vatten LJ. Prospective study of colorectal cancer risk and physical activity, diabetes, blood glucose and BMI: exploring the hyperinsulinaemia hypothesis. *Br J Cancer.* 2001 Feb 2;84(3):417-22.

55. Dulloo AG, Seydoux J, Girardier L, Chantre P, Vandermander J. Green tea and thermogenesis: interactions between catechin-polyphenols, caffeine and sympathetic activity. *Int J Obes Relat Metab Disord.* 2000 Feb;24(2):252-8.

56. Diepvens K, Westerterp K R, Westerterp-Plantenga MS. Obesity and thermogenesis related to the consumption of caffeine, ephedrine, capsaicin, and green tea. *Am J Physiol Regul Integr Comp Physiol.* 2007 Jan;292(1):R77-85.

57. Di Pierro F, Menghi AB, Barreca A, Lucarelli M, Calandrelli A. Greenselect Phytosome as an adjunct to a low-calorie diet for treatment of obesity: a clinical trial. *Altern Med Rev.* 2009 Jun;14(2):154-60.

58. Maki KC, Reeves MS, Farmer M, et al. Green tea catechin consumption enhances exercise-induced abdominal fat loss in overweight and obese adults. *J Nutr.* 2009 Feb;139(2):264-70.

59. Tsai CH, Chiu WC, Yang NC, Ouyang CM, Yen YH. A novel green tea meal replacement formula for weight loss among obese individuals: a randomized controlled clinica *J Nutr Biochem.* 2000 Jan;11(1):45-51.

60. Juhel C, Armand M, Pafumi Y, Rosier C, Vandermander J, Lairon D. Green tea extract (AR25) inhibits lipolysis of triglycerides in gastric and duodenal medium in vitro. *Int J Food Sci Nutr.* 2009 Sep 7:1-9.



BONE RESTORE

for Lifelong Bone Health

Maintaining strong, healthy bones is a cornerstone to successful aging. Unfortunately, even supplement users often fail to consume enough **calcium**, **vitamin D**, and other nutrients shown to help maintain healthy bones.

New research on **vitamin D** has led an increasing number of experts to advise men and women to consume **1000 IU** (and higher) of vitamin D each day to help maintain bone density.

The daily dose of **Bone Restore** provides **1200 elemental milligrams** of highly absorbable calcium, plus **1000 IU** of **vitamin D3**.

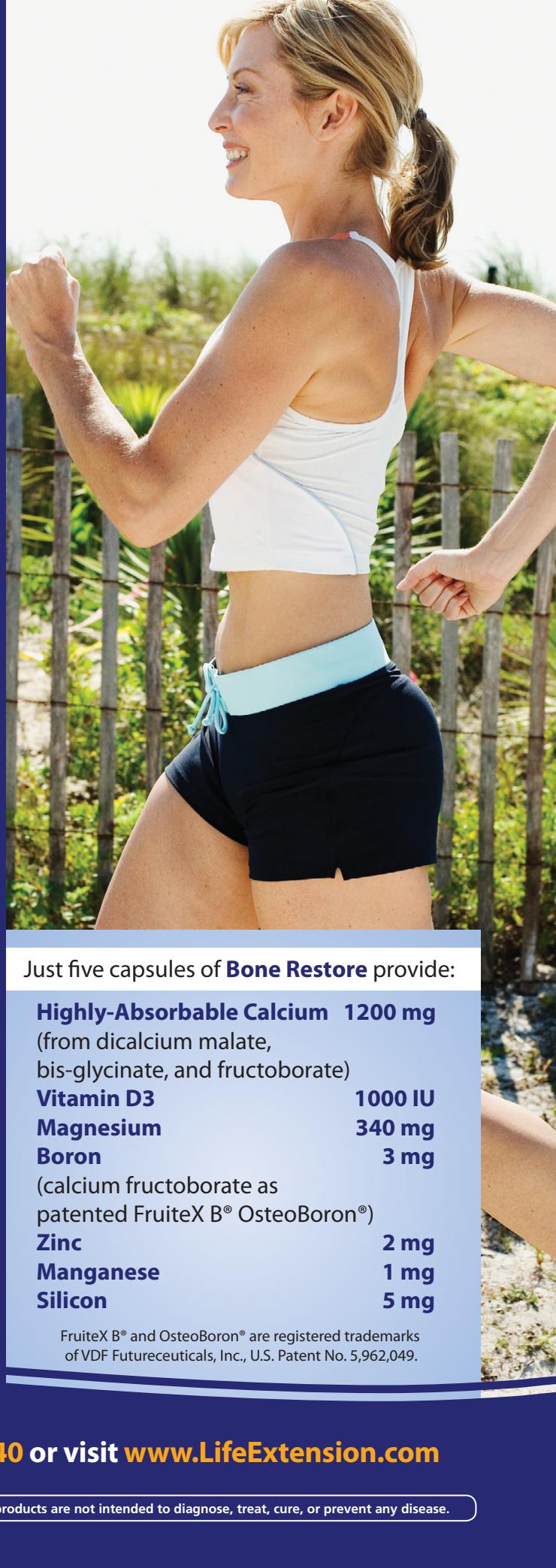
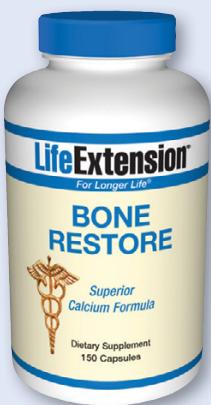
Bone Restore also contains ample **magnesium**, along with nutrients that enable calcium and magnesium to be incorporated in the bone matrix.

In addition, **Bone Restore** delivers a patented form of **boron** that is identical to natural plant forms found in food. Considered more bioavailable than other forms of boron, **FruiteX B® OsteoBoron®** supports healthy joints and bone.

Bone Restore comes in fast-release capsules, making the minerals and other nutrients immediately available for absorption.

The retail price for 150 capsules of **Bone Restore** is \$22.50. If a member buys four bottles during **Super Sale**, the price is reduced to just **\$13.16** per bottle.

Item #00811



Just five capsules of **Bone Restore** provide:

Highly-Absorbable Calcium	1200 mg
(from dicalcium malate, bis-glycinate, and fructoborate)	
Vitamin D3	1000 IU
Magnesium	340 mg
Boron	3 mg
(calcium fructoborate as patented FruiteX B® OsteoBoron®)	
Zinc	2 mg
Manganese	1 mg
Silicon	5 mg

FruiteX B® and OsteoBoron® are registered trademarks of VDF Futureceuticals, Inc., U.S. Patent No. 5,962,049.

To order **Bone Restore**, call **1-800-544-4440** or visit www.LifeExtension.com

A Multi-Faceted Approach to HEALTHY BODY Weight

It's not your fault! Public health agencies blame gluttonous behavior and lack of physical activity as the sole reasons for today's **obesity epidemic**. Ignored are a plethora of age-related **metabolic** changes that predispose us to weight gain, even when we try to cut back on caloric intake.

The good news is that you don't have to do it alone. Scientists have identified *natural compounds* that function via *multiple* mechanisms to combat the underlying factors involved in excess accumulation of body fat.

New Phase 3™ Sucrase Modulator

A recent study in the journal *Science* concluded that in primates a **calorie-restricted diet** can delay disease onset and age-related mortality.¹ While *Life Extension®* readers understand the benefits of calorie restriction, that doesn't make it any easier to cut back on one of the major calorie culprits in the American diet: *refined sugar*.

Sucrase is an enzyme that breaks down sucrose in the digestive tract for *absorption* into the bloodstream. **Phase 3™** is a new, patented compound that functions as a beneficial modulator of sucrase. By delaying the rapid absorption of sucrose, **Phase 3™** can help support the healthy release of insulin and sugar into the bloodstream in response to sucrose ingestion that so readily converts to body fat. **Phase 3™** contains a natural substance found in plants called **L-Arabinose** that provides an important new weapon in the battle to reduce the number of absorbed calories from sugar.

Green Tea Phytosome Cellular Energy Enhancer

The ability of **green tea extracts** to inhibit the breakdown and absorption of dietary fat has been the subject of research studies across the globe. Mounting scientific research has documented the **metabolic effects** of **green tea** polyphenols. A proprietary **phytosome complex** has shown an ability to increase the absorption of green tea polyphenols into the bloodstream better than conventional tea extracts — resulting in peak plasma levels of the critical green tea polyphenol *epigallocatechin-3-gallate*.²

Participants in a human clinical trial showed an average weight loss of **30 pounds** and a **10%** reduction in waist circumference in **90 days** when 300 mg/day of this new **green tea phytosome** was taken in conjunction with a reduced-calorie diet.² The *placebo* group that followed the same reduced-calorie diet lost only **9.9 pounds** and only **5%** of their waist size.



The Alpha-Amylase Enzyme

Aging reduces our ability to utilize the **carbohydrates** (and fats) that constitute what most would consider part of a healthy diet. The result is that as we grow older, our bloodstreams become chronically *bloated* with **glucose** and **triglycerides** in direct contrast to the youthful metabolic profile we have in our younger years. Emerging scientific research suggests that the alpha-amylase enzyme plays an undesirable role in the digestion of dietary carbohydrates and subsequent absorption of calories from starch and sugar.

A natural bean extract (*Phaseolus vulgaris*) moderates *alpha-amylase* activity. In a human trial in which all overweight participants were placed on a 2,000–2,200-calorie, carbohydrate-rich diet, those taking ***Phaseolus vulgaris*** lost **6.5 pounds** and **1.2 inches** in waist size in only **30 days** compared with **0.8 pounds** and **0.2 inches** in the placebo group.³

The Alpha-Glucosidase Enzyme

Another intestinal enzyme that enables carbohydrate absorption is **alpha-glucosidase**. A patented seaweed extract (**InSea™**) has demonstrated the ability to help maintain healthy levels of both *alpha-glucosidase* and *alpha-amylase*. When given to laboratory animals, this seaweed extract reduced after-meal (postprandial) **glucose** elevations by up to **90%** compared with non-supplemented animals.⁴

Remember, young healthy individuals rapidly convert ingested fats-sugars into energy. Age-related changes decrease our **metabolic capacity** to efficiently utilize dietary fats-sugars. It is thus paramount for aging people to reduce their *absorption of excess* calories. By taking nutrients before each meal that reduce the activity of carbohydrate-fat digesting enzymes, the calorie burden is significantly reduced.

To order Optimized Irvingia with Phase 3™ Calorie Control Complex, call toll-free 1-800-544-4440.

Leptin Sensitivity

Fat cells (adipocytes) secrete a hormone called *leptin* that tells our brain we have eaten enough. *Leptin* can also facilitate the breakdown of stored *triglycerides* in our adipocytes via the process of *lipolysis*. Heavy individuals have startlingly high blood levels of leptin, indicating that their cells have become *resistant* to the *leptin* that is supposed to prevent them from putting on so many fat pounds.

An extract from a West African food called *Irvingia gabonensis* has been shown to help support *leptin sensitivity* in overweight people. In a recently published study, *Irvingia* demonstrated beneficial effects upon leptin blood levels, followed by weight loss and inches off the waistline.⁵ In addition to supporting healthy *leptin sensitivity*, *Irvingia* has demonstrated the following beneficial effects on key aspects of metabolism:

- **Glycerol-3-phosphate dehydrogenase** is an enzyme involved in the complex biochemical process that converts ingested starch and sugar calories to stored body fat. *Irvingia* has been shown in studies involving fat cells to reduce the activity of *glycerol-3-phosphate dehydrogenase*,⁶ which may help reduce the impact of starch and sugar calories on body fat.
- Scientific research on fat cells suggests that *Irvingia* has **alpha-amylase-inhibiting** properties⁷ (like *InSea*^{2™} and *Phaseolus vulgaris*), which can help support a *slowing* of the rate of carbohydrate absorption from the intestines and a reduction of the caloric impact of starchy and sugary foods.⁸
- **Adiponectin** is a hormone involved in helping to maintain insulin sensitivity on the membranes of energy-producing cells. Big fat cells produce *less* adiponectin, and overweight people need to be especially concerned about maintaining healthy levels of adiponectin to support insulin sensitivity and metabolic fitness. Scientific data suggests that *Irvingia* helps support healthy adiponectin levels.⁶

The New Optimized *Irvingia* with Phase 3™ Calorie Control Complex

In reviewing the remarkable effects demonstrated by these *natural compounds*, one might think that any one of them might be a solution to their weight problem. The reality is that aging individuals often fall victim to *many molecular factors* that can sabotage the best weight-loss programs.

The new **Optimized *Irvingia*** formula provides a combination of nutrients that combat *age-related* fat accumulation via the following eight distinct mechanisms:

1. Delaying *digestion and absorption* of sucrose.
2. Enhancing *resting energy expenditure* at the cellular level.
3. Slowing the absorption of *dietary fat* from the intestines.
4. Moderating *alpha-amylase enzyme activity* to *reduce carbohydrate absorption* in the bloodstream.
5. Reducing *alpha-glucosidase enzyme activity* to *further slow the absorption* of starches and sugars into the bloodstream.
6. Supporting *leptin sensitivity* to reduce hunger and stimulate lipolysis.
7. Supporting youthful levels of *adiponectin* to help maintain healthy insulin sensitivity.
8. Moderating *glycerol-3-phosphate dehydrogenase enzyme activity* to reduce the amount of ingested starches that are converted to triglycerides and stored as fat.

Two capsules of **Optimized *Irvingia* with Phase 3™**

Calorie Control Complex provide:

Phase 3™ L-Arabinose and chromium complex.....550 mg (supplying 475 mg L-Arabinose and 100 mcg Chromium in a food bound state)

Green Tea Phytosome (decaffeinated) extract.....150 mg

Phase 2® Phaseolus vulgaris white kidney bean extract.... 445 mg

InSea^{2™} seaweed extract.....125 mg

Irvingia gabonensis extract150 mg

Summary of Human Studies with New Optimized *Irvingia* Ingredients

Ingredient	Study	Treatment vs Placebo
Green Tea Phytosome Extract	100 overweight subjects placed on a hypocaloric diet (men: 1,850 calories; women: 1,350 calories) randomized to receive 300 mg/day of green tea phytosome extract or placebo for 90 days ²	<u>Weight loss</u> 30.1 pounds vs. 9.9 pounds <u>Waist size reduction</u> 10% vs. 5% (14% vs. 7% in men)
Phase 2® White Kidney Bean Extract	60 overweight subjects placed on a 2,000-2,200 calorie, carbohydrate-rich diet and randomized to either 445 mg/day of white kidney bean extract or placebo for 30 days ³	<u>Weight loss</u> 6.5 pounds vs. 0.8 pounds <u>Waist size reduction</u> 1.2 inches vs. 0.2 inches
Integra-Lean® <i>Irvingia Gabonensis</i> Extract	102 overweight subjects randomized to either 150 mg of <i>Irvingia</i> twice daily or placebo for 10 weeks ⁵	<u>Weight loss</u> 28 pounds vs. 1.5 pounds <u>Waist size reduction</u> 6.4 inches vs. 2.1 inches
Phase 3™	50 non-diabetic subjects were used to study the suppression effects of L-Arabinose and chromium on capillary glucose ⁹	Consuming LA-Cr simultaneously with a 70 gram sucrose challenge suppressed the glucose response an average of -20% compared to control over four different time periods.

Note: Supplements should be taken in conjunction with a healthy diet and regular exercise program. Results may vary.



Item #01492

Directions are to take two capsules before the two heaviest meals of the day. A bottle of **120** capsules of **Optimized *Irvingia* with Phase 3™ Calorie Control Complex** retails for \$78. If a member buys four bottles during **Super Sale**, the price is reduced to **\$48.60 per bottle**.

InSea^{2™} is a trademark of innoVactiv, Inc. Integra-Lean[®] Irvingia is protected by U.S. Patent No. 7,537,790. Other patents pending. Phase 2® and Phase 3™ are used under license.

Caution: This product is designed to target several critical factors involved in age-related weight gain. Those who ingest more calories than what their body has the metabolic capacity to utilize will not see results. This

is because some people are ingesting so many excess calories that no matter how much their metabolic rate is increased, or how much improvement occurs in their post-meal blood sugar and serum triglyceride levels, or how much youthful insulin sensitivity and other body fat-regulating systems are restored, they are overwhelming the metabolic capacity to utilize these calories. This will result in excess calories being stored in adipocytes. One cannot consume limitless calories and expect to shed fat pounds by taking drugs, nutrients, and/or hormones that demonstrate weight-loss effects in clinical studies.

References:

1. *Science*. 2009 Jul 10;325(5937):201-4.
2. *Integr Nutr*. 2008;1(2):1-14.
3. *Int J Med Sci*. 2007;4:45-52.
4. http://www.innovativ.com/index.php?option=com_content&task=view&id=18&Itemid=5
5. *Lipids Health Dis*. 2009 Mar 2;8:7.
6. *Lipids Health Dis*. 2008 Nov 13;7:44.
7. *Ann Nutr Metab*. 1993;37(1):14-23.
8. *Lipids Health Dis*. 2008 Mar 31;7:12.
9. Submitted for publication, 2009.





BY JULIUS GOEPP, MD

Reverse Mitochondrial Damage

Potent Molecular Energizers for Lifelong Health

Progressive loss of function in the *mitochondria*—the cellular generators responsible for nearly all the body's energy output—speeds aging and death.

Mitochondrial *dysfunction* has been linked to an array of degenerative illnesses, ranging from diabetes and neurological disorders to heart failure.^{1,2}

In 2007, a group of researchers reported a **major (but little-known) breakthrough** in our understanding of *how* mitochondrial dysfunction unfolds—and what can be done to protect yourself against its lethal impact.³

They discovered that potentially deadly defects in human mitochondria, including molecular decay and membrane injury, begin to appear and can be detected **nearly a decade before the onset of permanent damage to the DNA.**³

More importantly, their analysis revealed that in its initial stages, mitochondrial dysfunction is **reversible**, enabling the life and health of cells to be **prolonged** at the molecular level. The key lies in **early** interventions to ensure *optimal* mitochondrial function before **irreversible** DNA damage occurs.

In this article, we review the latest research on a set of compounds that specifically target and enhance mitochondrial function through multiple modes of action. > >



The Cellular Death Spiral

Mitochondria are responsible for converting energy from the food you ingest into usable “currency.” Carbohydrates, fats, and proteins are broken down inside your cells into components that enter the cellular powerhouses known as mitochondria. Throughout this cellular journey, these “macronutrients” undergo a complex series of biochemical transformations that generate *adenosine triphosphate* (ATP), the molecular energy currency behind *all* biological functions. To give you an idea of ATP’s life-sustaining importance, your body converts a volume of ATP equal to your entire weight *every day*.

At the core of this energy conversion matrix lies the **electron transport chain**, a series of molecules embedded in the inner mitochondrial membrane. It serves as the “power line” through which needed chemical energy is released and transferred into vital ATP.

This **energy-intensive** process throws off an immense number of electrons within the mitochondria, resulting in constant exposure to **free radicals**—and rendering the mitochondria especially vulnerable to oxidative damage.⁴⁻⁶

The result is a cellular *death spiral*: the mitochondria gradually deteriorate, leading to a decrease in vital ATP production and a deadly increase in free

radical generation. Over time, this continuous free-radical onslaught destroys the mitochondria through progressive membrane damage and molecular decay.

As levels of oxidative damage from mitochondrial dysfunction steadily rise with age,^{7,8} the body’s antioxidant defenses gradually weaken *at the same time*, accelerating cellular senescence and death.^{4,9,10}

Left unchecked, this fatal cycle speeds the general decline in overall function that accompanies aging^{4,11} and contributes to the onset of degenerative disease.¹²⁻¹⁵

CoQ10’s Rejuvenating Power

Coenzyme Q10 (CoQ10) powerfully safeguards mitochondria from age-related decay and death through two principal pathways.

It plays an essential role in the **electron transport chain**, facilitating the efficient transfer of electrons into ATP for use in cellular function.¹⁶ CoQ10 resides primarily on the inner membranes of the mitochondria; **95%** of all cellular energy production depends on it.

CoQ10 also acts as a powerful free radical scavenger, neutralizing their lethal action and dramatically reducing oxidative damage. The more available CoQ10 in the mitochondria, the less free radical damage.¹⁶ This is one of the reasons why the highest CoQ10

concentrations are found in the most energy-intensive organs: the brain, heart, liver, and kidneys.¹⁷

CoQ10 levels in our vital organs, like the heart, steadily rise after birth and peak at about 20 years of age. After that, they undergo a continuous decline.¹⁸ Fortunately, three decades of cutting-edge research have shown us how to *restore* CoQ10 levels in the mitochondria to slow and even reverse the effects of aging.¹⁹⁻²²

In pre-clinical models, CoQ10 supplementation protects tissue from lethal DNA damage and increases lifespan.²³ It boosts mitochondrial function and total energy output in heart muscle in aging animals.²⁴ And in animal models, lifelong CoQ10 supplementation has been shown to decrease oxidative damage in skeletal muscle, increase native antioxidant enzymes, and favorably modify age-related changes in muscular energy metabolism.²⁵

Until 2007, the only form of CoQ10 available was *ubiquinone*. Unfortunately, the ubiquinone form of CoQ10 has limited absorption.²⁶ Another form of CoQ10, known as *ubiquinol*, remains up to eight times longer in the blood.^{27,28}

The Heart Health Warrior

Dense with mitochondria, the heart requires more energy than any other organ—and the greatest concentration of CoQ10.²⁹ This is especially true for aging individuals, even those with advanced chronic heart disease. Scores of studies show that chronic heart conditions, including **congestive heart failure** (CHF), are characterized by diminished levels of CoQ10 in heart tissue. Its therapeutic benefit has proven just as profound for these individuals.

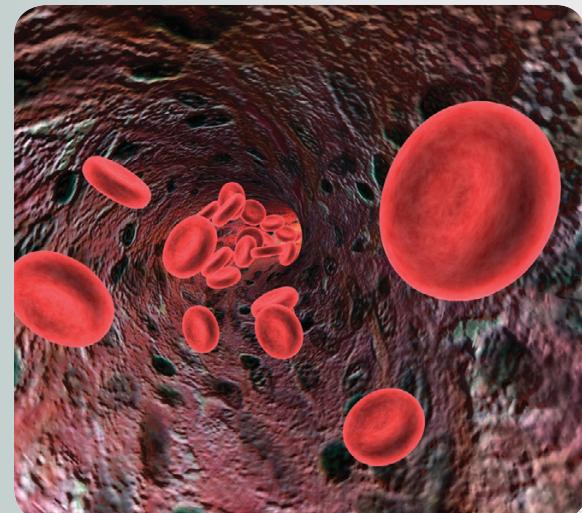
In a 2008 study, **standard** CoQ10 supplements failed to improve either CoQ10 levels or cardiac performance in individuals suffering from CHF, while **ubiquinol** succeeded on both fronts.³⁰

The study involved individuals with advanced CHF. Their hearts pumped less than half as well as normal, with low CoQ10 levels *despite* taking an average of **450 mg/day** of standard CoQ10. When the same people took ubiquinol (**580 mg/day** on average), their CoQ10 blood levels vaulted into the therapeutic range—and their hearts' pumping action improved by **77%**.

At the outset of this study, every participant suffered from category IV CHF (the most severe form), presenting continuous symptoms—even at rest—with severely limited activity. By the end of the study, the average CHF score had fallen to category II, indicated by mild symptoms (such as slight shortness of breath and/or angina) and minimal limitations during ordinary activity.³¹

Mitochondria Health

- Mitochondria are the cellular organelles that power every energy-requiring bodily process.
- Progressive loss of function in the *mitochondria*—the cellular power generators responsible for nearly all energy output in the body—speeds cell aging and death.
- Researchers recently discovered that signs of age-related mitochondrial damage appear nearly a decade before the onset of **permanent** DNA damage.
- They also found that mitochondrial decay and dysfunction are reversible.
- A handful of mitochondrial-energizing nutrients have been shown to offer powerful protection from mitochondrial damage and dysfunction.
- CoQ10 speeds mitochondrial electron transport, increases energy production, and protects tissues from mitochondrial decline.
- Shilajit, an ancient Indian adaptogen, enhances CoQ10's mitochondrial benefits and supports levels of the active ubiquinol form.
- R-alpha-lipoic acid further supports mitochondrial energy production.
- Acetyl-L-carnitine “feeds” energy-releasing molecules to mitochondria, improving their efficiency and preventing damage.

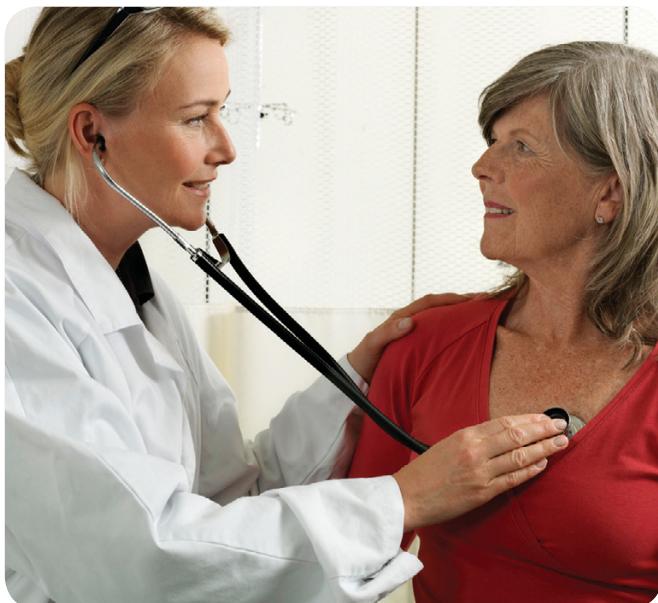


CoQ10 supplementation also increases heart muscle **contractility**—the strength of the heart's squeezing action—enabling the heart to pump more blood more efficiently, even in patients with advanced CHF.³²

Chronic CoQ10 deficiency has been linked to poor surgical outcomes in elderly patients compared to younger ones.^{21,33} By energizing cardiac mitochondria, CoQ10 exerts a powerful effect on cardiac performance in individuals with CHF. Supplementation with CoQ10 and other antioxidants and heart-energizing nutrients such as L-carnitine and taurine reduces distended heart volume in patients—a vital factor in reducing the risk of bypass surgery.³⁴

Following a heart attack, cardiac tissue is at great risk for further injury, including a second attack. In patients recovering from recent heart attacks, just **120 mg** of CoQ10 per day produced remarkable benefits.³⁵ After one year, only **25%** of supplemented patients suffered a cardiac event, compared with **45%** in the placebo group, and cardiac deaths were significantly fewer compared with placebo. Supplemented patients also had increased high-density lipoprotein (HDL) and dramatically lower measures of oxidative stress.

Mitochondrial dysfunction is linked to a broad range of degenerative illness, from diabetes and neurological disorders to heart disease.



CoQ10 also benefits people undergoing cardiac surgery, particularly older adults whose outcomes tend to be worse than younger people's, owing to declining mitochondrial function and density in heart tissue.²¹

Treating heart tissue with CoQ10 improves its metabolic stress response and speeds recovery after ischemia (loss of blood flow)—two major concerns after cardiac surgery.²¹ Oral CoQ10 therapy for one week before surgery improves mitochondrial energy efficiency and post-operative heart function, while reducing heart muscle damage and shortening hospital stays.²¹ A 2008 study also showed significantly fewer arrhythmias (abnormal heart beats), less need for medications to boost cardiac strength, and less need for blood transfusion in patients who received CoQ10 supplementation prior to cardiac surgery, compared to patients who did not receive CoQ10.³⁶

Potent Endothelial Defense

The lining of our blood vessels, or **endothelium**, regulates blood flow and pressure, and is easily damaged by oxidative stress and inflammation, which increases the risk of atherosclerosis and erectile dysfunction. CoQ10 powerfully protects endothelial function, an effect that is likely due to its uniquely beneficial effect on mitochondrial function.³⁷

One study of the **ubiquinol** form of CoQ10 showed that it protects against hypertension, improves endothelial function, and reduces cardiac enlargement in stroke-prone rats.³⁸ When humans with endothelial dysfunction took **300 mg/day** of CoQ10 orally for one month, their blood vessels relaxed more readily and they moved more oxygenated blood into tissues compared with placebo recipients.³⁹

People with type 2 diabetes are at particularly high risk for endothelial dysfunction, and have more heart attacks and strokes as a result.⁴⁰ CoQ10 supplementation is especially effective at improving endothelial function in this population.⁴¹ Diabetics (and others) often need to take statin-type lipid-lowering medications to control their cholesterol levels. Unfortunately, these drugs (also known as HMG Co A reductase inhibitors) are known to deplete CoQ10 and can cause muscle pain that may be related to this depletion.⁴² CoQ10 overcomes this problem and has been shown to improve endothelial function in diabetic patients on statins.⁴³



Muscular Energy Enhancement

Exercise can boost longevity and even increase mitochondrial density in the short term; however, exercise can also *damage* the mitochondria in the long term.^{44,45} The high rate of oxygen and electron flow that exercise requires can lead to chronically low ATP levels, which may exert negative effects during vigorous exercise.⁴⁶ CoQ10 supplementation can counteract such effects, enhancing the adaptive response of skeletal muscle following exercise.⁴⁷

CoQ10 supplementation before exercise increases muscle CoQ10 levels, reduces muscular oxidant stress, and may increase the amount of time you can exercise until exhaustion.⁴⁸ To take one dramatic example, CoQ10 supplementation of **300 mg/day** resulted in improved blood markers of exercise-induced muscle injury among elite Japanese Kendo athletes (a form of martial arts) practicing up to five-and-a-half hours per day.⁴⁹

CoQ10 at just **100 mg/day** even enhances performance of normally sedentary men during repeated bouts of exercising.⁵⁰ Supplementation of **300 mg/day** enabled adults to increase their velocity on a stationary bike compared with placebo, while reducing fatigue.⁵¹

CoQ10's remarkable energy-boosting effects can also reduce adverse effects associated with statin therapy, including fatigue, muscle pain, shortness of breath, memory loss, and nerve pain in the extremities.⁵² Patients with statin-induced fatigue who stopped the drug and took **240 mg/day** of CoQ10 saw a decrease in fatigue from **84%** to just **15%**; a drop in muscle pain from **64%** to **6%**, and a decline in shortness of breath from **58%** to **12%**. These are all manifestations of restored mitochondrial energy and function—and the study found no adverse consequences from discontinuing the statin drugs. (You should never abruptly discontinue **any** medication without discussing it with your doctor.)

System-Wide Protection

CoQ10 also has dramatic benefits in other tissues, particularly in the brain, eyes, and skin.

There's growing recognition of the role of "brain energetics," including mitochondrial health, in causing (or preventing) progressively fatal neurological conditions, including Alzheimer's disease, Parkinson's disease, and amyotrophic lateral sclerosis (ALS or Lou Gehrig's Disease).^{12,53} Animal studies show that CoQ10 supplementation increases brain levels of CoQ10, sustaining the brain's tremendous energy needs. At the same time it reduces brain injury and increases life span in mice with a neurodegenerative disease similar to ALS.⁵⁴

CoQ10 benefits peripheral nerves as well. People with diabetes often succumb to painful diabetic neuropathy and decreased ability to sense pressure, which can lead to disastrous injuries. Studies of diabetic rats with neuropathy show that CoQ10 improved nerve conduction velocity and strength of nerve impulses.⁵⁵

Nerve cells in the eye are faced with enormous energy demands—they must convert light into electrical impulses, while protecting themselves from the damaging effects of both.⁵⁶ Researchers now know that mitochondrial health is vital to sustaining the health of cells in the retina, where optical nerves are concentrated.⁵⁷ Unfortunately, CoQ10 levels in the retina decline rapidly with age, leaving delicate cells vulnerable.⁵⁸

In combination with acetyl-L-carnitine and omega-3 fatty acids, CoQ10 generated dramatic results in studies of individuals suffering from early age-related macular degeneration. Supplemented patients had a **10-fold lower risk** of worsening over a 12-month period, compared with those who received placebo.⁵⁹ Pre-clinical models suggest that CoQ10 may even protect retinal tissue from the effects of glaucoma.⁵⁶

Our skin shows the most immediate and visible signs of aging. Only recently have we learned how

Advanced Mitochondrial Threats: Glycation and Lipoxidation

The chemical reaction of glucose with proteins and fats that occurs over a lifetime produces **advanced glycation end-products** (AGEs) and **advanced lipoxidation end-products** (ALEs).¹⁰⁴ These deadly molecules cause oxidative and inflammatory damage to mitochondria, hastening mitochondrial dysfunction and aging.¹⁰⁵⁻¹⁰⁹ Specific compounds have been shown to provide *targeted mitochondrial defense* against **glycation** and the inflammation it produces.

- **Carnosine** is a nutrient comprised of two amino acids. It's a natural antioxidant and anti-glycation molecule proven to reduce reactive oxygen and nitrogen species resulting from chronic glucose exposure, while also binding to potentially dangerous metal ions (chelation). These features make it attractive as an anti-aging, anti-Alzheimer's agent.¹¹⁰⁻¹¹²
- **Luteolin** is a flavonoid with potent anti-inflammatory effects.^{113,114} It directly inhibits AGE formation at early, middle, and late stages in their development—more powerfully than standard chemical AGE-inhibitors.^{115,116} It also directly counters the sugar-induced mitochondrial damage caused by reduction in a survival protein called Bcl-2.¹¹⁷
- **Benfotiamine** is a fat-soluble form of thiamine (vitamin B1). Its higher bioavailability allows it to strongly increase glycation-fighting thiamine levels in blood and tissues in normal people and in people with either type 1 or 2 diabetes.¹¹⁸⁻¹²⁰ Benfotiamine powerfully reduces AGE production and damage to vascular endothelial cells under high-glucose conditions.^{121,122} It blocks three distinct pathways of sugar-induced tissue damage to protect against retinal damage in diabetes.¹²³
- **Pyridoxal -5'-phosphate** (PLP) is the biologically active form of vitamin B6. It is a powerful inhibitor of both protein and fat glycation.^{124,125} Glycation reductions by PLP are credited with reducing sugar-induced blood vessel and kidney damage from diabetes.^{126,127}

Each of these nutrients works through distinct pathways, acting as a “therapeutic cocktail” that provides maximum protection against glycation-induced toxicity and mitochondrial damage.¹²⁸

much this has to do with mitochondrial dysfunction in skin cells: skin biopsies from older people show substantially less mitochondrial function than those from younger people.⁶⁰

Increased oxidative damage from diminished mitochondrial function has been shown to trigger inflammation and launch protein-destroying enzymes into action. Over time this leads to a weakening of the delicate matrix of skin tissue, spots, wrinkles, dryness—even cancer.⁶¹ Many studies show that topical CoQ10 treatment inhibits inflammatory cytokines, reduces wrinkling enzyme production, and improves the appearance and radiation-resistance of older skin.⁶¹⁻⁶³ Boosting CoQ10 through oral supplementation also affords vital protection.⁶⁴

Mitochondrial Protection with a Potent Adaptogen

Long known to Ayurvedic practitioners for its healing power, **shilajit** is an organic substance harvested from biomass high in the Himalayas.^{65,66} It acts as a powerful **adaptogen**, providing broad systemic defense against stress and illness. Cutting-edge scientific analysis has isolated **humic substances** as the principal active ingredients that enhance mitochondrial energy flow.⁶⁷

In 2009, a series of landmark studies detailed for the first time how shilajit works on energy metabolism.

Mice subjected to strenuous exercise underwent expected ATP declines in muscle, blood, and brain tissue. **When supplemented with shilajit, ATP loss was sharply reduced.**⁶⁸ Other biochemical markers of energy status also dramatically improved in the supplemented animals—including levels of CoQ10, which fell twice as fast in control mice as in supplemented animals. When given in combination, CoQ10 and shilajit displayed a powerful **synergistic effect**. Energy parameters such as CoQ10 levels increased significantly more than with either supplement alone.

Further analysis brought some of its key *mechanisms of action* to light. **Shilajit** contains two primary components, **fulvic acid** and **DBPs** (dibenzo-apyrone). **Fulvic acid** independently stimulates mitochondrial energy metabolism, protects mitochondrial membranes from oxidative damage, and helps channel electron-rich **DBPs** into the mitochondria to support the electron transfer chain.^{69,70} Fulvic acid works as an electron “shuttle,” augmenting CoQ10 to speed electron flow within mitochondria.⁷¹⁻⁷³

The DBPs in shilajit serve as electron “reservoirs,” replenishing electrons lost by CoQ10 when it donates them to free radicals (thereby neutralizing them).^{70,74}

When laboratory mice are supplemented with oral CoQ10 alone, CoQ10 levels rise in heart, liver, and kidney tissue, as might be expected.⁷⁵ When DBPs from *shilajit* are added to the supplement, CoQ10 levels rise still further—as much as **29%** in the liver.⁷⁵

A recent study suggests that DBPs from shilajit preserve CoQ10 in its superior ***ubiquinol*** form.⁷⁵

Preliminary findings suggest that shilajit protects human tissue from lost energy in the form of ATP, while maximizing benefits from CoQ10, with dramatic improvement in exercise performance.⁷⁶ In an as-yet unpublished study, people who took shilajit **200 mg** once daily for 15 days registered **14%** higher post-exercise ATP levels in the blood—equivalent to levels in people who hadn’t exercised at all. The average number of steps they took on a standardized dynamic exercise test rose significantly, and their mean fitness scores increased by **15%**—without any intervening exercise training.

In pre-clinical studies, shilajit has been shown to possess a number of additional benefits, allowing it to work *in synergy* with CoQ10 to protect and support mitochondrial health:

- Preliminary unpublished studies showed that shilajit (**250 mg** twice daily for 90 days) lowered fasting **blood sugar** and a measure of **systemic inflammation** called the ESR (erythrocyte sedimentation rate), while increasing hemoglobin levels and platelet counts.^{77,78}
- Shilajit protected laboratory rats from developing chemically-induced **diabetes** through its free-radical scavenging properties.⁷⁰
- Shilajit *augmented* learning acquisition and memory retrieval in laboratory rats while reducing manifestations of anxiety during maze experiments.⁷⁰
- Shilajit reduced levels of the enzyme *acetylcholinesterase* that destroys the vital neurotransmitter **acetylcholine**. This effect may help to prevent or treat Alzheimer’s disease by maintaining levels of the neurotransmitter.⁷⁰
- Shilajit increased levels of the neurotransmitter **dopamine** in rat brains, making it an attractive candidate for treatment of Parkinson’s disease and other movement disorders.⁷⁰
- In pre-clinical studies, shilajit produced significant increases in the endogenous antioxidants **superoxide dismutase** (SOD), **catalase**, and **glutathione peroxidase** in brain tissue. Increased levels of these enzymes protect vulnerable brain cells against the oxidative damage that leads to brain aging and cognitive decline.⁷⁰

A Complementary Coenzyme

Lipoic acid is a naturally occurring compound found in mitochondria. Like CoQ10, it is a **coenzyme** required for proper function of the mitochondrial energy chain.⁴ Lipoic acid directly increases ATP production in mitochondria.⁷⁹ Clinical models indicate that lipoic acid may serve as a first-line defense for diseases involving impaired energy utilization, including diabetes and the nerve damage associated with it.⁸⁰⁻⁸²

R-alpha-lipoic acid is the most bioactive form of lipoic acid—and a powerful activator of mitochondrial energy complexes.^{83,84} Studies in aging animals support the use of R-alpha-lipoic acid to improve mitochondrial function, decrease oxidative damage, and increase metabolic rate, all of which otherwise become impaired with aging.⁴

R-alpha-lipoic acid has been proven effective in reducing symptoms of diabetic neuropathy, without significant adverse reactions.^{81,85} It also increases nerve conduction velocity in people with diabetic neuropathy, crucial to improved nerve signaling.⁸⁶



Experts attribute these effects to diminished fat oxidation in nerve cell membranes and improvements in local blood supply around nerves resulting from improved mitochondrial functioning.^{87,88}

R-alpha-lipoic acid displays many protective effects. It reverses the age-related increase in liver cell damage caused by exogenous toxins, helping to protect liver function.⁸⁹ It prevents brain cells from becoming depleted of the natural antioxidant *reduced glutathione*, an important intracellular antioxidant in the body. Deficiency of *reduced glutathione* can predispose people to liver failure, Parkinson's disease, and other neurodegenerative conditions.^{90,91} A therapeutic dose of **600 mg/day** even helped relieve migraine attack rates—an observation that may support the theory that migraines may be partially *caused* by impaired mitochondrial function.⁹²

As you might expect of a mitochondrial energy booster, lipoic acid may also play a role in helping to ward off cardiovascular disease. Three months of lipoic acid supplementation provided pain relief to patients with **peripheral vascular disease** (PVD), extending the time they could walk before pain occurred.⁹³ Combined therapy with acetyl-L-carnitine improved blood vessel relaxation and blood flow, while reducing blood pressure, in patients with coronary artery disease.⁹⁴ And combined supplementation is a very good idea, as we'll see next.

Your Mitochondrial Fat-Burner

L-carnitine is a molecule required for helping transport fatty acids into the mitochondria, where they can be burned as fuel. Acetyl-L-carnitine (ALC) is the *form* of carnitine optimally absorbed through oral delivery. It has also been shown to boost mitochondrial health, facilitating fuel delivery to the electron transport chain, where supplements like CoQ10, shilajit, and lipoic acid take over.

Total carnitine levels diminish with age, a decline that may also be accelerated by overeating and diabetes.⁹⁵ As with other mitochondrial energy optimizers, ALC supplementation possesses distinct benefits across numerous physiological systems.⁹⁶

A review of clinical studies shows that ALC may slow the natural course of Alzheimer's disease.⁹⁷ It has substantially increased Alzheimer's disease patients' responses to drug treatment, from **38%** to **50%** in one study.⁹⁸ ALC also protects brain tissue against destructive effects of hypoxia (low oxygen), by supporting cellular metabolism.⁹⁹ ALC and lipoic acid supplementation partially restored depleted brain mitochondrial activity in aged rats to that of young adults.¹⁰⁰



The combination of ALC with lipoic acid improved cognitive function in a mouse model of Alzheimer's disease.¹⁰¹ ALC alone has exhibited powerful effects, restoring aging animals' cardiac energy metabolism to that of young adults.¹⁰² In combination with lipoic acid, ALC helps maintain heart muscle function in aging animals as well.¹⁰³

Summary

Mitochondrial dysfunction is linked to a broad range of degenerative illness, from diabetes and neurological disorders to heart disease. Researchers have discovered that age-related mitochondrial dysfunction—which can ultimately lead to DNA damage and cell death—may be prevented and even *reversed*. The key lies in early and sustained interventions that support optimal mitochondrial health and function. CoQ10 in its superior *form* as **ubiquinol** may restore mitochondrial function. The organic adaptogen **shilajit** acts in synergy with ubiquinol, further enhancing mitochondrial function. R-alpha-lipoic acid and acetyl-L-carnitine have been shown in clinical studies to provide additional mitochondrial support. ●

If you have any questions on the scientific content of this article, please call a Life Extension® Health Advisor at 1-866-864-3027.

References

- Conley KE, Amara CE, Jubrias SA, Marcinek DJ. Mitochondrial function, fibre types and ageing: new insights from human muscle in vivo. *Exp Physiol.* 2007 Mar;92(2):333-9.
- Lesnfsky EJ, Moghaddas S, Tandler B, Kerner J, Hoppel CL. Mitochondrial dysfunction in cardiac disease: ischemia—reperfusion, aging, and heart failure. *J Mol Cell Cardiol.* 2001 Jun;33(6):1065-89.
- Conley KE, Marcinek DJ, Villarin J. Mitochondrial dysfunction and age. *Curr Opin Clin Nutr Metab Care.* 2007 Nov;10(6):688-92.
- Hagen TM, Ingersoll RT, Lykkesfeldt J, et al. (R)-alpha-lipoic acid-supplemented old rats have improved mitochondrial function, decreased oxidative damage, and increased metabolic rate. *FASEB J.* 1999 Feb;13(2):411-8.
- Shigenaga MK, Hagen TM, Ames BN. Oxidative damage and mitochondrial decay in aging. *Proc Natl Acad Sci U S A.* 1994 Nov 8;91(23):10771-8.
- Genova ML, Pich MM, Bernacchia A, et al. The mitochondrial production of reactive oxygen species in relation to aging and pathology. *Ann N Y Acad Sci.* 2004 Apr;1011:86-100.
- Sohal RS, Weindruch R. Oxidative stress, caloric restriction, and aging. *Science.* 1996 Jul 5;273(5271):59-63.
- Hagen TM, Yowe DL, Bartholomew JC, et al. Mitochondrial decay in hepatocytes from old rats: membrane potential declines, heterogeneity and oxidants increase. *Proc Natl Acad Sci U S A.* 1997 Apr 1;94(7):3064-9.
- Sanz N, Diez-Fernandez C, Alvarez A, Cascales M. Age-dependent modifications in rat hepatocyte antioxidant defense systems. *J Hepatol.* 1997 Sep;27(3):525-34.
- Erdinclar DS, Seven A, Inci F, Beger T, Candan G. Lipid peroxidation and antioxidant status in experimental animals: effects of aging and hypercholesterolemic diet. *Clin Chim Acta.* 1997 Sep 8;265(1):77-84.
- DiMauro S, Tanji K, Bonilla E, Pallotti F, Schon EA. Mitochondrial abnormalities in muscle and other aging cells: classification, causes, and effects. *Muscle Nerve.* 2002 Nov;26(5):597-607.
- Sullivan PG, Brown MR. Mitochondrial aging and dysfunction in Alzheimer's disease. *Prog Neuropsychopharmacol Biol Psychiatry.* 2005 Mar;29(3):407-10.
- Choksi KB, Nuss JE, Boylston WH, Rabek JP, Papaconstantinou J. Age-related increases in oxidatively damaged proteins of mouse kidney mitochondrial electron transport chain complexes. *Free Radic Biol Med.* 2007 Nov 15;43(10):1423-38.
- Baines CP. The mitochondrial permeability transition pore and ischemia-reperfusion injury. *Basic Res Cardiol.* 2009 Mar;104(2):181-8.
- Di Lisa F, Kaludercic N, Carpi A, Menabo R, Giorgio M. Mitochondria and vascular pathology. *Pharmacol Rep.* 2009 Jan-Feb;61(1):123-30.
- Sohal RS, Forster MJ. Coenzyme Q, oxidative stress and aging. *Mitochondrion.* 2007 Jun;7 Suppl:S103-11.
- Aberg F, Appelkvist EL, Dallner G, Ernster L. Distribution and redox state of ubiquinones in rat and human tissues. *Arch Biochem Biophys.* 1992 Jun;295(2):230-4.
- Kalen A, Appelkvist EL, Dallner G. Age-related changes in the lipid compositions of rat and human tissues. *Lipids.* 1989 Jul;24(7):579-84.
- Bliznakov EG. Immunological senescence in mice and its reversal by coenzyme Q10. *Mech Ageing Dev.* 1978 Mar;7(3):189-97.
- Rosenfeldt FL, Pepe S, Linnane A, et al. The effects of ageing on the response to cardiac surgery: protective strategies for the ageing myocardium. *Biogerontology.* 2002;3(1-2):37-40.
- Rosenfeldt FL, Pepe S, Linnane A, et al. Coenzyme Q10 protects the aging heart against stress: studies in rats, human tissues, and patients. *Ann N Y Acad Sci.* 2002 Apr;959:355-39; discussion 463-35.
- Aejmelaeus R, Metsa-Ketela T, Laippala P, Solakivi T, Alho H. Ubiquinol-10 and total peroxyl radical trapping capacity of LDL lipoproteins during aging: the effects of Q-10 supplementation. *Mol Aspects Med.* 1997;18 Suppl:S11320.
- Quiles JL, Ochoa JJ, Huertas JR, Mataix J. Coenzyme Q supplementation protects from age-related DNA double-strand breaks and increases lifespan in rats fed on a PUFA-rich diet. *Exp Gerontol.* 2004 Feb;39(2):189-94.
- Ochoa JJ, Quiles JL, Huertas JR, Mataix J. Coenzyme Q10 protects from aging-related oxidative stress and improves mitochondrial function in heart of rats fed a polyunsaturated fatty acid (PUFA)-rich diet. *J Gerontol A Biol Sci Med Sci.* 2005 Aug;60(8):970-5.
- Ochoa JJ, Quiles JL, Lopez-Frias M, Huertas JR, Mataix J. Effect of lifelong coenzyme Q10 supplementation on age-related oxidative stress and mitochondrial function in liver and skeletal muscle of rats fed on a polyunsaturated fatty acid (PUFA)-rich diet. *J Gerontol A Biol Sci Med Sci.* 2007 Nov;62(11):1211-8.
- Kaikkonen J, Tuomainen TP, Nyysonen K, Salonen JT. Coenzyme Q10: absorption, antioxidative properties, determinants, and plasma levels. *Free Radic Res.* 2002 Apr;36(4):389-97.
- Hosoe K, Kitano M, Kishida H, Kubo H, Fujii K, Kitahara M. Study on safety and bioavailability of ubiquinol (Kaneka QH) after single and 4-week multiple oral administration to healthy volunteers. *Regul Toxicol Pharmacol.* 2007 Feb;47(1):19-28.
- Shults CW, Flint Beal M, Song D, Fontaine D. Pilot trial of high dosages of coenzyme Q10 in patients with Parkinson's disease. *Exp Neurol.* 2004 Aug;188(2):491-4.
- Soukoulis V, Dihu JB, Sole M, et al. Micronutrient deficiencies an unmet need in heart failure. *J Am Coll Cardiol.* 2009 Oct 27;54(18):1660-73.
- Langsjoen PH, Langsjoen AM. Supplemental ubiquinol in patients with advanced congestive heart failure. *Biofactors.* 2008;32(1-4):119-28.
- Available at: http://www.abouthf.org/questions_stages.htm. Accessed October 9, 2009.
- Belardinelli R, Mucaj A, Licalaprice F, et al. Coenzyme Q10 improves contractility of dysfunctional myocardium in chronic heart failure. *Biofactors.* 2005;25(1-4):137-45.
- Morisco C, Trimarco B, Condorelli M. Effect of coenzyme Q10 therapy in patients with congestive heart failure: a long-term multicenter randomized study. *Clin Investig.* 1993;71(8 Suppl):S134-6.
- Jeejeebhoy F, Keith M, Freeman M, et al. Nutritional supplementation with MyoVive repletes essential cardiac myocyte nutrients and reduces left ventricular size in patients with left ventricular dysfunction. *Am Heart J.* 2002 Jun;143(6):1092-100.
- Singh RB, Neki NS, Kartikey K, et al. Effect of coenzyme Q10 on risk of atherosclerosis in patients with recent myocardial infarction. *Mol Cell Biochem.* 2003 Apr;246(1-2):75-82.
- Makhija N, Sendasgupta C, Kiran U, et al. The role of oral coenzyme Q10 in patients undergoing coronary artery bypass graft surgery. *J Cardiothorac Vasc Anesth.* 2008 Dec;22(6):832-9.
- Kuettner A, Pieper A, Koch J, Enzmann F, Schroeder S. Influence of coenzyme Q(10) and cerivastatin on the flow-mediated vasodilation of the brachial artery: results of the ENDOTACT study. *Int J Cardiol.* 2005 Feb 28;98(3):413-9.
- Graham D, Huynh NN, Hamilton CA, et al. Mitochondria-targeted antioxidant MitoQ10 improves endothelial function and attenuates cardiac hypertrophy. *Hypertension.* Aug 2009;54(2):322-8.
- Tiano L, Belardinelli R, Carnevali P, Principi F, Seddaiu G, Littarru GP. Effect of coenzyme Q10 administration on endothelial function and extracellular superoxide dismutase in patients with ischaemic heart disease: a double-blind, randomized controlled study. *Eur Heart J.* 2007 Sep;28(18):2249-55.
- Beckman JA, Creager MA, Libby P. Diabetes and atherosclerosis: epidemiology, pathophysiology, and management. *JAMA.* 2002 May 15;287(19):2570-81.
- Watts GF, Playford DA, Croft KD, Ward NC, Mori TA, Burke V. Coenzyme Q(10) improves endothelial dysfunction of the brachial artery in Type II diabetes mellitus. *Diabetologia.* 2002 Mar;45(3):420-6.
- Maroff L, Thompson PD. The role of coenzyme Q10 in statin-associated myopathy: a systematic review. *J Am Coll Cardiol.* 2007 Jun 12;49(23):2231-7.

43. Hamilton SJ, Chew GT, Watts GF. Coenzyme Q10 improves endothelial dysfunction in statin-treated type 2 diabetic patients. *Diabetes Care.* 2009 May;32(5):810-2.

44. Di Meo S, Venditti P. Mitochondria in exercise-induced oxidative stress. *Biol Signals Recept.* 2001 Jan-Apr;10(1-2):125-40.

45. Powers SK, Jackson MJ. Exercise-induced oxidative stress: cellular mechanisms and impact on muscle force production. *Physiol Rev.* 2008 Oct;88(4):1243-76.

46. Yegutkin GG, Samburski SS, Mortensen SP, Jalkanen S, Gonzalez-Alonso J. Intravascular ADP and soluble nucleotidases contribute to acute prothrombotic state during vigorous exercise in humans. *J Physiol.* 2007 Mar 1;579(Pt 2):553-64.

47. Hellsten Y, Nielsen JJ, Lykkesfeldt J, et al. Antioxidant supplementation enhances the exercise-induced increase in mitochondrial uncoupling protein 3 and endothelial nitric oxide synthase mRNA content in human skeletal muscle. *Free Radic Biol Med.* 2007 Aug 1;43(3):353-61.

48. Cooke M, Iosia M, Buford T, et al. Effects of acute and 14-day coenzyme Q10 supplementation on exercise performance in both trained and untrained individuals. *J Int Soc Sports Nutr.* 2008;5:8.

49. Kon M, Tanabe K, Akimoto T, et al. Reducing exercise-induced muscular injury in kendo athletes with supplementation of coenzyme Q10. *Br J Nutr.* 2008 Oct;100(4):903-9.

50. Gokbel H, Gul I, Belviranli M, Okudan N. The Effects Of Coenzyme Q10 Supplementation on Performance During Repeated Bouts of Supramaximal Exercise in Sedentary Men. *J Strength Cond Res.* 2009 Jul 28.

51. Mizuno K, Tanaka M, Nozaki S, et al. Antifatigue effects of coenzyme Q10 during physical fatigue. *Nutrition.* 2008 Apr;24(4):293-9.

52. Langsjoen PH, Langsjoen JO, Langsjoen AM, Lucas LA. Treatment of statin adverse effects with supplemental Coenzyme Q10 and statin drug discontinuation. *Biofactors.* 2005;25(1-4):147-52.

53. Kidd PM. Neurodegeneration from mitochondrial insufficiency: nutrients, stem cells, growth factors, and prospects for brain rebuilding using integrative management. *Altern Med Rev.* 2005 Dec;10(4):268-93.

54. Matthews RT, Yang L, Browne S, Baik M, Beal MF. Coenzyme Q10 administration increases brain mitochondrial concentrations and exerts neuroprotective effects. *Proc Natl Acad Sci U S A.* 1998 Jul 21;95(15):8892-7.

55. Ayaz M, Tuncer S, Okudan N, Gokbel H. Coenzyme Q(10) and alpha-lipoic acid supplementation in diabetic rats: conduction velocity distributions. *Methods Find Exp Clin Pharmacol.* 2008 Jun;30(5):367-74.

56. Russo R, Cavaliere F, Rombola L, et al. Rational basis for the development of coenzyme Q10 as a neurotherapeutic agent for retinal protection. *Prog Brain Res.* 2008;173:575-82.

57. Feher J, Papale A, Mannino G, Gualdi L, Balacco Gabrieli C. Mitotropic compounds for the treatment of age-related macular degeneration. The metabolic approach and a pilot study. *Ophthalmologica.* 2003 Sep-Oct;217(5):351-7.

58. Qu J, Kaufman Y, Washington I. Coenzyme Q10 in the human retina. *Invest Ophthalmol Vis Sci.* 2009 Apr;50(4):1814-8.

59. Feher J, Kovacs B, Kovacs I, et al. Improvement of visual functions and fundus alterations in early age-related macular degeneration treated with a combination of acetyl-L-carnitine, n-3 fatty acids, and coenzyme Q10. *Ophthalmologica.* 2005 May-Jun;219(3):154-66.

60. Prahrl S, Kueper T, Biernoth T, et al. Aging skin is functionally anaerobic: importance of coenzyme Q10 for anti aging skin care. *Biofactors.* 2008;32(1-4):245-55.

61. Inui M, Ooe M, Fujii K, Matsunaka H, Yoshida M, Ichihashi M. Mechanisms of inhibitory effects of CoQ10 on UVB-induced wrinkle formation in vitro and in vivo. *Biofactors.* 2008;32(1-4):237-43.

62. Hoppe U, Bergemann J, Diembeck W, et al. Coenzyme Q10, a cutaneous antioxidant and energizer. *Biofactors.* 1999;9(2-4):371-8.

63. Blatt T, Lenz H, Koop U, et al. Stimulation of skin's energy metabolism provides multiple benefits for mature human skin. *Biofactors.* 2005;25(1-4):179-85.

64. Passi S, De Pità O, Grandinetti M, Simotti C, Littarru GP. The combined use of oral and topical lipophilic antioxidants increases their levels both in sebum and stratum corneum. *Biofactors.* 2003;18(1-4):289-97.

65. Schepetkin IA, Xie G, Jutila MA, Quinn MT. Complement-fixing activity of fulvic acid from Shilajit and other natural sources. *Phytother Res.* 2009 Mar;23(3):373-84.

66. Goel RK, Banerjee RS, Acharya SB. Antiulcerogenic and antiinflammatory studies with shilajit. *J Ethnopharmacol.* 1990 Apr;29(1):95-103.

67. Agarwal SP, Khanna R, Karmarkar R, Anwer MK, Khar RK. Shilajit: a review. *Phytother Res.* 2007 May;21(5):401-5.

68. Bhattacharyya S, Pal D, Gupta AK, Ganguly P, Majumder UK, Ghosal S. Beneficial effect of processed shilajit on swimming exercise induced impaired energy status of mice. *Pharmacologyonline.* 2009;1:817-25.

69. Piotrowska D, Dlugosz A, Witkiewicz K, Pajak J. The research on antioxidative properties of TOLPA Peat Preparation and its fractions. *Acta Pol Pharm.* 2000 Nov;57 Suppl:127-9.

70. Ghosal S. *Shilajit in Perspective.* Oxford, U.K.: Narosa Publishing House; 2006.

71. Visser SA. Effect of humic substances on mitochondrial respiration and oxidative phosphorylation. *Sci Total Environ.* 1987 Apr;62:347-54.

72. Royer RA, Burgos WD, Fisher AS, Unz RF, Dempsey BA. Enhancement of biological reduction of hematite by electron shuttling and Fe(II) complexation. *Environ Sci Technol.* 2002 May 1;36(9):1939-46.

73. Kang SH, Choi W. Oxidative degradation of organic compounds using zero-valent iron in the presence of natural organic matter serving as an electron shuttle. *Environ Sci Technol.* 2009 Feb 1;43(3):878-83.

74. Islam A, Ghosh R, Banerjee D, Nath P, Mazumder U, Ghosal S. Biotransformation of 3-hydroxydibenzo-pyrone into 3,8 dihydroxydibenzo-pyrone and aminoacyl conjugates by *Aspergillus niger* isolated from native "shilajit." *Electronic Journal of Biotechnology.* 2008 Jul 15;11(3):2-10.

75. Bhattacharyya S, Pal D, Banerjee D, et al. Shilajit dibenzo-pyrone: Mitochondria targeted antioxidants. *Pharmacologyonline.* 2009; 2:690-8.

76. Pal D, Bhattacharya S. Pilot Study on the Improvement of Human Performance with ReVitalETTM as Energy Booster: Part-IV. 2006. Data on file. Natreon, Inc.

77. Clinical study for evaluation of safe use in purified and standardized shilajit in normal volunteers. J. B. Roy State Ayurvedic Medical College and Hospital, Kolkata. 2007. Data on file. Natreon, Inc.

78. Clinical study for evaluation of plasma antioxidant capacity and safe use of purified and standardized Shilajit (ReVitalET) in normal volunteers. J. B. Roy State Ayurvedic Medical College and Hospital, Kolkata. 2007. Data on file. Natreon, Inc.

79. Zimmer G, Mainka L, Kruger E. Dihydrolipoic acid activates oligomycin-sensitive thiol groups and increases ATP synthesis in mitochondria. *Arch Biochem Biophys.* 1991 Aug 1;288(2):609-13.

80. Jacob S, Henriksen EJ, Schiemann AL, et al. Enhancement of glucose disposal in patients with type 2 diabetes by alpha-lipoic acid. *Arzneimittelforschung.* 1995 Aug;45(8):872-4.

81. Ziegler D, Hanefeld M, Ruhnau KJ, et al. Treatment of symptomatic diabetic peripheral neuropathy with the anti-oxidant alpha-lipoic acid. A 3-week multicentre randomized controlled trial (ALADIN Study). *Diabetologia.* 1995 Dec;38(12):1425-33.

82. Sachse G, Willms B. Efficacy of thioctic acid in the therapy of peripheral diabetic neuropathy. *Horm Metab Res Suppl.* 1980;9:105-7.

83. Loffelhardt S, Bonaventura C, Locher M, Borbe HO, Bisswanger H. Interaction of alpha-lipoic acid enantiomers and homologues with the enzyme components of the mammalian pyruvate dehydrogenase complex. *Biochem Pharmacol.* 1995 Aug 25;50(5):637-46.

84. Carlson DA, Smith AR, Fischer SJ, Young KL, Packer L. The plasma pharmacokinetics of R-(+)-lipoic acid administered as sodium R-(+)-lipoate to healthy human subjects. *Altern Med Rev.* 2007 Dec;12(4):343-51.

85. Tankova T, Koev D, Dakovska L. Alpha-lipoic acid in the treatment of autonomic diabetic neuropathy (controlled, randomized, open-label study). *Rom J Intern Med.* 2004;42(2):457-64.

86. Negrisanu G, Rosu M, Bolte B, Lefter D, Dabelea D. Effects of 3-month treatment with the antioxidant alpha-lipoic acid in diabetic peripheral neuropathy. *Rom J Intern Med.* 1999 Jul-Sep;37(3):297-306.

87. Androne L, Gavan NA, Veresiu IA, Orasan R. In vivo effect of lipoic acid on lipid peroxidation in patients with diabetic neuropathy. *In Vivo.* 2000 Mar-Apr;14(2):327-30.

88. Haak E, Usadel KH, Kusterer K, et al. Effects of alpha-lipoic acid on microcirculation in patients with peripheral diabetic neuropathy. *Exp Clin Endocrinol Diabetes.* 2000;108(3):168-74.

89. Hagen TM, Vinarsky V, Wehr CM, Ames BN. (R)-alpha-lipoic acid reverses the age-associated increase in susceptibility of hepatocytes to tert-butylhydroperoxide both in vitro and in vivo. *Antioxid Redox Signal.* 2000 Fall;2(3):473-83.

90. Bharat S, Cochran BC, Hsu M, Liu J, Ames BN, Andersen JK. Pre-treatment with R-lipoic acid alleviates the effects of GSH depletion in PC12 cells: implications for Parkinson's disease therapy. *Neurotoxicology.* 2002 Oct;23(4-5):479-86.

91. Suh JH, Wang H, Liu RM, Liu J, Hagen TM. (R)-alpha-lipoic acid reverses the age-related loss in GSH redox status in post-mitotic tissues: evidence for increased cysteine requirement for GSH synthesis. *Arch Biochem Biophys.* 2004 Mar 1;423(1):126-35.

92. Magis D, Ambrosini A, Sandor P, Jacquy J, Laloux P, Schoenen J. A randomized double-blind placebo-controlled trial of thioctic acid in migraine prophylaxis. *Headache.* 2007 Jan;47(1):52-7.

93. Vincent HK, Bourguignon CM, Vincent KR, Taylor AG. Effects of alpha-lipoic acid supplementation in peripheral arterial disease: a pilot study. *J Altern Complement Med.* 2007 Jun;13(5):577-84.

94. McMackin CJ, Widlansky ME, Hamburg NM, et al. Effect of combined treatment with alpha-Lipoic acid and acetyl-L-carnitine on vascular function and blood pressure in patients with coronary artery disease. *J Clin Hypertens (Greenwich).* 2007 Apr;9(4):249-55.

95. Noland RC, Koves TR, Seiler SE, et al. Carnitine insufficiency caused by aging and overnutrition compromises mitochondrial performance and metabolic control. *J Biol Chem.* 2009 Aug 21;284(34):22840-52.

96. Rosca MG, Lemieux H, Hoppel CL. Mitochondria in the elderly: Is acetyl-carnitine a rejuvenator? *Adv Drug Deliv Rev.* 2009 Aug 29.

97. Carta A, Calvani M. Acetyl-L-carnitine: a drug able to slow the progress of Alzheimer's disease? *Ann N Y Acad Sci.* 1991;640:228-32.

98. Bianchetti A, Rozzini R, Trabucchi M. Effects of acetyl-L-carnitine in Alzheimer's disease patients unresponsive to acetylcholinesterase inhibitors. *Curr Med Res Opin.* 2003;19(4):350-3.

99. Corbucci GG, Melis A, Piga M, Marchionni A, Calvani M. Influence of acetyl-carnitine on some mitochondrial enzymic activities in the human cerebral tissue in conditions of acute hypoxia. *Int J Tissue React.* 1992;14(4):183-94.

100. Long J, Gao F, Tong L, Cotman CW, Ames BN, Liu J. Mitochondrial decay in the brains of old rats: ameliorating effect of alpha-lipoic acid and acetyl-L-carnitine. *Neurochem Res.* Apr 2009 Apr;34(4):755-63.

101. Shenk JC, Liu J, Fischbach K, et al. The effect of acetyl-L-carnitine and R-alpha-lipoic acid treatment in ApoE4 mouse as a model of human Alzheimer's disease. *J Neurol Sci.* 2009 Aug 15;283(1-2):199-206.

102. Lesnfsky EJ, He D, Moghaddas S, Hoppel CL. Reversal of mitochondrial defects before ischemia protects the aged heart. *FASEB J.* 2006 Jul;20(9):1543-5.

103. Hagen TM, Moreau R, Suh JH, Visioli F. Mitochondrial decay in the aging rat heart: evidence for improvement by dietary supplementation with acetyl-L-carnitine and/or lipoic acid. *Ann N Y Acad Sci.* 2002 Apr;959:491-507.

104. Rahbar S. Novel inhibitors of glycation and AGE formation. *Cell Biochem Biophys.* 2007;48(2-3):147-57.

105. Adeghate E. Molecular and cellular basis of the aetiology and management of diabetic cardiomyopathy: a short review. *Mol Cell Biochem.* 2004 Jun;261(1-2):187-91.

106. Schleicher E, Friess U. Oxidative stress, AGE, and atherosclerosis. *Kidney Int Suppl.* 2007 Aug;(106):S17-26.

107. Tan AL, Forbes JM, Cooper ME. AGE, RAGE, and ROS in diabetic nephropathy. *Semin Nephrol.* 2007 Mar;27(2):130-43.

108. Gasser A, Forbes JM. Advanced glycation: implications in tissue damage and disease. *Protein Pept Lett.* 2008;15(4):385-91.

109. Peppa M, Uribarri J, Vlassara H. Aging and glycoxidant stress. *Hormones (Athens).* 2008 Apr-Jun;7(2):123-32.

110. Reddy VP, Garrett MR, Perry G, Smith MA. Carnosine: a versatile antioxidant and antiglycating agent. *Sci Aging Knowledge Environ.* 2005 May 4;2005(18):pe12.

111. Hipkiss AR. Would carnosine or a carnivorous diet help suppress aging and associated pathologies? *Ann N Y Acad Sci.* 2006 May;1067:369-74.

112. Hipkiss AR. Could carnosine or related structures suppress Alzheimer's disease? *J Alzheimers Dis.* 2007 May;11(2):229-40.

113. Kotanidou A, Xagorari A, Bagli E, et al. Luteolin reduces lipopolysaccharide-induced lethal toxicity and expression of proinflammatory molecules in mice. *Am J Respir Crit Care Med.* 2002 Mar 15;165(6):818-23.

114. Kim JS, Jobin C. The flavonoid luteolin prevents lipopolysaccharide-induced NF-kappaB signalling and gene expression by blocking IkappaB kinase activity in intestinal epithelial cells and bone-marrow derived dendritic cells. *Immunology.* 2005 Jul;115(3):375-87.

115. Wu CH, Yen GC. Inhibitory effect of naturally occurring flavonoids on the formation of advanced glycation endproducts. *J Agric Food Chem.* 2005 Apr 20;53(8):3167-73.

116. Psotova J, Chlopickova S, Miketova P, Hrbac J, Simanek V. Chemoprotective effect of plant phenolics against anthracycline-induced toxicity on rat cardiomyocytes. Part III. Apigenin, baicalein, kaempferol, luteolin and quercetin. *Phytother Res.* 2004 Jul;18(7):516-21.

117. Wu CH, Wu CF, Huang HW, Jao YC, Yen GC. Naturally occurring flavonoids attenuate high glucose-induced expression of proinflammatory cytokines in human monocytic THP-1 cells. *Mol Nutr Food Res.* 2009 Aug;53(8):984-95.

118. Volvert ML, Seyen S, Piette M, et al. Benfotiamine, a synthetic S-acyl thiamine derivative, has different mechanisms of action and a different pharmacological profile than lipid-soluble thiamine disulfide derivatives. *BMC Pharmacol.* 2008;8:10.

119. Du X, Edelstein D, Brownlee M. Oral benfotiamine plus alpha-lipoic acid normalises complication-causing pathways in type 1 diabetes. *Diabetologia.* 2008 Oct;51(10):1930-2.

120. Stirban A, Negrean M, Stratmann B, et al. Benfotiamine prevents macro- and microvascular endothelial dysfunction and oxidative stress following a meal rich in advanced glycation end products in individuals with type 2 diabetes. *Diabetes Care.* 2006 Sep;29(9):2064-71.

121. Pomerio F, Molinar Min A, La Selva M, Allione A, Molinatti GM, Porta M. Benfotiamine is similar to thiamine in correcting endothelial cell defects induced by high glucose. *Acta Diabetol.* 2001;38(3):135-8.

122. Marchetti V, Menghini R, Rizza S, et al. Benfotiamine counteracts glucose toxicity effects on endothelial progenitor cell differentiation via Akt/FoxO signaling. *Diabetes.* 2006 Aug;55(8):2231-7.

123. Hammes HP, Du X, Edelstein D, et al. Benfotiamine blocks three major pathways of hyperglycemic damage and prevents experimental diabetic retinopathy. *Nat Med.* 2003 Mar;9(3):294-9.

124. Khatami M, Sultani Z, David I, Li W, Rockey JH. Inhibitory effects of pyridoxal phosphate, ascorbate and aminoguanidine on nonenzymatic glycosylation. *Life Sci.* 1988;43(21):1725-31.

125. Higuchi O, Nakagawa K, Tsuzuki T, Suzuki T, Oikawa S, Miyazawa T. Aminophospholipid glycation and its inhibitor screening system: a new role of pyridoxal 5'-phosphate as the inhibitor. *J Lipid Res.* 2006 May;47(5):964-74.

126. Nakamura S, Niwa T. Pyridoxal phosphate and hepatocyte growth factor prevent dialysate-induced peritoneal damage. *J Am Soc Nephrol.* 2005 Jan;16(1):144-50.

127. Nakamura S, Li H, Adijiang A, Pischetsrieder M, Niwa T. Pyridoxal phosphate prevents progression of diabetic nephropathy. *Nephrol Dial Transplant.* 2007 Aug;22(8):2165-74.

128. Mehta R, Shangari N, O'Brien PJ. Preventing cell death induced by carbonyl stress, oxidative stress or mitochondrial toxins with vitamin B anti-AGE agents. *Mol Nutr Food Res.* 2008 Mar;52(3):379-85.

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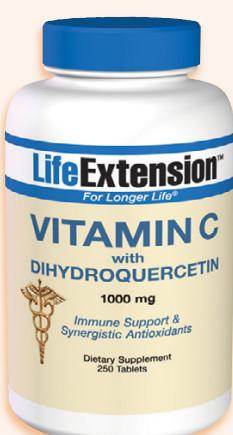
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References:

1. *PLoS Med.* 2005 Sep;2(9):e307; author reply e309.
2. *Am J Clin Nutr.* 1988 Sep;48(3):601-4.
3. *J Food Tech.* 1969;4:255-67.



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References:

1. Hormones (Athens). 2008 Apr-Jun;7(2):123-32.
2. Protein Pept Lett. 2008;15(4):385-91.
3. J Alzheimers Dis. 2007 May;11(2):229-40.
4. Ann N Y Acad Sci. 2006 May;1067:369-74.
5. Sci Aging Knowledge Environ. 2005 May 4;2005(18):pe12.
6. Life Sci. 2007 Nov 30;81(23-24):1602-14.
7. J Nutr. 2006 Jun;136(6):1517-21.
8. Biochem Pharmacol. 2005 Jan 15;69(2):241-8.
9. Immunology. 2005 Jul;115(3):375-87.
10. Am J Respir Crit Care Med. 2002 Mar 15;165(6):818-23.
11. Eur J Pharmacol. 2006 Jul 10;541(1-2):95-105.
12. Nat Med. 2003 Mar;9(3):294-9.
13. Acta Diabetol. 2001;38(3):135-8.
14. Diabetes. 2006 Aug;55(8):2231-7.
15. Diabetes Metab Res Rev. 2008 Jul-Aug;24(5):371-7.
16. J Lipid Res. 2006 May;47(5):964-74.
17. Biochem Biophys Acta. 2001 Feb;14;153(2):110-9.
18. J Am Soc Nephrol. 2005 Jan;16(1):144-50.
19. Life Sci. 1988;43(21):1725-31.
20. Biochem Biophys Res Commun. 1996 Apr 16;221(2):422-9.
21. FASEB J. 1999 Feb;13(2):411-8.
22. Antioxid Redox Signal. 2000 Fall;2(3):473-83.
23. Biochem Mol Biol Int. 1995 Oct;37(2):361-70.
24. Nerenberg Res. 1995 Jan;20(1):1-9.

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1 BOTTLE	\$110	\$74.25
4 BOTTLES	\$98	\$66.15
10 BOTTLES	\$85	\$57.38

The encapsulated version of Life Extension Mix used by many members provides 1 mg of copper. These capsules are also available without copper. The suggested dosage is 14 capsules per day in divided doses with meals. Capsules are easier to swallow for some people than tablets.

LIFE EXTENSION MIX™ 14.81 OZ POWDER

	RETAIL PRICE EACH BOTTLE	SUPER SALE MEMBER PRICE EACH BOTTLE
1 BOTTLE	\$98	\$66.15
4 BOTTLES	\$86	\$58.05
10 BOTTLES	\$72	\$48.60

The powder version of Life Extension Mix contains 1 mg of copper. This powder version is also available without copper. The suggested dose is three scoops per day in divided doses with meals. Because there are so many different ingredients, the powder can be difficult for some people to easily mix. Some people use a blender to combine the Life Extension Mix™ powder with whey protein and other powdered supplements.

To order your supply of LIFE EXTENSION MIX™ during Super Sale, call 1-800-544-4440 or visit www.LifeExtension.com/SuperSale

The Latest Advance in COQ10 Technology

Life Extension® was the first to introduce **coenzyme Q10** to the United States way back in **1983**. Since then, we have consistently introduced more **potent** and better **absorbable** forms of this critical nutrient.

The **new Super Ubiquinol with Enhanced Mitochondrial Support™** contains an organic compound called **PrimaVie® shilajit** that research shows **doubles** levels of CoQ10 in the **mitochondria**.¹

Shilajit has been shown to help restore and sustain **cellular energy**. The latest studies reveal that when **shilajit** is **combined** with **CoQ10**, **cellular energy** gains substantially increase.

In a breakthrough preliminary study, the combination of CoQ10 and **shilajit** produced a **56%** increase in cellular energy production in the **brain—40% better** than CoQ10 alone. In **muscle** there was a **144%** increase, or **27% better** than CoQ10 alone.²

Researchers have found that **shilajit** works to boost CoQ10's beneficial effects by:

1. Stabilizing CoQ10 in its superior **ubiquinol** form, thereby **prolonging** its action at the cellular level.^{3,4}
2. Facilitating more efficient **delivery** of CoQ10 into the **mitochondria**, resulting in greater cellular energy output.⁵⁻⁹

Scientific analysis shows that **shilajit** itself is rich in **essential** compounds that promote mitochondrial metabolism. Part of **shilajit**'s beneficial effects derives from its ability to help the mitochondria convert **fats** and **sugars** into adenosine triphosphate, or **ATP**—the body's main source of energy.⁵⁻⁹

PrimaVie® is a registered trademark of Natreon, Inc.
Kaneka QH® is a registered trademark of Kaneka Corporation.

References:

1. Systemic CoQ level in animals: Part II. Unpublished study. Natreon, Inc.; 2007.
2. *Pharmacologyonline*. 2009;1:817-25.
3. *Pharmacologyonline*. 2009;2:690-8.
4. *Electronic Journal of Biotechnology*. 2008 Jul 15;11(3).
5. Ghosal S. *Shilajit in Perspective*. Alpha Science International Limited; 2006.
6. *Sci Total Environ*. 1987 Apr;62:347-54.
7. *Environ Sci Technol*. 2002 Jul 15;36(14):3170-5.
8. *Environ Sci Technol*. 2002 May 1;36(9):1939-46.
9. *Environ Sci Technol*. 2009 Feb 1;43(3):878-83.

The **new Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™**

Combining **ubiquinol** CoQ10 with **shilajit** generates a powerful **synergy** that supports more youthful cellular energy production than CoQ10 alone.^{2,4,5} What's more, Life Extension has added this novel ingredient to its CoQ10 formulations without increasing the price! So you get a more effective CoQ10 at the same cost!

The retail price for 60 100-mg softgels of **Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™** is \$62. If a member buys four bottles during **Super Sale**, the price is reduced to **\$37.80** per bottle. **Item #01426**

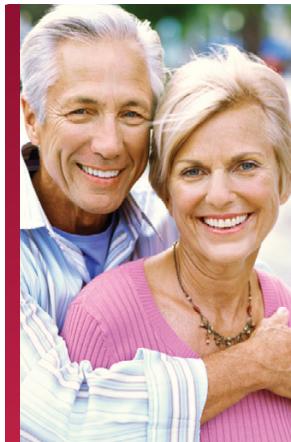
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To order the new
Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™
call **1-800-544-4440** or visit
www.LifeExtension.com.



Item #01426

Item #01425



Discount Prices For Premium-Quality Products

Life Extension® members are longevity enthusiasts, willing to take extraordinary steps to stave off disease, aging, and death. When members buy products from the **Life Extension Foundation Buyers Club**, they are assured of receiving the highest quality products based on the latest scientific studies that demonstrate benefits.

The **large discounts** available to Foundation members enable them to purchase **premium-quality** supplements at prices substantially below those charged by commercial companies.

Here are some examples of **savings** members enjoy during the annual **SUPER SALE**:

		SUPER SALE Our Low Retail Price	Member-Volume Discount Price Per Bottle
Calorie Restriction Mimetic Formula , 60 vegetarian capsules, Item #01419	\$36		\$22.28
Provides in two capsules, 250 mg of trans resveratrol , the highest dose (3 mg) of pterostilbene , more quercetin , plus beneficial grape seed and black tea polyphenols , all to help favorably support the healthy gene expression observed in response to calorie restriction			(four-bottle purchase)
Super Omega-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract , 120 softgels, Item #01482	\$32		\$16.81
Super-refined highly concentrated EPA/DHA fish oil plus sesame lignans and more potent olive fruit extract			(ten-bottle purchase)
Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™ , 100 mg, 60 softgels, Item #01426	\$62		\$35.10
Preferred "ubiquinol" combined with shilajit, an organic compound shown to double levels of CoQ10 in the mitochondria			(ten-bottle purchase)
Optimized Irvingia with Phase 3™ Calorie Control Complex* , 120 vegetarian capsules, Item #01492	\$78		\$48.60
Combats age-related fat accumulation via eight distinct mechanisms			(four-bottle purchase)
Vitamin D3 with Sea-Iodine™ , 5000 IU, 60 vegetarian capsules, Item #01372	\$14		\$8.44
High-potency vitamin D plus 1000 mcg of natural iodine			(four-bottle purchase)
Ultra Natural Prostate Formula , 60 softgels, Item #01475	\$38		\$21.60
Now with more potent Saw Palmetto to support normal urinary flow and help support healthy prostate function			(twelve-bottle purchase)
Prelox® Natural Sex for Men® , 60 tablets, Item #01373	\$50		\$29.90
Supports healthy endothelial function and blood flow for maximum male performance			(four-bottle purchase)
DHEA (Dehydroepiandrosterone) , 25 mg, 100 capsules, Item #00335	\$15		\$8.44
Anti-aging benefits for overall health			(four-bottle purchase)
Arthro-Immune Joint Support , 60 vegetarian capsules, Item #01404	\$30		\$18.23
Combines two clinically proven plant extracts to support healthy joint structure and function			(four-bottle purchase)
Natural Stress Relief , 30 vegetarian capsules, Item #00987	\$28		\$16.20
With lemon balm extract to reduce occasional anxiety and sleeplessness, and pure L-theanine to promote relaxation without drowsiness			(four-bottle purchase)
Mitochondrial Energy Optimizer , 120 capsules, Item #01368	\$86		\$52.65
To maintain healthy cellular function and protein structural integrity			(four-bottle purchase)

To order call toll-free 1-800-544-4440

These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

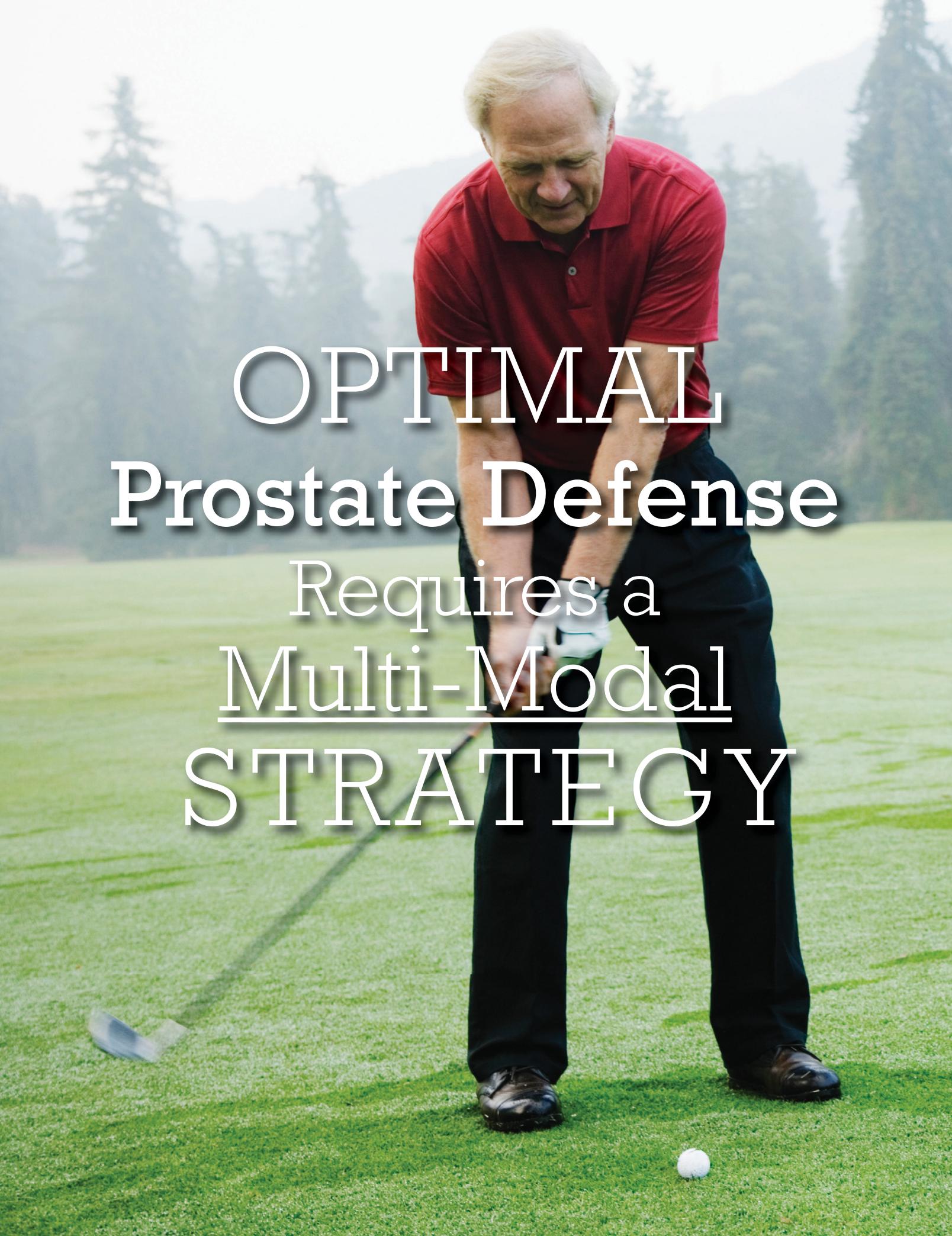
*Note: Supplements should be taken in conjunction with a healthy diet and regular exercise program. Results may vary.

		Our Low Retail Price	SUPER SALE Member-Volume Discount Price Per Bottle
Life Extension Mix™, 315 tablets, Item #01455 High-potency multi-nutrient formula now with twice as much Vitamin D and iodine, plus pterostilbene to favorably regulate gene expression	\$98	\$46.91 (ten-bottle purchase)	
Super R-Lipoic Acid, 300 mg, 60 vegetarian capsules, Item #01208 Double the efficacy of ordinary alpha-lipoic acid ... supplies 210 mg of stabilized R-lipoic acid	\$49	\$29.90 (four-bottle purchase)	
Optimized Tryptopure® Plus, 90 vegetarian capsules, Item #01202 For natural support of healthy serotonin levels in the brain	\$40	\$24.30 (four-bottle purchase)	
Cognitex with Pregnenolone & NeuroProtection Complex, 90 softgels, Item #00922 Sharp-PS® GOLD (PS-DHA), vinpocetine, phosphatidylserine, glyceryl-phosphoryl-choline, pregnenolone and uridine for optimal brain function (also available without pregnenolone)	\$74	\$43.20 (twelve-bottle purchase)	
Pomegranate Extract with CocoaGold™, 30 vegetarian capsules, Item #01256 Each capsule standardized to provide the equivalent of 12.3 ounces of pomegranate juice and 50 mg cocoa polyphenols.	\$24	\$14.18 (four-bottle purchase)	
Acetyl-L-Carnitine Arginate, 100 capsules, Item #00788 Patented form of carnitine to support healthy neurite growth for optimal brain health	\$59	\$34.42 (four-bottle purchase)	
Triple Action Cruciferous Vegetable Extract, 60 vegetarian capsules, Item #01468 Comprehensive cruciferous plant extract formulation, now with apigenin, for optimal cellular health	\$24	\$14.85 (four-bottle purchase)	
Agave Digestive-Immune Support, 360 grams powder, Item #01417 Clinically advanced prebiotic blend supports growth of beneficial bacteria in the colon for healthy immune response	\$30	\$18.23 (four-bottle purchase)	
Bone Restore, 150 capsules, Item #00811 High-potency bone protection formula with FruiteX B® OsteoBoron®	\$22.50	\$13.16 (four-bottle purchase)	
Vital Greens Mix, 309 grams powder, Item #01098 An organic blend of whole plant foods, this green "super food" provides protein, fiber, vitamins, minerals, and mushroom extracts	\$48	\$29.70 (four-bottle purchase)	
Fast-Acting Joint Formula, 30 capsules, Item #00965 One-per-day joint support	\$39	\$24.30 (four-bottle purchase)	
Super MiraForte with Standardized Lignans, 120 capsules, Item #01315 Supports healthy testosterone levels in men	\$62	\$37.80 (four-bottle purchase)	
SAMe, 400 mg, 20 tablets, Item #00557 S-adenosyl-methionine from Europe in double-strength potency to restore sense of well-being	\$35	\$18.90 (six-box purchase)	
Gamma E Tocopherol with Sesame Lignans, 60 softgels, Item #00759 Potent, free radical-quenching vitamin E formula	\$32	\$19.58 (four-bottle purchase)	
Super Bio-Curcumin® 60 vegetarian capsules, Item #00407 Super-absorbable formulation promotes healthy lipid & joint function, and healthy DNA	\$30	\$17.89 (four-bottle purchase)	
Super Booster Softgels with Advanced K2 Complex, 60 softgels, Item #01380 Critical, oil-based nutrients including gamma-tocopherol, sesame lignans, lycopene, lutein, ginkgo, chlorophyllin, selenium and both forms of vitamin K2	\$42	\$25.65 (four-bottle purchase)	
Endothelial Defense with GliSODin® and CocoaGold™, 60 vegetarian capsules, Item #01297 Pomegranate concentrate plus two other nutrients that have been shown to help maintain healthy endothelial function and arterial circulation	\$54	\$32.40 (four-bottle purchase)	

Order online at www.LifeExtension.com/SuperSale

These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

*Note: Supplements should be taken in conjunction with a healthy diet and regular exercise program. Results may vary.

A man in a red polo shirt and dark pants is putting a golf ball into a hole on a green. He is leaning forward with his right leg extended. The background shows a forest of tall evergreen trees.

OPTIMAL Prostate Defense Requires a Multi-Modal STRATEGY

A photograph of two men on a golf course. The man on the left is wearing a blue polo shirt, an orange vest, and brown pants, and is holding a golf club. The man on the right is wearing a white t-shirt and dark pants, and is also holding a golf club. They are standing on a green grassy field with trees in the background.

BY JULIUS GOEPP, MD

As an aging man, your odds of suffering **benign prostate enlargement** (BPH) and/or **prostate cancer** are exceedingly high.

BPH will afflict 50% of men over the course of their lives.^{1,2} At advanced stages, BPH can almost completely obstruct the urethral canal, leading to a host of lower urinary tract symptoms.^{3,4}

Prostate cancer remains the second leading cause of cancer death in men, accounting for more than **28,000 deaths** and **186,000 new cases** in 2008 alone.⁵

This article reviews eight clinically supported interventions with outstanding safety records to protect prostate health. > >



Saw Palmetto—Front Line Defense!

Used by over **2 million** men in the United States, saw palmetto (*Serenoa repens*) remains the front line fighter in natural prostate defense, with a substantial body of clinical investigation and support.⁶ Extracts from the ripe red berries of the plant are rich in bioactive plant-based compounds, including **beta-sitosterol** and a host of vital free fatty acids.⁷⁻⁹

The natural ingredients in saw palmetto benefit the prostate in several related ways. They *inhibit* enzymes that convert testosterone into dihydrotestosterone (DHT),¹⁰ a hormone that increases prostate growth and may promote cancer.¹¹ They bind to DHT receptors on prostate cells, further reducing DHT's potential negative impact.¹⁰ And they block certain transmitters that can aggravate lower urinary tract symptoms.^{10,12,13}

These multiple mechanisms account for saw palmetto's therapeutic effects in managing BPH. In a summary of **18** clinical trials involving **2,939 men**, saw palmetto alleviated lower urinary tract symptoms

(LUTS) and urine flow measures more effectively than placebo.¹⁴ The same analysis also demonstrated that men taking saw palmetto had significantly less nighttime urination (*nocturia*). The analysis also indicated additional benefit by combining saw palmetto with other phytonutrient extracts, including **nettle root** and bark of the African plum tree *Pygeum africanum*.

When placed head-to-head against prescription drugs like *finasteride* (Proscar[®]) and *tamsulosin* (Flomax[®]), saw palmetto was as effective in improving urinary symptom scores and peak urinary flow rates.^{6,15,16} Of even greater interest, men given saw palmetto experienced a lower incidence of **associated sexual dysfunction** compared to those given pharmaceuticals!¹⁵

In fact, few adverse effects have ever been confirmed with saw palmetto extract.¹⁷

The complex of plant-based chemicals (*phytosterols*) in the saw palmetto berry has been shown to exert a suppressive effect on prostate cancer cells. Saw palmetto's high **beta-sitosterol** content—along with other constituents within its phytosterol complex—*inhibit* prostate cancer cell proliferation by selectively arresting cell growth and inducing programmed cell death (*apoptosis*).^{9,18,19} The **carotenoids** contained in saw palmetto also act in tandem with this phytosterol complex to attack cancer cell membranes and slow tumor growth.^{8,19,20} Despite these impressive findings, saw palmetto should not be considered a primary treatment for prostate cancer.

The Complementary Power of Nettle Root

In numerous studies, the root of stinging nettle (*Urtica dioica*) has been shown to exert favorable effects on prostate health that complement those of saw palmetto. Stinging nettle appears to modulate hormone activity in prostate tissue. It also possesses powerful anti-inflammatory and antimicrobial effects—with very low toxicity.^{24,25} Animal studies further indicate its ability to restrict prostate tissue growth.²⁶

Controlled clinical research has demonstrated nettle extract's power to counter and even *reverse* the effects of benign prostatic hypertrophy (BPH). In a double-blind, placebo-controlled study of **558** patients with BPH, nettle extract improved LUTS by **81%**, compared with just **16%** of controls.²⁷ It also significantly increased peak urine flow rates compared with placebo.

When combined with saw palmetto, nettle root extract displays even more impressive effects on BPH. One large study found that the combination was as

effective as the drug *finasteride* (Proscar®) at reducing symptom scores and increasing urine flow, but produced far fewer adverse events.²⁸ Other studies have demonstrated the combination's superior performance over placebo in trials lasting up to 96 weeks—with virtually no side effects.²⁹

Nettle root extract may also fight prostate cancer, reducing cancer cell proliferation without affecting normal tissue.³⁰ It blocks the enzymes that cancer cells need for rapid turnover.³¹ The *lectins* in nettle root extract—proteins involved in cell recognition—display a preference for cancer cells over healthy ones, enhancing nettle root's ability to attack malignant cells and induce production of the cancer-suppressing cytokine interleukin-2 (IL-2).³² As with saw palmetto, nettle root extract alone is not sufficiently effective to be used as a curative treatment for prostate cancer.

Potent Prevention with Flax and Norway Spruce Lignans

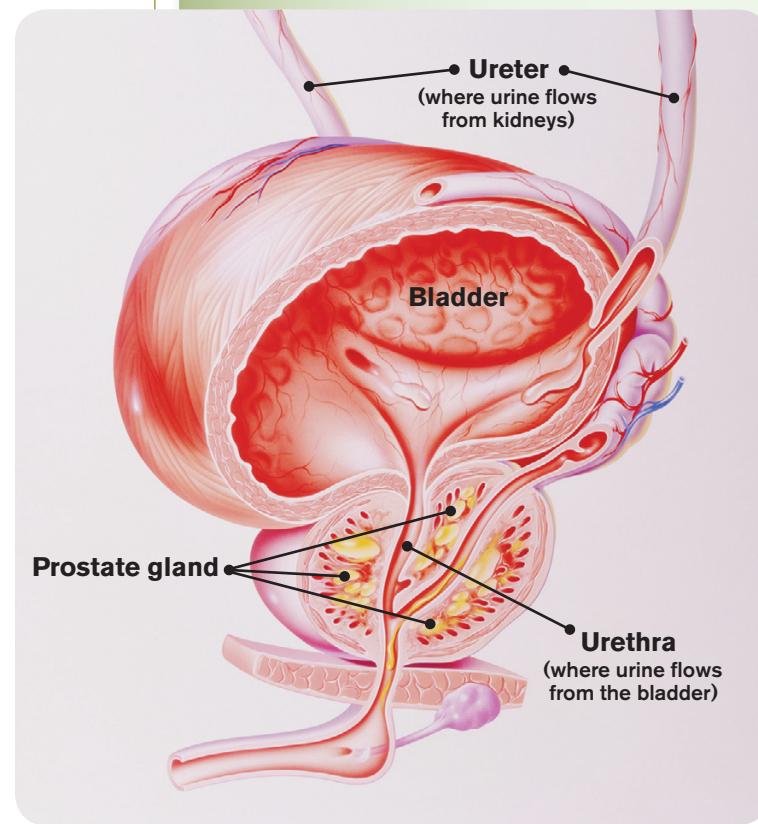
Plants evolved *lignans* to defend against disease. As it happens, their health-promoting benefits are passed on to us when ingested.³³ Resident bacteria in our colon convert these plant lignans into the mammalian lignan *enterolactone*, a phytoestrogen that has been shown to suppress numerous cancers, particularly hormone-dependent types—including prostate and breast cancer.³⁴⁻³⁶ Lignans from flax, Norway spruce (*Picea abies*), and other sources may help prevent both BPH and prostate cancer.

Recent studies have revealed that *flax seed* lignan extract produces improvements in LUTS and quality of life, while also lowering plasma cholesterol and glucose concentrations.^{37,38}

The evidence for their preventive power against prostate cancer is equally compelling. Foods high in plant lignans are associated with lower prostate cancer risk.³⁹ Enterolactone derived from dietary plant lignans has been shown to induce apoptosis (cell death) in cultured human prostate cells.^{40,41} In a clinical setting, flax seed-supplemented diets generated favorable reductions in tumor proliferation rates in men with prostate cancer in as little as **30 days**.⁴² Lignans derived from the Norway spruce have demonstrated powerful anti-cancer effects in an animal model of human prostate cancer, including smaller tumor volume and increased apoptosis.⁴³

Prostate Defense

- Adverse prostate conditions will affect at least half of all men after the age of 50, and 1 in 6 men will be diagnosed with prostate cancer during his lifetime—claiming as many as 28,000 lives each year in the United States.
- Technological advances in extraction techniques optimize the already well-known benefits of saw palmetto extracts.
- Saw palmetto extracts possibly fight BPH and prostate cancer by directly blocking hormone conversion in prostate tissue, and by modifying expression of genes that contribute to cancer development.
- A diverse array of complementary, natural interventions operate across multiple pathways to disrupt prostate diseases at every stage in their development.
- Both alone and in combination, they can be as effective as pharmaceuticals—*without* undesirable side effects, including sexual dysfunction.



Advanced Technology Preserves Saw Palmetto's Bioactive Compounds

The harsh chemical processes and low-pressure techniques often used to extract the saw palmetto berry's bioactive ingredients paradoxically *destroy* many of them. An advanced high-pressure CO₂ extraction technology has been developed that delivers *intact* a far greater proportion of saw palmetto's beneficial, high molecular-weight compounds. The result is a carotenoid-rich extract that most closely reflects the composition of mature saw palmetto berries compared to typical saw palmetto extracts. Carotenoids have demonstrated protective effects against various prostate disorders.²¹⁻²³



Novel Boswellia Extract

A novel extract from *Boswellia serrata*, also known as Indian frankincense, has demonstrated potential in prostate health protection.⁴⁴ It acts as a powerful **5-lipoxygenase** inhibitor. The enzyme **5-lipoxygenase** (5-LOX) inflicts numerous adverse effects⁴⁵ that have been implicated in prostate cancer development. Boswellia extract also favorably modulates gene expression, powerfully suppressing production of pro-inflammatory cytokines involved in BPH and prostate cancer.^{46,47} The anti-inflammatory effects of boswellia extracts have been demonstrated in human studies,⁴⁸ making it an attractive component of BPH prevention.

Research further indicates that boswellia is particularly effective in controlling proliferation of prostate cancer cells. Inhibitors of 5-LOX induce apoptosis in prostate cancer cells through the aptly-named "death receptors" that trigger cancer cell suicide.^{49,50} This extract also blocks prostate cancer growth and proliferation in cell cultures.⁵¹ These effects are due in part to its ability to decrease the androgen receptors that many prostate cancers require to survive.⁵² Boswellia extracts also inhibit new blood vessel growth (*angiogenesis*), depriving malignant tissues of the nutrients and oxygen they need to metastasize.⁵³

Pygeum Africanum

The bark of the African plum tree, *Pygeum africanum*, contains powerful compounds that support the health of prostate and bladder tissue.⁵⁴ Pygeum extracts were proven effective against BPH in numerous open and placebo-controlled studies in the 1990's, and pygeum is now recognized in Europe as a standard therapeutic option.⁵⁵ Pygeum and beta-sitosterol in doses of **50-100 mg** twice daily improve lower urinary tract symptoms and significantly increase urine flow rates, while decreasing residual urine volume in the bladder that can lead to urinary tract infections.⁵⁶⁻⁵⁹

Pygeum extracts also appear to offset the sexual dysfunction that often accompanies BPH and results in overall improvement in quality of life.⁶⁰ One unique mode of action seems to be that pygeum extracts actually inhibit proliferation of the muscle and fibrous tissue in the prostate that help contribute to BPH in the first place.^{61,62} Adverse effects associated with pygeum are rare and mild,⁶³ though some gastrointestinal upset has been reported.

Numerous studies have recently emerged demonstrating pygeum's prostate cancer-fighting potential. Pygeum extracts block the male hormone (*androgen*) receptors that prostate cancer cells need to thrive.⁶⁴ As with the other chemoprotective compounds reviewed in this article, pygeum has been shown to inhibit cancer cell growth and proliferation, while stimulating desired apoptosis (programmed death of cancer cells).⁶⁵ And pygeum's androgen-blocking constituents reduce cancer cells' potential to invade healthy prostate tissue, reducing spread of dangerous tumors.⁶⁶

Saw palmetto extract is one of several natural compounds with clinically proven value in reducing symptoms of BPH and risk factors for prostate cancer.



Novel Flower Pollen Compound

Over two decades of research have confirmed that a specific compound of flower **pollens**, called **cernitin**, possesses unique effects on prostate tissue.⁶⁷⁻⁷⁰ It relaxes certain smooth muscle tissues in the urinary tract,⁷¹ thus potentially alleviating lower urinary tract symptoms associated with BPH. It has been shown to significantly alleviate symptoms of bladder obstruction in men with BPH, while reducing residual urine volume and shrinking the size of the prostate itself.⁶⁷ One major study found cernitin worked as well as six prescription drugs commonly used for BPH.⁷²

Cernitin significantly decreased inflammatory cytokines and tissue inflammation in a rat model of nonbacterial prostatitis, a painful chronic condition that afflicts many men.⁷³ This may render it an ideal

candidate both for the management of BPH and for a complication of chronic prostatitis called **chronic pelvic pain syndrome**, which has proven difficult to treat with standard medication.^{74,75}

In **2009**, a multicenter, randomized, double-blind, placebo-controlled study showed that cernitin significantly improved measures of pain and quality of life in men with category III prostatitis/chronic pelvic pain syndrome, a prevalent condition for which no standardized treatment exists. Overall prostatitis symptom scores were also significantly reduced compared with placebo, *without* major side effects.⁷⁶

Higher doses of cernitin appear to provide faster relief of symptoms.⁷⁷

Unique Mineral Protection

The trace element **boron** combats prostate cancer through a unique combination of underlying mechanisms. Calcium channel signaling is a major regulator of cancer cell proliferation, and yet has received little attention in cancer prevention. Higher boron levels in the **blood** lower the risk of prostate cancer by reducing intracellular calcium signals and storage.⁷⁸ At normal concentrations, boron operates *selectively*, inhibiting prostate cancer cell proliferation while allowing healthy prostate cells to grow.⁷⁹

In animal models boron supplementation inhibited growth of human prostate cancer cells and reduced the size of implanted tumors, reducing local expression of a vital growth factor—a cancer-fighting mechanism unique to boron.⁸⁰ Another study demonstrated that when combined with **phytosterols**, boron amplifies the suppression of cancer cell growth.⁸¹

Several large human studies have shown that **higher dietary boron intake** lowers prostate cancer risk. In one study, men with the highest boron intake **halved** their prostate cancer risk compared to those with the least intake.⁸²

Symptoms of Benign Prostate Hypertrophy

- Urinary hesitancy
- Urinary retention
- Painful urination
- Frequent urination
- Urinary tract infections
- Ejaculatory dysfunction

A Role for Saw Palmetto in Preventing Hair Loss?

Highly-purified saw palmetto extracts benefit the prostate gland by blocking DHT production, regulating prostate cell growth. Compelling new evidence suggests that these same hormonal effects block and even partially reverse hair loss in men with common male pattern baldness!

In 2002, a group of leading-edge scientists recognized that saw palmetto's DHT-blocking action might help in male pattern baldness. In a placebo-controlled, double-blind study, **60%** of men receiving the active supplement showed significant improvement.⁹⁵ A follow-up study suggests that in conjunction with specific anti-inflammatory compounds, saw palmetto may reduce expression of inflammatory genes in hair follicle cells, slashing hair loss risk.⁹⁶

The causes of male pattern baldness are complex.^{100,101} In addition to the shrinkage of hair follicles accelerated by higher DHT levels,^{97,98} sustained microscopic inflammation of hair follicles and remodeling of connective tissue may contribute to making hair loss permanent. Saw palmetto has demonstrated significant reduction of inflammatory markers.^{96,102} Thus, if used before hair loss is advanced, saw palmetto may be an option for addressing the underlying causes of male pattern baldness.



Lycopene

Lycopene is an antioxidant carotenoid found in tomatoes, watermelon, pink grapefruit, and guava. Multiple studies have found that higher intake of lycopene is associated with decreased cancer risk.⁸³⁻⁸⁷ It inhibits prostate cancer cell growth in culture, halting the reproductive cell cycle and inducing apoptosis.^{85,88} Lycopene possesses two unique chemopreventive features:

1. It interferes with internal sex hormone-regulated signaling among cancer cells to prevent coordinated growth.⁸⁹
2. It disrupts DNA synthesis in growing prostate cell cultures, but also protects healthy DNA from damage.^{90,91}

In an early human trial, lycopene improved post-surgical survival rates in aggressively treated patients with advanced prostate cancer, reducing levels of PSA, and shrinking both the primary tumor and metastases.⁹² More recent large-scale studies of men with prostate cancer showed that lycopene—alone or in combination with phytosterols and antioxidants—slowed and even halted rising PSA levels.^{93,94} A 2008 human study demonstrated that lycopene slows disease progression in men with BPH.²²

Summary

Fifty percent of men will eventually suffer enlarged prostate from benign prostatic hyperplasia (BPH), and 1 in 6 men will be diagnosed with prostate cancer during his lifetime. These age-related changes are far from inevitable. Saw palmetto extract is one of several natural compounds with clinically proven value in reducing symptoms of BPH and risk factors for prostate cancer. Each works by uniquely different mechanisms to confer multiple benefits and effective protection. Through a combination of direct impact on prostate tissue and powerful modulation of gene expression, these compounds operate through synergistic and complementary modes of action, attacking BPH and prostate cancer on multiple fronts. Compared to single-target drugs that provide only partial relief (and come with undesirable side effects), these natural interventions provide aging men with a safe, low-cost, alternative—for lifelong prostate health. ●

If you have any questions on the scientific content of this article, please call a Life Extension Health Advisor at 1-866-864-3027.

Prostate Drugs: Limited Benefits, Sexual Side Effects

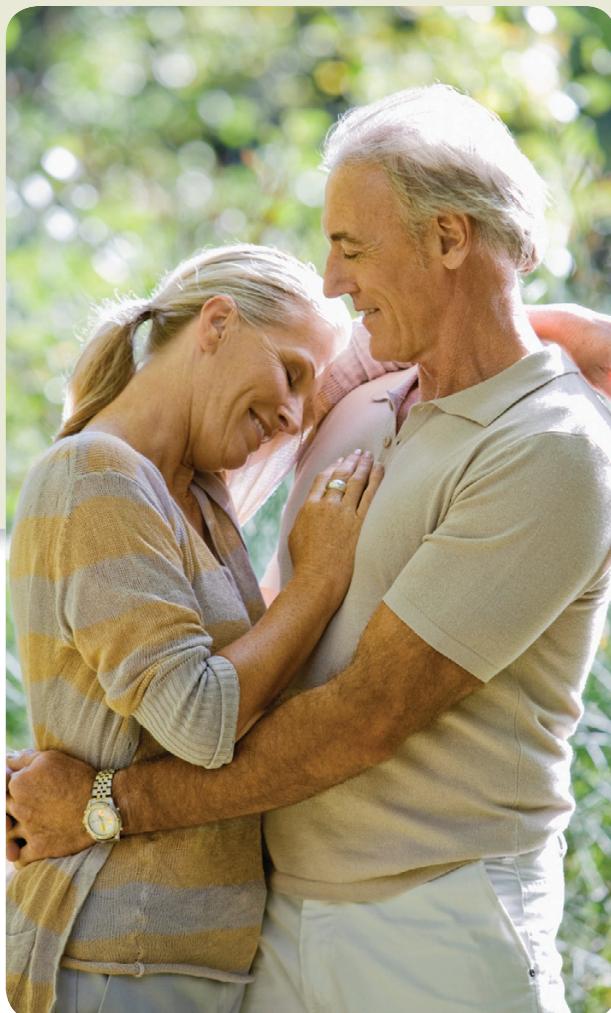
Unlike the multi-targeted benefits of natural compounds, prescription prostate drugs work through narrow, single-targeted mechanisms of action—with a significant array of side effects. These drugs fall into two principal categories.

5-Alpha Reductase Inhibitors. These drugs inhibit the enzyme that converts testosterone into its active dihydrotestosterone (DHT) form. Saw palmetto extracts work by the same mechanism, as well as many others—*without* the side effects of prescription drugs. Finasteride (Proscar®) and dutasteride (Avodart®) are the main players. Both are effective at reducing lower urinary tract symptoms (LUTS) in benign prostatic hypertrophy (BPH).¹⁰³ Finasteride is also approved for use in male pattern baldness (marketed as Propecia®).⁹⁹ Both are also undergoing clinical evaluation for prostate cancer prevention, though to date their efficacy have been uncertain or modest at best.¹⁰⁴

While generally considered safe from a medical standpoint, these drugs have a troubling side effect profile, particularly related to their tendency to cause sexual problems such as erectile dysfunction (ED), ejaculatory dysfunction, decreased libido, and breast enlargement.¹⁰⁵⁻¹⁰⁷ ED is the most common and most troubling side effect.¹⁰⁸ Because BPH itself can cause ED, it can be a difficult decision for men and their physicians about whether to start medication or not. A typical mainstream medical approach when faced with side effects from one drug is to add yet another drug¹⁰⁹ (usually drugs belonging to the following category).

Alpha-adrenergic blockers. These prevent adrenaline from acting on adrenaline receptors in prostate tissue that contribute to BPH. Their sexual side effects are somewhat less pronounced than 5-alpha reductase blockers, with the exception of their impact on ejaculation. In a study of healthy volunteers, tamsulosin (Flomax®) markedly decreased ejaculation volume in almost **90%** of subjects—and **35%** were completely unable to ejaculate.¹⁰⁷

Physicians who prescribe both categories of drugs generally recognize the importance of the sexual side effects that often accompany these drugs. Patients are advised to “consider the sexual dimension” in making treatment choices.^{107,110,111} Another approach is to consider the multi-targeted benefits of nutraceuticals, which are demonstrably free of these side effects—and may offer comparable benefits.^{6,15,16}



References

1. Baazeem A, Elhilali MM. Surgical management of benign prostatic hyperplasia: current evidence. *Nat Clin Pract Urol.* 2008 Oct;5(10):540-9.
2. Vahlensieck W, Jr. With alpha blockers, finasteride and nettle root against benign prostatic hyperplasia. Which patients are helped by conservative therapy? *MMW Fortschr Med.* 2002 Apr 18;144(16):33-6.
3. Roumeguere T, Zouaoui Boudjeltia K, et al. Is there a rationale for the chronic use of phosphodiesterase-5 inhibitors for lower urinary tract symptoms secondary to benign prostatic hyperplasia? *BJU Int.* 2009 Aug;104(4):511-7.

4. Hellstrom WJ, Giuliano F, Rosen RC. Ejaculatory dysfunction and its association with lower urinary tract symptoms of benign prostatic hyperplasia and BPH treatment. *Urology*. 2009 Jul;74(1):15-21.
5. Colli JL, Amling CL. Chemoprevention of prostate cancer: what can be recommended to patients? *Curr Urol Rep*. 2009 May;10(3):165-71.
6. Hizli F, Uygur MC. A prospective study of the efficacy of Serenoa repens, tamsulosin, and Serenoa repens plus tamsulosin treatment for patients with benign prostate hyperplasia. *Int Urol Nephrol*. 2007;39(3):879-86.
7. Abe M, Ito Y, Oyunzul L, Oki-Fujino T, Yamada S. Pharmacologically relevant receptor binding characteristics and 5 α reductase inhibitory activity of free fatty acids contained in saw palmetto extract. *Biol Pharm Bull*. 2009 Apr;32(4):646-50.
8. Petrangeli E, Lenti L, Buchetti B, et al. Lipido-sterolic extract of Serenoa repens (LSEsr, Permixon) treatment affects human prostate cancer cell membrane organization. *J Cell Physiol*. 2009 Apr;219(1):69-76.
9. Scholtysek C, Kruckiewicz AA, Alonso JL, Sharma KP, Sharma PC, Goldmann WH. Characterizing components of the Saw Palmetto Berry Extract (SPBE) on prostate cancer cell growth and traction. *Biochem Biophys Res Commun*. 2009 Feb 13;379(3):795-8.
10. Gerber GS. Saw palmetto for the treatment of men with lower urinary tract symptoms. *J Urol*. 2000 May;163(5):1408-12.
11. Comhaire F, Mahmoud A. Preventing diseases of the prostate in the elderly using hormones and nutriceuticals. *Aging Male*. 2004 Jun;7(2):155-69.
12. Sultan C, Terraza A, Devillier C, et al. Inhibition of androgen metabolism and binding by a liposterolic extract of "Serenoa repens B" in human foreskin fibroblasts. *J Steroid Biochem*. 1984;23:515.
13. Goepel M, Hecker U, Krege S, Rübben H, Michel MC. Saw palmetto extracts potently and non-competitively inhibit human alpha1-adrenoceptors in vitro. *Prostate*. Feb 1999;38(3):208-15.
14. Wilt TJ, Ishani A, Rutks I, MacDonald R. Phytotherapy for benign prostatic hyperplasia. *Public Health Nutr*. 2000 Dec;3(4A):459-72.
15. Fong YK, Milani S, Djavan B. Role of phytotherapy in men with lower urinary tract symptoms. *Curr Opin Urol*. 2005 Jan;15(1):45-8.
16. Engelmann U, Walther C, Bondarenko B, Funk P, Schlaefke S. Efficacy and safety of a combination of sabal and urtica extract in lower urinary tract symptoms. A randomized, double-blind study versus tamsulosin. *Arzneimittelforschung*. 2006;56(3):222-9.
17. Avins AL, Bent S, Staccone S, et al. A detailed safety assessment of a saw palmetto extract. *Complement Ther Med*. 2008 Jun;16(3):147-54.
18. Yang Y, Ikezoe T, Zheng Z, Taguchi H, Koeffler HP, Zhu WG. Saw Palmetto induces growth arrest and apoptosis of androgen-dependent prostate cancer LNCaP cells via inactivation of STAT 3 and androgen receptor signaling. *Int J Oncol*. 2007 Sep;31(3):593-600.
19. Baron A, Mancini M, Caldwell E, Cabrelle A, Bernardi P, Pagano F. Serenoa repens extract targets mitochondria and activates the intrinsic apoptotic pathway in human prostate cancer cells. *BJU Int*. 2009 May;103(9):1275-83.
20. Wadsworth TL, Worstell TR, Greenberg NM, Roselli CE. Effects of dietary saw palmetto on the prostate of transgenic adenocarcinoma of the mouse prostate model (TRAMP). *Prostate*. 2007 May 1;67(6):661-73.
21. Anderson ML. A preliminary investigation of the enzymatic inhibition of 5 α -reduction and growth of prostatic carcinoma cell line LNCap-FCG by natural astaxanthin and Saw Palmetto lipid extract in vitro. *J Herb Pharmacother*. 2005;5(1):17-26.
22. Schwarz S, Obermuller-Jevic UC, Hellmis E, Koch W, Jacobi G, Biesalski HK. Lycopene inhibits disease progression in patients with benign prostate hyperplasia. *J Nutr*. 2008 Jan;138(1):49-53.
23. Liu AG, Volker SE, Jeffery EH, Erdman JW. Feeding Tomato and Broccoli Powders Enriched with Bioactives Improves Bioactivity Markers in Rats. *J Agric Food Chem*. 2009 Aug 3.
24. Gansser D, Spitteler G. Plant constituents interfering with human sex hormone-binding globulin. Evaluation of a test method and its application to *Urtica dioica* root extracts. *Z Naturforsch C*. 1995 Jan-Feb;50(1-2):98-104.
25. Chrubasik JE, Roufogalis BD, Wagner H, Chrubasik S. A comprehensive review on the stinging nettle effect and efficacy profiles. Part II: *urticae radix*. *Phytomedicine*. 2007 Aug;14(7-8):568-79.
26. Lichius JJ, Renneberg H, Blaschek W, Aumuller G, Muth C. The inhibiting effects of components of stinging nettle roots on experimentally induced prostatic hyperplasia in mice. *Planta Med*. 1999 Oct;65(7):666-8.
27. Safarinejad MR. *Urtica dioica* for treatment of benign prostatic hyperplasia: a prospective, randomized, double-blind, placebo-controlled, crossover study. *J Herb Pharmacother*. 2005;5(4):1-11.
28. Søkeland J. Combined sabal and urtica extract compared with finasteride in men with benign prostatic hyperplasia: analysis of prostate volume and therapeutic outcome. *BJU Int*. 2000 Sep;86(4):439-42.
29. Lopatkin N, Sivkov A, Schläfke S, Funk P, Medvedev A, Engelmann U. Efficacy and safety of a combination of Sabal and Urtica extract in lower urinary tract symptoms--long-term follow-up of a placebo-controlled, double-blind, multicenter trial. *Int Urol Nephrol*. 2007;39(4):1137-46.
30. Konrad L, Müller HH, Lenz C, Laubinger H, Aumuller G, Lichius JJ. Antiproliferative effect on human prostate cancer cells by a stinging nettle root (*Urtica dioica*) extract. *Planta Med*. 2000 Feb;66(1):44-7.
31. Durak I, Biri H, Devrim E, Sozen S, Avci A. Aqueous extract of *Urtica dioica* makes significant inhibition on adenosine deaminase activity in prostate tissue from patients with prostate cancer. *Cancer Biol Ther*. 2004 Sep;3(9):855-7.
32. Kayser K, Bubenzier J, Kayser G, et al. Expression of lectin, interleukin-2 and histopathologic blood group binding sites in prostate cancer and its correlation with integrated optical density and syntactic structure analysis. *Anal Quant Cytol Histol*. 1995 Apr;17(2):135-42.



33. Lamblin F, Hano C, Fliniaux O, Mesnard F, Fliniaux MA, Laine E. Interest of lignans in prevention and treatment of cancers. *Med Sci (Paris)*. 2008 May;24(5):511-9.

34. Chen LH, Fang J, Sun Z, et al. Enterolactone inhibits insulin-like growth factor-1 receptor signaling in human prostatic carcinoma PC-3 cells. *J Nutr*. 2009 Apr;139(4):653-9.

35. Wang LQ. Mammalian phytoestrogens: enterodiol and enterolactone. *J Chromatogr B Analyt Technol Biomed Life Sci*. 2002 Sep 25;777(1-2):289-309.

36. Hu C, Yuan YV, Kits TD. Antioxidant activities of the flaxseed lignan secoisolariciresinol diglucoside, its aglycone secoisolariciresinol and the mammalian lignans enterodiol and enterolactone in vitro. *Food Chem Toxicol*. 2007 Nov;45(11):2219-27.

37. Zhang W, Wang X, Liu Y, et al. Effects of dietary flaxseed lignan extract on symptoms of benign prostatic hyperplasia. *J Med Food*. 2008 Jun;11(2):207-14.

38. Zhang W, Wang X, Liu Y, et al. Dietary flaxseed lignan extract lowers plasma cholesterol and glucose concentrations in hypercholesterolaemic subjects. *Br J Nutr*. 2008 Jun;99(6):1301-9.

39. Heald CL, Ritchie MR, Bolton-Smith C, Morton MS, Alexander FE. Phyto-oestrogens and risk of prostate cancer in Scottish men. *Br J Nutr*. 2007 Aug;98(2):388-96.

40. Hedelin M, Klint A, Chang ET, et al. Dietary phytoestrogen, serum enterolactone and risk of prostate cancer: the cancer prostate Sweden study (Sweden). *Cancer Causes Control*. 2006 Mar;17(2):169-80.

41. Chen LH, Fang J, Li H, Demark-Wahnefried W, Lin X. Enterolactone induces apoptosis in human prostate carcinoma LNCaP cells via a mitochondrial-mediated, caspase-dependent pathway. *Mol Cancer Ther*. 2007 Sep;6(9):2581-90.

42. Demark-Wahnefried W, Polascik TJ, George SL, et al. Flaxseed supplementation (not dietary fat restriction) reduces prostate cancer proliferation rates in men presurgery. *Cancer Epidemiol Biomarkers Prev*. 2008 Dec;17(12):3577-87.

43. Bylund A, Saarinen N, Zhang JX, et al. Anticancer effects of a plant lignan 7-hydroxymatairesinol on prostate cancer model in vivo. *Exp Biol Med (Maywood)*. 2005 Mar;230(3):217-23.

44. Ernst E. Frankincense: systematic review. *BMJ*. 2008;337:a2813.

45. Bishnoi M, Patil CS, Kumar A, Kulkarni SK. Protective effects of nimesulide (COX Inhibitor), AKBA (5-LOX Inhibitor), and their combination in aging-associated abnormalities in mice. *Methods Find Exp Clin Pharmacol*. 2005 Sep;27(7):465-70.

46. Roy S, Khanna S, Shah H, et al. Human genome screen to identify the genetic basis of the anti-inflammatory effects of Boswellia in microvascular endothelial cells. *DNA Cell Biol*. 2005 Apr;24(4):244-55.

47. Roy S, Khanna S, Krishnaraju AV, et al. Regulation of vascular responses to inflammation: inducible matrix metalloproteinase-3 expression in human microvascular endothelial cells is sensitive to antiinflammatory Boswellia. *Antioxid Redox Signal*. 2006 Mar-Apr;8(3-4):653-60.

48. Sengupta K, Alluri KV, Satish AR, et al. A double blind, randomized, placebo controlled study of the efficacy and safety of 5-Loxin for treatment of osteoarthritis of the knee. *Arthritis Res Ther*. 2008;10(4):R85.

49. Pommery N, Taverne T, Telliez A, et al. New COX-2/5-LOX inhibitors: apoptosis-inducing agents potentially useful in prostate cancer chemotherapy. *J Med Chem*. 2004 Dec 2;47(25):6195-206.

50. Lu M, Xia L, Hua H, Jing Y. Acetyl-keto-beta-boswellie acid induces apoptosis through a death receptor 5-mediated pathway in prostate cancer cells. *Cancer Res*. 2008 Feb 15;68(4):1180-6.

51. Buchele B, Zugmaier W, Estrada A, et al. Characterization of 3alpha-acetyl-11-keto-alpha-boswellic acid, a pentacyclic triterpenoid inducing apoptosis in vitro and in vivo. *Planta Med*. 2006 Nov;72(14):1285-9.

52. Yuan HQ, Kong F, Wang XL, Young CY, Hu XY, Lou HX. Inhibitory effect of acetyl-11-keto-beta-boswellic acid on androgen receptor by interference of Sp1 binding activity in prostate cancer cells. *Biochem Pharmacol*. 2008 Jun 1;75(11):2112-21.

53. Pang X, Yi Z, Zhang X, et al. Acetyl-11-keto-beta-boswellic acid inhibits prostate tumor growth by suppressing vascular endothelial growth factor receptor 2-mediated angiogenesis. *Cancer Res*. 2009 Jul 15;69(14):5893-900.

54. Wang D, Li Y, Hou G, et al. Pygeum africanum: effect on oxidative stress in early diabetes-induced bladder. *Int Urol Nephrol*. 2009 Jul 16.

55. Breza J, Dzurny O, Borowka A, et al. Efficacy and acceptability of tadenan (Pygeum africanum extract) in the treatment of benign prostatic hyperplasia (BPH): a multicentre trial in central Europe. *Curr Med Res Opin*. 1998;14(3):127-39.

56. Klippen KF, Hiltl DM, Schipp B. A multicentric, placebo-controlled, double-blind clinical trial of beta-sitosterol (phytosterol) for the treatment of benign prostatic hyperplasia. German BPH-Phyto Study group. *Br J Urol*. 1997 Sep;80(3):427-32.

57. Barlet A, Albrecht J, Aubert A, et al. Efficacy of Pygeum africanum extract in the medical therapy of urination disorders due to benign prostatic hyperplasia: evaluation of objective and subjective parameters. A placebo-controlled double-blind multicenter study. *Wien Klin Wochenschr*. 1990 Nov 23;102(22):667-73.

58. Berges RR, Windeler J, Trampisch HJ, Senge T. Randomised, placebo-controlled, double-blind clinical trial of beta-sitosterol in patients with benign prostatic hyperplasia. Beta-sitosterol Study Group. *Lancet*. 1995 Jun 17;345(8964):1529-32.

59. Berges RR, Kassen A, Senge T. Treatment of symptomatic benign prostatic hyperplasia with beta-sitosterol: an 18-month follow-up. *BJU Int*. 2000 May;85(7):842-6.

60. Carani C, Salvioli V, Scuteri A, et al. Urological and sexual evaluation of treatment of benign prostatic disease using Pygeum africanum at high doses. *Arch Ital Urol Nefrol Androl*. 1991 Sep;63(3):341-5.

61. Edgar AD, Levin R, Constantinou CE, Denis L. A critical review of the pharmacology of the plant extract of Pygeum africanum in the treatment of LUTS. *Neurourol Urodyn*. 2007;26(4):458-63; discussion 464.

62. Boulbes D, Soutelle L, Costa P, et al. Pygeum africanum extract inhibits proliferation of human cultured prostatic fibroblasts and myofibroblasts. *BJU Int*. 2006 Nov;98(5):1106-113.

63. Chatelain C, Autet W, Brackman F. Comparison of once and twice daily dosage forms of Pygeum africanum extract in patients with benign prostatic hyperplasia: a randomized, double-blind study, with long-term open label extension. *Urology*. 1999 Sep;54(3):473-8.

64. Schleich S, Papaioannou M, Baniahmad A, Matusch R. Extracts from Pygeum africanum and other ethnobotanical species with antiandrogenic activity. *Planta Med*. 2006 Jul;72(9):807-13.

65. Shenouda NS, Sakla MS, Newton LG, et al. Phytosterol Pygeum africanum regulates prostate cancer in vitro and in vivo. *Endocrine*. 2007 Feb;31(1):72-81.

66. Papaioannou M, Schleich S, Prade I, et al. The natural compound atric acid is an antagonist of the human androgen receptor inhibiting cellular invasiveness and prostate cancer cell growth. *J Cell Mol Med*. 2008 Jul 4.

67. Buck AC, Cox R, Rees RW, Ebeling L, John A. Treatment of outflow tract obstruction due to benign prostatic hyperplasia with the pollen extract, cernilton. A double-blind, placebo-controlled study. *Br J Urol*. 1990 Oct;66(4):398-404.

68. MacDonald R, Ishani A, Rutks I, Wilt TJ. A systematic review of Cernilton for the treatment of benign prostatic hyperplasia. *BJU Int*. 2000 May;85(7):836-41.

69. Shaplygin LV, Sivakov AA. Use of cernilton in the therapy of prostatic adenoma and chronic prostatitis. *Urologia*. 2007 May-Jun(3):35-7, 39.

70. Aslamazov EG, Akhylediani ND, Vinarov AZ, Aliaev Iu G. Cernilton in the treatment of prostatic adenoma and chronic prostatitis. *Urologia*. 2007 Jan-Feb;(1):52, 54-6.

71. Nakase K, Takenaga K, Hamanaka T, Kimura M. Inhibitory effect and synergism of cernitin pollen extract on the urethral smooth muscle and diaphragm of the rat. *Nippon Yakurigaku Zasshi*. 1988 Jun;91(6):385-92.

72. Li NC, Wu SL, Jin J, et al. Comparison of different drugs on the treatment of benign prostate hyperplasia. *Zhonghua Wai Ke Za Zhi*. 2007 Jul 15;45(14):947-50.

73. Asakawa K, Nandachi N, Satoh S, et al. Effects of cernitin pollen extract (Cernilton) on inflammatory cytokines in sex-hormone-induced nonbacterial prostatitis rats. *Hinyokika Kiyo*. 2001 Jul;47(7):459-65.

74. Potts JM. Therapeutic options for chronic prostatitis/chronic pelvic pain syndrome. *Curr Urol Rep.* 2005 Jul;6(4):313-7.

75. Nickel JC. Treatment of chronic prostatitis/chronic pelvic pain syndrome. *Int J Antimicrob Agents.* 2008 Feb;31 Suppl 1:S112-6.

76. Wagenlehner FM, Schneider H, Ludwig M, Schnitker J, Braehler E, Weidner W. A pollen extract (cernilton) in patients with inflammatory chronic prostatitis-chronic pelvic pain syndrome: a multicentre, randomised, prospective, double-blind, placebo-controlled Phase 3 study. *Eur Urol.* 2009 Jun 3.

77. Xu J, Qian WQ, Song JD. A comparative study on different doses of cernilton for preventing the clinical progression of benign prostatic hyperplasia. *Zhonghua Nan Ke Xue.* 2008 Jun;14(6):533-7.

78. Henderson K, Stella SL, Kobylewski S, Eckhert CD. Receptor activated Ca(2+) release is inhibited by boric acid in prostate cancer cells. *PLoS One.* 2009;4(6):e6009.

79. Barranco WT, Eckhert CD. Boric acid inhibits human prostate cancer cell proliferation. *Cancer Lett.* 2004 Dec 8;216(1):21-9.

80. Gallardo-Williams MT, Chapin RE, King PE, et al. Boron supplementation inhibits the growth and local expression of IGF-1 in human prostate adenocarcinoma (LNCaP) tumors in nude mice. *Toxicol Pathol.* 2004 Jan-Feb;32(1):73-8.

81. Barranco WT, Hudak PF, Eckhert CD. Evaluation of ecological and in vitro effects of boron on prostate cancer risk (United States). *Cancer Causes Control.* 2007 Feb;18(1):71-7.

82. Cui Y, Winton MI, Zhang ZF, et al. Dietary boron intake and prostate cancer risk. 2004 Apr;11(4):887-92.

83. Rao AV, Fleshner N, Agarwal S. Serum and tissue lycopene and biomarkers of oxidation in prostate cancer patients: a case-control study. *Nutr Cancer.* 1999;33(2):159-64.

84. Giovannucci E, Rimm EB, Liu Y, Stampfer MJ, Willett WC. A prospective study of tomato products, lycopene, and prostate cancer risk. *J Natl Cancer Inst.* 2002 Mar 6;94(5):391-8.

85. Kim L, Rao AV, Rao LG. Effect of lycopene on prostate LNCaP cancer cells in culture. *J Med Food.* 2002 Winter;5(4):181-7.

86. Etminan M, Takkouche B, Caamano-Isorna F. The role of tomato products and lycopene in the prevention of prostate cancer: a meta-analysis of observational studies. *Cancer Epidemiol Biomarkers Prev.* 2004 Mar;13(3):340-5.

87. Wu K, Erdman JW, Jr, Schwartz SJ, et al. Plasma and dietary carotenoids, and the risk of prostate cancer: a nested case-control study. *Cancer Epidemiol Biomarkers Prev.* 2004 Feb;13(2):260-9.

88. Hwang ES, Bowen PE. Cell cycle arrest and induction of apoptosis by lycopene in LNCaP human prostate cancer cells. *J Med Food.* 2004 Fall;7(3):284-9.

89. Siler U, Barella L, Spitzer V, et al. Lycopene and vitamin E interfere with autocrine/paracrine loops in the Dunning prostate cancer model. *FASEB J.* 2004 Jun;18(9):1019-21.

90. Barber NJ, Zhang X, Zhu G, et al. Lycopene inhibits DNA synthesis in primary prostate epithelial cells in vitro and its administration is associated with a reduced prostate-specific antigen velocity in a phase II clinical study. *Prostate Cancer Prostatic Dis.* 2006;9(4):407-13.

91. Ellinger S, Ellinger J, Muller SC, Stehle P. Tomatoes and lycopene in prevention and therapy—is there an evidence for prostate diseases? *Aktuelle Urol.* 2009 Jan;40(1):37-43.

92. Ansari MS, Gupta NP. A comparison of lycopene and orchidectomy vs orchidectomy alone in the management of advanced prostate cancer. *BJU Int.* 2003 Sep;92(4):375-8; discussion 378.

93. Schroder FH, Roobol MJ, Boeve ER, et al. Randomized, double-blind, placebo-controlled crossover study in men with prostate cancer and rising PSA: effectiveness of a dietary supplement. *Eur Urol.* 2005 Dec;48(6):922-30; discussion 930-921.

94. Vaishampayan U, Hussain M, Banerjee M, et al. Lycopene and soy isoflavones in the treatment of prostate cancer. *Nutr Cancer.* 2007;59(1):1-7.

95. Prager N, Bickett K, French N, Marcovici G. A randomized, double-blind, placebo-controlled trial to determine the effectiveness of botanically derived inhibitors of 5-alpha-reductase in the treatment of androgenetic alopecia. *J Altern Complement Med.* 2002 Apr;8(2):143-52.

96. Chittur S, Parr B, Marcovici G. Inhibition of Inflammatory Gene Expression in Keratinocytes Using a Composition Containing Carnitine, Thioctic Acid and Saw Palmetto Extract. *Evid Based Complement Alternat Med.* 2009 Aug 19.

97. Trueb RM. Molecular mechanisms of androgenetic alopecia. *Exp Gerontol.* 2002 Aug-Sep;37(8-9):981-90.

98. Trueb RM. Association between smoking and hair loss: another opportunity for health education against smoking? *Dermatology.* 2003;206(3):189-91.

99. Kaufman KD, Olsen EA, Whiting D, et al. Finasteride in the treatment of men with androgenetic alopecia. Finasteride Male Pattern Hair Loss Study Group. *J Am Acad Dermatol.* 1998 Oct;39(4 Pt 1):578-89.

100. Kaufman KD, Dawber RP. Finasteride, a Type 2 5alpha-reductase inhibitor, in the treatment of men with androgenetic alopecia. *Expert Opin Investig Drugs.* 1999 Apr;8(4):403-15.

101. Ihm CW, Hong SS, Mun JH, Kim HU. Histopathological pictures of the initial changes of the hair bulbs in alopecia areata. *Am J Dermatopathol.* 2004 Jun;26(3):249-53.

102. Vela Navarrete R, Garcia Cardoso JV, Barat A, Manzarbeitia F, López Farré A. BPH and inflammation: pharmacological effects of Permixon on histological and molecular inflammatory markers. Results of a double blind pilot clinical assay. *Eur Urol.* 2003 Nov;44(5):549-55.

103. Aggarwal S, Thareja S, Verma A, Bhardwaj TR, Kumar M. An overview on 5alpha-reductase inhibitors. *Steroids.* 2009 Oct 30.

104. Thüer D, Pfister D, Epplen R, Brehmer B, Heidenreich A. Do alpha-reductase inhibitors prevent prostate cancer? 2008 Practice Guideline from the American Society of Clinical Oncology and American Urological Association. *Pol Arch Med Wewn.* 2009 Oct;119(10):648-53.

105. Kuritzky L. Noninvasive management of lower urinary tract symptoms and sexual dysfunction associated with benign prostatic hyperplasia in the primary care setting. *Compr Ther.* 2005 Fall;31(3):194-208.

106. Rosen RC, Giuliano F, Carson CC. Sexual dysfunction and lower urinary tract symptoms (LUTS) associated with benign prostatic hyperplasia (BPH). *Eur Urol.* 2005 Jun;47(6):824-37.

107. Giuliano F. Impact of medical treatments for benign prostatic hyperplasia on sexual function. *BJU Int.* 2006 Apr;97 Suppl 2:34-8; discussion 44-35.

108. Erdemir F, Harbin A, Hellstrom WJ. 5-alpha reductase inhibitors and erectile dysfunction: the connection. *J Sex Med.* 2008 Dec;5(12):2917-24.

109. Marihart S, Harik M, Djavan B. Dutasteride: a review of current data on a novel dual inhibitor of 5alpha reductase. *Rev Urol.* 2005 Fall;7(4):203-10.

110. Miner M, Rosenberg MT, Perelman MA. Treatment of lower urinary tract symptoms in benign prostatic hyperplasia and its impact on sexual function. *Clin Ther.* 2006 Jan;28(1):13-25.

111. Watson V, Ryan M, Brown CT, Barnett G, Ellis BW, Emberton M. Eliciting preferences for drug treatment of lower urinary tract symptoms associated with benign prostatic hyperplasia. *J Urol.* 2004 Dec;172(6 Pt 1):2321-5.

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References

1. *Int J Impot Res* 2008 Dec; Suppl 2:S9-14.
2. *Int J Impot Res.* 2008 Mar;20(2):173-80.
3. *Phytother Res.* 2009 Mar;23(3):297-302.
4. *J Sex Marital Ther.* 2003 May;29(3):207-13.
5. *European Bulletin of Drug Research.* 2005;13(1): 7-13.
6. *Akush Ginekol (Sofia).* 2007;46(5):7-12.
7. Rohdewald P. Pycnogenol, French maritime pine bark extract. In: Coates P, ed. *Encyclopedia of Dietary Supplements.* New York: Marcel Dekker; 2004.
8. *Urology.* 2006 Dec;68(6):1350-4.

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- **Saw palmetto extract with DeepExtract™.** A number of normal biological events in aging men affect prostate function and structure. Saw palmetto has been shown to interfere with DHT activity in the prostate, inhibit the alpha-adrenergic receptor activity (to support normal urinary flow), and help control inflammatory actions in the prostate gland.^{1,2} **DeepExtract™** is a patented, ultra-high pressure CO₂ technology engineered to recover **far greater amounts of bioactive, high-molecular weight compounds** from the saw palmetto berry during the extraction process. The result is a superior formula containing key ingredients in higher concentrations than many other extracts.³
- **Standardized lignans.** **Enterolactone** has demonstrated anti-estrogen and anti-DHT effects that are of particular importance for the aging prostate gland.⁴⁻⁶ **Lignans** from flax and Norway spruce convert to **enterolactone** in the intestine that is then absorbed into the bloodstream where it provides significant biological effects.
- **5-LOXIN® boswellia extract.** Published studies show that normal aging and poor diet cause levels of a dangerous enzyme (*5-lipoxygenase*) to increase, which can affect prostate cells.⁷ **5-LOXIN®** is a patented extract from the boswellia plant that has been shown to suppress *5-lipoxygenase* in addition to other pro-inflammatory factors.⁸⁻¹²
- **Nettle root extract (*Urtica dioica*).** Testosterone converts to estrogen at higher rates as men age. Prostate cells are sensitive to estrogen's growth stimulatory effects. Nettle root helps support prostate cells against excess estrogen levels.¹³
- **Pygeum.** Normal aging results in levels of certain prostaglandins increasing in the prostate gland. *Pygeum africanum* extract helps suppress these prostaglandins, keeping the prostate gland placid and promoting prostate comfort.¹⁴
- **Beta-sitosterol.** The most biologically active constituent of pygeum is beta-sitosterol. Beta-sitosterol from other plant sources is included in this prostate support formula.¹⁵
- **Cernitin®.** This European pollen extract has been shown to relax smooth muscle tone of the urethra, counteract DHT, and help regulate inflammatory reactions.¹⁵⁻¹⁹



**The daily dose of two softgels of
Ultra Natural Prostate Formula provides:**

USPlus® Saw Palmetto	320 mg	Phytosterol Complex	678 mg
(Co, DeepExtract™)		(standardized to 26.6% beta-sitosterol)	
Nettle root extract	240 mg	Boron	3 mg
(<i>Urtica dioica</i>)		(from citrate, glycinate, and aspartate)	
Pygeum extract	100 mg	Cernitin®	252 mg
(11.7% pygeum sterols)		HMRLignan™	20.15 mg
5-LOXIN®	70 mg	Norway Spruce and	
		ActiFlax™ Flax Lignan extract	
Lycopene extract	10 mg		

- **Boron.** In addition to helping protect bones, this mineral has been shown to slow elevation of prostate-specific antigen (PSA) — a benefit seeming to occur as a result of boron's positive effect on the presence of protein-degrading enzymes in the prostate gland.^{20,21}
- **Lycopene.** This carotenoid associated with tomato's red color helps maintain healthy DNA gene function in prostate cells.²²⁻²⁶

Ultra Natural Prostate Formula provides scientifically validated standardized plant extracts that have been shown to promote healthy prostate function. No other prostate protection formula provides such a broad array of nutrients to support the multiple factors involved in maintaining a healthy prostate gland.

The retail price for 60 softgels of this standardized extract formula is \$38. If a member buys four bottles during **Super Sale**, the price is reduced to **\$23.63** per bottle. If a one-year supply is purchased, the cost drops to only **\$21.60** per bottle.

Not available for export.

References

1. *J Urol.* 2000 May;163(5):1408-12.
2. *Curr Opin Urol.* 2005 Jan;15(1):45-8. Review.
3. Valensa International. Data on file. 2009.
4. *Br J Nutr.* 2007 Aug;98(2):388-96.
5. *Cancer Causes Control.* 2006 Mar;17(2):169-80.
6. *Cancer Epidemiol Biomarkers Prev.* 2005 Jan;14(1):213-20.
7. *Poult Sci.* 2007 May;86(5):1012-6.
8. *Phytother Res.* 2007 Jan;21(1):89-95.
9. *J Nat Prod.* 2000 Aug;63(8):1058-61.
10. *Planta Med.* 2000b Dec;66(8):781-2.
11. *Integr Cancer Ther.* 2002 Mar;1(1):7-37; discussion 37. Review.
12. *Z Naturforsch [C].* 1995 Jan-Feb;50(1-2):98-104.
13. *Curr Med Res Opin.* 1998;14(3):127-39.
14. *World J Urol.* 2002 Apr;19(6):426-35.
15. *Br J Urol.* 1989 Nov;64(5):496-9.
16. *Public Health Nutr.* 2000 Dec;3(4A):459-72.
17. *Planta Med.* 1986 Apr;(2):148-51.
18. *Pharmacometrics.* 1986;31:1-11.
19. *Folia Pharmacol Jpn.* 1988;91:385-92.
20. *Oncol Rep.* 2004 Apr;11(4):887-92.
21. *Toxicol Pathol.* 2004 Jan-Feb;32(1):73-8.
22. *BJU Int.* 2003 Sep;92(4):375-8; discussion 378. Erratum in: *BJU Int.* 2004 Mar;93(4):655.
23. *J Med Food.* 2002 Winter;5(4):181-7.
24. *Exp Biol Med (Maywood).* 2002 Nov;227(10):881-5.
25. *Am J Epidemiol.* 2002 Jun 1;155(11):1023-32.
26. *J Natl Cancer Inst.* 2002 Mar;94(5):391-8.

To order Ultra Natural Prostate Formula, call 1-800-544-4440 or visit www.LifeExtension.com

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SUPER FOODS

BY JON FINKEL

The Raw Power of Strawberries

It's common knowledge that certain berries are nutrition **power-houses**, but when it comes to fighting a vast array of diseases and health problems, few berries stack up to the strawberry in terms of sheer health potency. From combating inflammation and controlling diabetes, to protecting cognitive function and improving heart health, strawberries can bolster your body's health in countless ways.¹

An in vitro cell study using an extract from strawberry leaves on leukemia cells found significant cancer-killing activity, while freeze-dried strawberries slowed the growth of two varieties of cervical cancer cells grown in culture.

Strawberries' Story

Strawberries are one of the most popular berries in the world and are grown primarily in the United States, New Zealand, Japan, Australia, Italy, and Canada. There are over six hundred varieties of strawberry, all of which have the unique appearance of wearing their seeds on the outside instead of on the inside. The most common variety is called the Garden Strawberry, which is the one you are most likely to see in your local grocery store.¹





A Major Source of Antioxidants

At the core of the arsenal of benefits that the strawberry can provide is its dense *phenol* content. The most visible of its phenols are the *anthocyanins*, which give the fruit's flesh its bright red color.² Anthocyanins are widely recognized **antioxidants** that studies have shown help defend cell structures in the body and protect the body's organs. Another phenol found in high doses in strawberries is *ellagitannin*, which has anti-proliferative and antioxidant properties. The combination of anthocyanins and ellagitanins, along with its high vitamin C, folate, and potassium content push it to the top of the list of antioxidant providers among fruit.³ In fact, strawberries are second only to plums as the richest fruit in *phe-*

nols and antioxidants, being especially high in **cancer- and heart disease-fighting flavonoids, quercetin, catechin, and kaempferol.**¹

Anti-Inflammatory Properties

Life Extension Magazine[®] readers are familiar with vitamin C's positive effects when it comes to battling inflammation. While vitamin C is usually associated with orange-colored fruits, readers may be surprised to know that strawberries are a *tremendous source of* vitamin C. In fact, eight strawberries contain more vitamin C than an orange.¹

In a study of 20,000 people published in the *Annals of the Rheumatic Diseases*, subjects who ate the least amount of foods high in vitamin C were more than three times more likely to develop

inflammation leading to arthritis than people who ate the most vitamin C-rich foods.⁴

In addition, strawberries have potent anti-inflammatory properties because the high phenol content decreases the activity of the enzyme *cyclooxygenase*, otherwise known as COX. As a COX-inhibitor, strawberries can offer enhanced protection against *crippling* inflammation-related diseases like rheumatoid arthritis and osteoarthritis.¹

Cancer Protection

Numerous studies have shown that strawberries offer protection against a wide range of cancers, including *breast cancer* and *colon cancer*.⁵ An in vitro cell study using an extract from strawberry leaves

Picking the Perfect Strawberry

Avoid strawberries that are dull or have green or yellow patches. Strawberries don't ripen further once they are picked.

Use your sense of smell when you select strawberries at the store. When strawberries are ready to eat, they will have a flavorful "strawberry" scent.

Select medium-sized strawberries for their optimal flavor.

Ripe strawberries are firm to the touch. Avoid mushy or "soft" strawberries.

If you're selecting pre-packed strawberries, make sure they aren't packed too tightly to avoid buying damaged fruit.

Nutrient	Amount per one cup	Daily Value (%)
Vitamin C	81.65 mg	136.1
Manganese	0.42 mg	21.0
Dietary fiber	3.31 g	13.2
Iodine	12.96 mcg	8.6
Potassium	239.04 mg	6.8
Folate	25.49 mcg	6.4
Vitamin B2 (riboflavin)	0.10 mg	5.9
Vitamin B5 (pantothenic acid)	0.49 mg	4.9
Omega-3 fatty acids	0.11 g	4.6
Vitamin B6 (pyridoxine)	0.08 mg	4.0
Vitamin K	3.17 mcg	4.0
Magnesium	14.40 mg	3.6
Copper	0.07 mg	3.5



on leukemia cells found significant **cancer-killing activity**, while freeze-dried strawberries slowed the growth of two varieties of cervical cancer cells grown in culture. Strawberries, even in freeze-dried *form*, have shown an ability to inhibit tumor growth and to slow the progress of tumors as they begin to develop, according to a study using rats with esophageal cancer.¹

Heart Health

Folate is an integral part of any diet aimed at maintaining optimal heart health. In a study done by the Institute of Biochemistry in Ancona, Italy, researchers found that when subjects consumed strawberries daily for two weeks, an average **3.4%** increase in serum folate levels was observed, mean-

ing regular strawberry intake could have a beneficial effect on folate status and cardiovascular wellness in healthy people.⁶ ●

If you have any questions on the scientific content of this article, please call a Life Extension® Health Advisor at 1-866-864-3027.

References

1. Grotto D. *101 Foods That Could Save Your Life*. New York, New York: Bantam; 2007:313-6.
2. *J Agric Food Chem*. 2006 May 31;54(11):4069-75.
3. *J Agric Food Chem*. 2005 May 18;53(10):4032-40.
4. *Ann Rheum Dis*. 2004 Jul;63(7):843-7.
5. *J Agric Food Chem*. 2006 Feb 22;54(4):1248-55.
6. *Biofactors*. 2008;34(1):47-55.

Advanced Colon Care With **AGAVE** **DIGESTIVE-IMMUNE SUPPORT**

Balancing the ecosystem of the colon is crucial to whole body health. This balance depends on inhibiting **harmful bacteria** and promoting the growth of beneficial bacteria that support immune function. While *stress, diet and age* are important factors, a strong digestive system is *essential* — it accounts for 60% to 70% of our body's immunity.¹ Life Extension®'s **Agave Digestive-Immune Support** helps **beneficial bacteria** proliferate in the colon for healthy immune response.

A clinically *advanced* prebiotic blend, **Agave Digestive-Immune Support** nourishes beneficial bacteria, such as *bifidobacteria*, to *enhance* immune response and suppress growth of unwanted microorganisms. This supports the body's natural response to infection.

Among the several hundred species of bacteria, *bifidobacteria* are among the most essential to colon health. Prebiotics deliver the specific nutrients they need in an *undigested* form. This enables beneficial bacteria to flourish, providing a higher level of protection and support.

AGAVE DIGESTIVE-IMMUNE SUPPORT contains *three* complementary prebiotic ingredients:

AGAVE INULIN/FOS *Inulins* are the nutrients beneficial bacteria thrive on. The inulins in **Agave** are complex. They last longer in the digestive tract, allowing beneficial bacteria to distribute *further* into the colon.

FIBER/SUNFIBER® PARTIALLY HYDROLYZED GUAR GUM (PHGG) favorably modifies intestinal pH for more efficient *absorption* of essential nutrients, including magnesium and calcium.

SHORT-CHAIN FRUCTOOLIGOSACCHARIDES (FOS) are a preferred food source for *bifidobacteria*. FOS also increases stool bulk, supports normal bowel function, and promotes regularity.

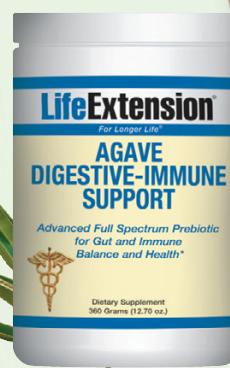
Those who want to obtain the benefits of this *digestion-enhancing* formula can order the new **Agave Digestive-Immune Support**. The **suggested dose** is **1 scoop per day** mixed into 4–8 ounces of any beverage or soft food.

A jar containing **360 grams of powder** retails for \$30, but when a member buys four jars during **Super Sale**, the price is reduced to **\$18.23** per jar.

Reference:

1. *Clin Exp Immunol.* 2008 Sep;153 Suppl 1:3-6.

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WELLNESS PROFILE

BY STEPHEN LAIFER

The Annie Appleseed Foundation

Hope through Alternative Therapies

Doctors don't have all the answers when it comes to battling *lethal* diseases like cancer. Too often, oncologists and other cancer specialists have closed minds when it comes to therapies that might lie outside the bounds of what they learned in medical school textbooks. It's an ironic state of affairs, since physicians—of all people—should understand the fact that there is no one-size-fits-all approach to disease management.

Of course, not all doctors appreciate having their ironclad approaches challenged. For example, one of AAP's underlying tenets is that antioxidants can prove useful in conjunction with conventional therapy.

The Annie Appleseed Project (AAP), based in Delray Beach, Florida, was formed in response to this approach. Anyone who has been diagnosed with cancer knows that it's probably the single most frightening word they will ever hear. Adding to the fear are the sudden confusing decisions that need to be made about which therapies or treatments might be right for their

particular situation. The goal of AAP is to help alleviate some of that fear by providing solid information, advocacy and awareness for people with cancer about complementary and alternative medicine (CAM). Although the project began as AAP founder Ann Fonfa's personal quest to get answers about CAM therapies, it has grown to become a nationally recognized organization.

WELLNESS PROFILE

Seeds Planted

Fonfa was diagnosed with breast cancer in January 1993. Because of chemical sensitivities, she opted not to take chemotherapy—but she knew she had to find some viable alternatives. “Back then, anything that wasn’t chemo or radiation was considered alternative,” Fonfa recalls. “Even yoga was considered dangerous at the time.”

Fonfa quickly agreed to a mastectomy—after three lumpectomies. “In the early 90s, patients typically knew nothing other than that you either kept or lost your breast.” Six months later, Fonfa had recurrence on the chest wall and was considered stage IV. “I objected, since I had no signs of metastatic disease, but was told that was the way it was done.”

After dabbling in a variety of alternative therapies, Fonfa was offered the chance to work with a Chinese herbalist who also had a PhD from Harvard. “This struck me as a good combination: someone with a foot in each camp, so to speak.” Fonfa had meanwhile tried her own brand of CAM by this point: high-dose vitamin A therapy. “By the third week, I noticed the tumors were smaller. I saw an oncologist who, after I told him the news, simply asked me to get dressed. He didn’t even look at the tumors. In later discussions with other oncologists, I found that they already knew about the benefits of vitamin A. I wonder to this day why women aren’t told about this.”

On the path to gaining knowledge for herself, Fonfa encountered others in a similar situation. She decided to found a study group in New York City to which she began inviting speakers, reading books, and obtaining literature

from clinics in places not tied in with the American medical establishment. Eventually, Cancer Care Inc. invited her to talk to their social workers about the patient perspective on alternative medicine. That prompted her to create an outline summarizing what she had learned.

“I realized I had tons of information in the form of summaries I had been making of each meeting I’d attended since 1994,” Fonfa explains. The Annie Appleseed website (www.annieappleseedproject.org) went online in June, 1999. Nowadays, relevant content is added on a weekly basis, currently totaling some 8,000 web pages and links. In addition to reports from conferences worldwide, the site posts studies that help explain the benefits of natural and non-toxic substances. More than two million people have visited in the last twelve months, and thousands

more have heard Fonfa speak at meetings around the US, asking challenging questions that many doctors don’t want to hear.

Seedling Takes Root

“Early funding for AAP came from family, personal friends, other advocates and advocacy groups,” says Fonfa. Even now, she says the organization receives more than 60% of its funding from donations under \$100. Still, growth has been phenomenal: in the last three years, the project has begun offering its own annual educational conference and some foundations have given larger grants. “We have few steady large donors, however,” she laments.

Fonfa still does much of the hands-on work herself. “I have yet to find anyone crazy enough to work full time for no pay,” she says. “We do have terrific volun-



WELLNESS PROFILE

teers who help us with creating print material, distributing materials all over the country, and attending meetings to obtain written summaries so that others can view the information gained." The project also bolsters its primary mission by attending many meetings each year to make its presence known. "It's crucial that the scientific, research and medical community be made aware of our advocacy," Fonfa says. "We speak up a lot, and we ask questions."

Fonfa feels CAM has had a slow road to acceptance in large part because Western science is overly focused on naming and trying to understand exactly why something works. "While we welcome such information, the information itself—the *why*—doesn't change *what* is happening," she points out. "Even if the workings of a natural substance or technique cannot be fully explained, that does not diminish the reality of its effect. From its inception, the Annie Appleseed Project was intended to challenge the existing treatment paradigm." An important part of this is questioning research subjects and methods, proposing instead new directions that take a serious look at CAM and other techniques. "Our hope is that this can ultimately result in a truly integrative oncology."

Of course, not all doctors appreciate having their iron-clad approaches challenged. For example, one of AAP's underlying tenets is that antioxidants can



prove useful in conjunction with conventional therapy. "The preponderance of evidence favors this theory," states Fonfa. "Relatively few oncologists share this perspective, however." She adds that individual patients are still faced with challenging their oncology team if they question what they are told or ask about alternatives. And for those cancer centers that do have integrative sections or offer newer options, programs typically have

to pay out of pocket to utilize the benefits.

Nevertheless, Fonfa sees AAP's ideas slowly gaining ground despite such huge hurdles. "Younger doctors are often more open to CAM, but things change slowly in the medical world," she concedes. "While increasing numbers of hospitals have nutritionists on staff, not everyone gets a recommendation to visit them. But at least yoga is considered a good idea now!" The project intervenes in these areas by suggesting patients download relevant articles and studies from its extensive online library. With these in hand, patients are better equipped to open a possibly difficult dialogue with their doctor.

The goal of AAP is providing solid information, advocacy and awareness for people with cancer about complementary and alternative medicine (CAM).

WELLNESS PROFILE



Tree Bears Fruit

AAP's role as a disseminator of vital information continues to expand. Fonfa has presented posters, papers, and workshops at many conferences worldwide. She was a featured speaker in Hyderabad, India in 2006, and has given presentations at the National Breast Cancer Coalition's annual meeting. AAP has sponsored two national meetings—"Evidence-based Complementary and Alternative Cancer Therapies"—in 2008 and 2009, and has exhibited at 30 other meetings including the American Society for Clinical Oncology, Oncology Nursing Society, Ovarian National Cancer Alliance, Prostate Forum, and American Psychosocial Oncology Society. Even politicians

have been consulting the AAP website as a policy-planning resource.

This year will see the third annual Evidence-Based CAM for Cancer Advocates in January, 2010, to be held in West Palm Beach. Fonfa says the Annie Appleseed Project, as the premier resource on CAM from the patient perspective, is thrilled to be able to offer this educational conference. Patients can meet others who have used complementary or alternative modalities, while practitioners will in turn hear ways to help patients make more informed choices. All attendees will learn how to make healthy lifestyle choices during treatment, after treatment, and into long-term survivorship.

For Fonfa, achieving that survivorship has been a long road of self-discovery. "I took my first dose of Chinese herbs in April 1999. That day I broke out with severe hives. I was sent to a dermatologist, who recommended antihistamines. I woke in the middle of the night with terrible tightness in my chest, which I discovered is a primary possible negative side effect of antihistamines. Since I had specifically mentioned my double mastectomy, I thought the dermatologist might have warned me!"

Fonfa was afraid to touch the herbs again for a solid month. Finally, she called the Chinese herbalist and went forward once more—but first she had an MRI to document the extent of invasive lobular carcinoma in her chest wall. "I made a tea of the herbs from June 1999 until August 2001. I was afraid to stop, even after the herbalist advised me that I was done." Finally, Fonfa had another MRI. "The doctor who was reading it couldn't find anything, and was confused as to why I'd had it done."

Fonfa met with her surgeon on September 12, 2001. "I was living in New York City, and the whole world knows what happened the day before. Fonfa says her doctor told her officially what she really already knew: she was cancer-free. Outside the hospital, Fonfa began to cry, thinking how she had just gotten a reprieve when so many others had died. "Instantly, three women came up to embrace me. They thought I was responding to the visible plume of smoke over lower Manhattan—but here I was, crying because I would live." ●

For more information on the Annie Appleseed Project, visit www.annieappleseedproject.org.



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References: 1. <http://www.womenshealth.gov/faq/varicose-spider-veins.cfm>. 2. *Angiology*. 2003 Jul-Aug;54 Suppl 1:533-44. 3. *Eur J Vasc Endovasc Surg* 2004;28:484-93.

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\$324

FEMALE LIFE EXTENSION PANEL (LC322535)

CBC/Chemistry Profile
DHEA-S
Estradiol (an estrogen)
Homocysteine
C-Reactive Protein (high-sensitivity)
Progesterone
Free Testosterone
Total Testosterone

\$269

FEMALE HORMONE ADD-ON PANEL (LCADDF)

Pregnenolone, Total Estrogens, and TSH
To provide an even more in-depth analysis of a woman's hormone status, Life Extension has created this panel as an addition to the Female Life Extension Panel. This panel provides valuable information about the thyroid, total estrogen status, and the mother hormone that acts as a precursor to all other hormones.

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LIFE EXTENSION FEMALE WEIGHT LOSS PANEL (LCWLF)

CBC/Chemistry Profile, DHEA-S, Free Testosterone, Total Testosterone, Estradiol, Progesterone, C-Reactive Protein, Homocysteine, TSH, Free T4, Free T3

\$324

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TSH, T4, Free T3, Free T4.

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CBC/Chemistry Profile, DHEA-S, Dihydrotestosterone (DHT), Estradiol, PSA, Pregnenolone, Total and Free Testosterone, and TSH. Continual monitoring of hormone levels is necessary for men seeking to maintain optimal blood level values.

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IGF-1, FSH, LH, TSH, T4 free, Cortisol, Fasting Insulin, Sex Hormone Binding Globulin, and Hemoglobin A1C.

\$668

DR. LICHTEN COMPREHENSIVE FEMALE PANEL (LICHTE01F)

FEMALE LIFE EXTENSION PANEL PLUS:
IGF-1, FSH, LH, TSH, T4 free, Cortisol, Fasting Insulin, Sex Hormone Binding Globulin, and Hemoglobin A1C.

\$668

THE CBC/CHEMISTRY PROFILE (LC381822) OVER 40 PARAMETERS TESTED CARDIOVASCULAR RISK PROFILE

Total Cholesterol Cholesterol/HDL Ratio
HDL Cholesterol Estimated CHD Risk
LDL Cholesterol Glucose
Triglycerides Iron
LIVER FUNCTION PANEL
AST (SGOT) Total Bilirubin
ALT (SGPT) Alkaline Phosphatase
LDH

\$35

KIDNEY FUNCTION PANEL

BUN BUN/Creatinine Ratio
Creatinine Uric Acid
BLOOD PROTEIN LEVELS
Total Protein Globulin
Albumin Albumin/Globulin Ratio

BLOOD COUNT/RED AND WHITE BLOOD CELL PROFILE

Red Blood Cell Count Monocytes
White Blood Cell Count Lymphocytes
Eosinophils Platelet Count
Basophils Hemoglobin
Polys (Absolute) Hematocrit
Lymphs (Absolute) MCV
Monocytes (Absolute) MCH
Eos (Absolute) MCHC
Baso (Absolute) Polynucleated Cells
RDW

BLOOD MINERAL PANEL

Calcium Sodium
Potassium Chloride
Phosphorus Iron

NOTE: CBC/Chemistry profile is included in the Male and Female Life Extension panels.

24-HOUR URINARY PROFILES NEW

URINARY COMPLUS PANEL (LCM4080)

\$472.50

This 24-hour urinary steroid hormone profile contains a combination of 30 hormones and metabolites. This panel may be used to evaluate bio-identical hormone replacement therapy.

URINARY COMPREHENSIVE PANEL (LCM4001)

\$360

This 24-hour urinary steroid hormone profile contains a combination of 17 hormones and metabolites. This panel may be used to evaluate bio-identical hormone replacement therapy.

URINARY ADRENAL PROFILE (LCM4000)

\$315

This 24-hour urine test measures a combination of 12 steroid hormones and metabolites synthesized by the adrenal cortex and may prove useful in the evaluation of endocrine disorders, stress, adrenal insufficiency, and more.

URINARY ESTROGEN PROFILE (LCM4061)

\$281.25

This 24-hour urinary steroid hormone profile contains a combination of 8 hormones and metabolites. This panel may be used to evaluate bio-identical hormone replacement therapy.

* This test requires samples to be shipped to the lab on dry ice for customers using a Blood Draw Kit and will incur an additional \$35 charge. If the customer is having blood drawn at a LabCorp facility, this extra charge does not apply.



MOST POPULAR SINGLE TESTS

Life Extension Member Pricing

HORMONES

- CORTISOL (LC004051)**
This test is to measure adrenal function.
- ADRENOCORTICOTROPIC HORMONE (ACTH)* (LC004440)**
A pituitary function test useful in evaluating adrenocortical dysfunction.
- DHEA-SULFATE (LC004697)**
This test shows if you are taking the proper amount of DHEA. This test normally costs \$100 or more at commercial laboratories.
- DIHYDROTESTOSTERONE (DHT)* (LC500142)**
Measures serum concentrations of DHT.
- ESTRADIOL (LC004515)**
For men and women. Determines the proper amount in the body.
- INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 3 (IGFBP3) (LC140152)**
Elevated levels in hypertensive individuals have been associated with a nine-fold increase of carotid arteriosclerosis.
- INSULIN FASTING (LC004333)**
Can predict those at risk of diabetes, obesity, and heart and other diseases.
- PREGNENOLONE* (LC140707)**
Used to determine ovarian failure, hirsutism, adrenal carcinoma, and Cushing's syndrome.
- PROGESTERONE (LC004317)**
Primarily for women. Determines the proper amount in the body.
- SEX HORMONE BINDING GLOBULIN (SHBG) (LC082016)**
This test is used to monitor SHBG levels which are under the positive control of estrogens and thyroid hormones, and suppressed by androgens.
- SOMATOMEDIN C (IGF-1) (LC010363)**
Indicates growth hormone secretion levels. Low levels have been associated with arteriosclerosis as well as all-cause mortality.
- TOTAL AND FREE TESTOSTERONE (LC140103)**
Determines whether testosterone replacement should be considered as a therapy for depression, abdominal obesity, low energy, poor mental performance, or loss of libido.
- URINARY 2/16 ESTROGEN METABOLITE RATIO (LCU73000)**
This ratio is relevant as a risk factor for estrogen-sensitive diseases including breast and cervical cancers, osteoporosis, and recurrent respiratory papillomatosis.

\$39

NEW
\$91

\$61

\$99

\$33

\$47

\$42

\$116

\$55

\$33

\$75

\$99

\$128

CARDIAC RISK

- Lp-PLA2 (PLAC TEST) (LC141275)**
This test is used to aid in predicting risk for coronary heart disease, and ischemic stroke associated with atherosclerosis. Lp-PLA2 is a cardiovascular risk factor that provides unique information about the stability of the plaque inside your arteries.
- C-REACTIVE PROTEIN (HIGH-SENSITIVITY) (LC120766)**
Measures inflammation factors in arteries. Recent studies indicate that C-reactive protein may be the most accurate risk factor for predicting heart attack and stroke.
- APOLIPOPROTEIN ASSESSMENT - (APO A1 + APO B + RATIO) (LC216010)**
This ratio correlates with risk of coronary artery disease and is useful in the presence of borderline elevations of cholesterol.
- FIBRINOGEN* (LC001610)**
High levels of this blood-clotting factor increase the risk of heart attack and stroke.
- HOMOCYSTEINE (LC706994)**
Can indicate if you are likely to have a heart attack or stroke. Even if you take folic acid, you still may have dangerously high levels of this artery-clotting metabolic debris that can be lowered with high doses of TMG and vitamin B6.
- VAP™ TEST (LC804500) (UPDATED)**
The VAP cholesterol test provides a more comprehensive coronary heart disease (CHD) risk assessment than the conventional lipid profile. Direct measurements, not estimations, are provided for total cholesterol, LDL, HDL, VLDL, and cholesterol subclasses.
- MALE HEALTH**
 - PSA (PROSTATE-SPECIFIC ANTIGEN) (LC010322)**
Can provide an early warning sign for prostate disorders and possible cancer.
 - FREE-PSA (INCLUDES TOTAL PSA)* (LC480780)**
Recommended to determine if an elevated PSA is indicative of prostate cancer.
- GENERAL HEALTH**
 - CELIAC DISEASE ANTIBODY SCREEN (LC334971)**
This test measures deamidated gliadin IgA, tissue transglutaminase IgA, and serum IgA.
 - VITAMIN D (250H) (LC081950)**
This test is used to rule out vitamin D deficiency as a cause of bone disease. It can also be used to identify hypercalcemia.
 - VITAMIN C* (LC001479)**
Levels of vitamin C are used to measure nutritional status.
 - FOOD SAFE ALLERGY TEST (LCM73001)**
This test measures delayed (IgG) food allergies for 45 common foods.

\$125

\$42

NEW
\$55

\$31

\$64

\$185

\$31

\$61

\$47

NEW
\$47

\$174

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BLOOD TESTS
FROM VIRTUALLY
ANYWHERE IN THE US!**

TERMS AND CONDITIONS

This blood test service is for informational purposes only and no specific medical advice will be provided. National Diagnostics, Inc., and the Life Extension Foundation contract with a physician who will order your test(s), but will not diagnose or treat you. Both the physician and the testing laboratory are independent contractors and neither National Diagnostics, Inc., nor the Life Extension Foundation® will be liable for their acts or omissions. Always seek the advice of a trained health professional for medical advice, diagnosis, or treatment. When you purchase a blood test from Life Extension/National Diagnostics, Inc., you are doing so with the understanding that you are privately paying for these tests. There will be absolutely no billing to Medicare, Medicaid, or private insurance. I have read the above Terms and Conditions and understand and agree to them.

Signature of Life Extension Member

X

Life Extension Foundation Members only

MEMBER NO.

Male Female

Name

Date of Birth
(required) / /

Address

City

State Zip

Phone

Credit Card No.

Expiration Date /

Mail your order form to:

Life Extension
NATIONAL DIAGNOSTICS, INC.

1100 West Commercial Blvd.,
Ft. Lauderdale, FL 33309

Phone your order to: 1-800-208-3444

Fax your order to: 1-866-728-1050

For non-member pricing call 1-800-208-3444

Products

AMINO ACIDS

Acetyl-L-Carnitine
Acetyl-L-Carnitine-Arginate
Branched Chain Amino Acids
D, L-Phenylalanine Capsules
GABA Powder
Glycine Capsules
Glycine Powder
Arginine Capsules
L-Arginine Free Base Powder
Arginine/L-Ornithine Capsules
L-Carnitine Capsules
L-Carnitine Powder
L-Cysteine Capsules
L-Glutathione, L-Cysteine & C
L-Glutamine Capsules
L-Glutamine Powder
L-Lysine Capsules
L-Lysine Powder
L-Methionine Powder
L-Tyrosine Powder
L-Tyrosine Tablets
Mega L-Glutathione Capsules
N-Acetyl-Cysteine Capsules
Optimized Carnitine with GlycoCarn®
PharmaGABA
Super Carnosine Capsules
Taurine Capsules
Tryptopure® Tryptophan
(Optimized) Tryptopure® Plus

BONE & JOINT HEALTH

ArthroMax™ with Theaflavins
Bone-Up™
Bone Restore™
Bone Strength Formula w/KoAct™
Chondroitin Sulfate
Chondrox
Fast Acting Joint Formula
Glucosamine Chondroitin Capsules

BRAIN HEALTH

Acetyl-L-Carnitine
Acetyl-L-Carnitine-Arginate
CDP Choline Capsules
Cognitex with NeuroProtection Complex
Cognitex with Pregnenolone & NeuroProtection Complex
DMAE
DMAE-Ginkgo
DMAE Powder (37% DMAE)
Ginkgo Biloba Certified Extract™
Hyperzine A
Lecithin with B5 and BHA
Lecithin Granules
Methylcobalamin Lozenges
Optimized Ashwagandha Extract
Phosphatidylserine Capsules
Rhodiola Extract
Super Ginkgo Extract
Vinpocetine

DIGESTIVE

Agave Digestive-Immune Support
Bromelain Powder
Carnosothe w/PicroProtect
Digest RC™
Enhanced Super Digestive Enzymes (also available with Probiotics)
Florastor
Intact Digest
Life Flora™
Natural EsophaGuard
NutraFlora (FOS) Powder
N-Zimes™
Pancreatin
Papain Powder
Primal Defense
Regimint™

DURK AND SANDY PRODUCTS

Blast™
Dual-C
Inner Power™
Memory Upgrade

EYE CARE

Bilberry Extract
Blackcurrant Freeze Dried Extract
Brite Eyes III
Lutein Plus Powder
Overcast Polarized Sunglasses
Solarshield Sunglasses
Super Zeaxanthin with Lutein & Meso-Zeaxanthin
(Plus Astaxanthin also available)
Vision Optimizer

FIBER

Apple Pectin Powder
Chitosan
Enhanced Fiber Food Powder
Hi-Lignan® Nutri-Flax®
SlimStyles® PGX
WellBetX PGX® Soluble Fiber Blend

HAIR CARE

Dr. Proctor's Advanced Hair Formula
Dr. Proctor's Shampoo
Life Extension Shampoo and Conditioner
Super-Absorbable Tocotrienols

HEART HEALTH

Advanced Lipid Control
D-Ribose Capsules
D-Ribose Powder
Endothelial Defense
Enhanced CoQ10 with Brewer's Yeast
Fibrinogen Resist
Forskolin
Homocysteine Resist
Low Dose Aspirin (Enteric Coated)
Natural BP Management
Peak ATP® with GlycoCarn®
Policosanol
Red Yeast Rice
Super Absorbable CoQ10™ with d-Limonene
Super Omega-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract
Super Ubiquinol CoQ10
Super Ubiquinol CoQ10 with Enhanced Mitochondrial™ Support
Strynol™
Theaflavin Standardized Extract
TMG Powder
TMG Tablets
Vascular Protect

HERBAL/PHYTO PRODUCTS

Artichoke Leaf Extract
Astaxanthin
Berry Complete
Blackcurrant Juice Concentrate
Blueberry Extract
Blueberry Extract w/Pomegranate and CocoaGold™
Butterbur Extract w/Standardized Rosmarinic Acid
Calcium D-Glucarate
Chloroplex
Cilantro Herbal Extract
Citrus Bioflavonoid
CocoaGold™
Grapeseed Extract with Resveratrol & Pterostilbene
Green Tea Leaves
Hesperidin Complex Powder
Hyperzine A with Natural Vitamin E
Kyolic® Garlic Formula 105
Kyolic® Reserve
Lycopene
Mega Green Tea Extract
Mega Green Tea Extract (Decaffeinated)
Mega Lycopene
Mega Silymarin with Isosilybin B Nutrim
Optimized Ashwagandha Extract
Phyto-Food
Pomegranate Extract
Pomegranate Extract with CocoaGold™
Pomegranate Juice Concentrate
ProGreens®
Pure-Gar™

Pycnogenol

Optimized Quercetin
Resveratrol with Pterostilbene
Rhodiola Extract
Rosmarinic Acid Extract
Rutin Powder
Silibinin Plus
Silymarin
SODzyme™ with Glisodin®
Stevia Extract
Super Bio-Curcumin®
Super Curcumin with Bioperine®
Super Polyphenol Extracts w/CocoaGold™
Super Ginkgo Extract
Triple Action Cruciferous Vegetable Extract
Venotone
Whole Grape Extract

HORMONES

7-Keto® DHEA
DHEA
DHEA Complete
GH Pituitary Support Day Formula
GH Pituitary Support Night Formula
Melatonin
Melatonin Timed Release
Natural Estrogen with Pomegranate Extract
Pregnenolone
ProFem Cream
Pure IGF
Super Miraforte with Standardized Lignans

IMMUNE ENHANCEMENT

Agave Digestive-Immune Support
AHCC® (Active Hexose Correlated Compound)
Buffered Vitamin C Powder
Echinacea
Enhanced Life Extension Whey Protein
Immune Protect with PARACTIN®
Lactoferrin
Maitake SX-Fraction
Norwegian Shark Liver Oil
Primal Defense™
ProBoost™ Thymic Protein A
Pure Gar™
Sambu® Guard
Theralac Capsules
Thymic Immune Factors
Vitamin C with Dihydroquercetin
Zinc Lozenges with Vitamin C

INFLAMMATORY REACTIONS

Arthro-Immune Joint Support
ArthroMax™ with Theaflavins
Barlean's Kids DHA
Boswella
Boswella™ Topical Cream
Bromelain (Specially-coated)
Coromega Kids Brain and Body (DHA)
DHA 240
Emulsified Norwegian Cod Liver Oil
Fast Acting Joint Formula
Korean Angelica
5-LOXIN®
Mega EPA/DHA
Mega GLA with Sesame Lignans
MSM
Natural Relief 1222™ Cream
NKO Krill Oil
Perilla Oil
Serraflazyme
Shark Cartilage
SODzyme™ with Glisodin® and Wolfberry
Super MaxEPA
Super Omega-3 EPA/DHA with Sesame Lignans & Olive Fruit Extract
Udo's Choice Oil
Zyflamend Easy

LIVER HEALTH

Branch Chain Amino Acids
HepatoPro
Mega Silymarin with Isosilybin B
N-Acetyl Cysteine
SAMe
Silibinin Plus
Silymarin

Products

MINERALS

Biosil
Bone Restore
Bone Strength Formula w/KoAct™
Bone-Up™
Boron Capsules
Calcium Citrate with D3
Chromium Ultra
Copper
Iodoral
Iron Protein Plus
Magnesium
Magnesium Citrate
Mineral Formula for Men
Mineral Formula for Women
Only Trace Minerals
OptiZinc
Sea-Iodine™
Selenium
Se-Methylselenocysteine
Vanadyl Sulfate
Zinc/Vitamin C Lozenges

MISCELLANEOUS

Blood Pressure Monitor Arm Cuff Medium
Cell Sensor Gauss Meter™
Empty Gelatin Capsules
The Capsule Filler Machine
The Pill Cutter and Grinder

MITOCHONDRIAL SUPPORT

Acetyl-L-Carnitine
Acetyl-L-Carnitine-Arginate
ChronoForte with Luteolin
Mitochondrial Energy Optimizer
Optimized Carnitine with GlycoCarn®
Super Absorbable CoQ10™ with d-Limonene
Super Alpha Lipoic Acid with Biotin
Super R-Lipoic Acid
Super Ubiquinol CoQ10 with Enhanced Mitochondrial™ Support

MOOD RELIEF

L-Theanine
Natural Stress Relief
Optimized TryptoPure™ Plus
SAMe
St. John's Wort Extract
Tryptopure™ L-Tryptophan

MOUTH CARE

Advanced Oral Hygiene
Dr. Tung's Tongue Cleaner
Life Extension Mouthwash w/Pomegranate
Life Extension Toothpaste
MistOral III™ with CoQ10

MULTIVITAMIN

Children's Formula Life Extension Mix
Comprehensive Nutrient Pack
Life Extension Booster
Life Extension Mix™ Capsules
Life Extension Mix™ Powder
Life Extension Mix™ Tablets
Life Extension Mix™ w/o Copper Capsules
Life Extension Mix™ w/o Copper Powder
Life Extension Mix™ w/o Copper Tablets
Life Extension Mix™ w/Extra Niacin
Life Extension Mix™ w/Extra Niacin w/o Copper
Life Extension Mix™ w/Stevia Powder
Life Extension Mix™ w/Stevia w/o Copper Powder
Life Extension One-Per-Day
Life Extension Two-Per-Day
Super Booster Softgels w/Advanced K2 Complex
Vital Greens Mix

PET CARE

Life Extension Cat Mix
Life Extension Dog Mix

PROSTATE & URINARY HEALTH

BetterWOMAN®
Cran-Max
5-LOXIN®
(Water-Soluble) Pumpkin Seed Extract

Super Saw Palmetto with Beta-Sitosterol
Super Saw Palmetto/Nettle Root Formula w/Beta-Sitosterol
Ultra Natural Prostate Formula

SKIN CARE

Anti-Aging Mask
Anti-Glycation Serum
Antioxidant Facial Mist
Antioxidant Rejuvenating Foot Cream
Antioxidant Rejuvenating Foot Scrub
Antioxidant Rejuvenating Hand Cream
Antioxidant Rejuvenating Hand Scrub
Anti-Redness & Blemish Lotion
Cellulite Suppress™ Formula
Corrective Clearing Mask
DHEA Skin Creme
Derma Whey
Dual-Action MicroDerm Abrasion
Essential Plant Lipids Reparative Serum
Face Master® Platinum
Face Rejuvenating Antioxidant Cream
Enhanced FernBlock® with Sendara™
Fine Line-Less
Hair Suppress Formula
Healing Mask
Hyaluronic Facial Moisturizer
Hydroderm®
Lavilin Underarm Deodorant
Life Extension Sun Protection Spray
Lifting & Tightening Complex
Mild Facial Cleanser
NaPCA w/Aloe Vera
Neck Rejuvenating Antioxidant Cream
New Face Solution
Peel Off Cleansing Mask
Pigment Correcting Cream
(Ultra) Rejuvenex®
Rejuvenex® Body Lotion
Rejuvenex® Factor
Rejuvenating Serum
Skin Lightening Serum
Skin Stem Cell Serum
Total Sun Protection Cream
Triple Action Under Eye Rejuvenator
Ultra Rejuvenex®
Ultra RejuveNight® w/ Progesterone
Ultra RejuveNight® w/o Progesterone
Ultra Lip Plumper
Ultra Wrinkle Relaxed
Under Eye Refining Serum
Under Eye Rescue Cream
Vitamin C Serum
Vitamin K Healing Cream

SOY

Natural Estrogen w/Pomegranate
Soy Power Powder
Soy Protein Concentrate
Super Absorbable Soy Isoflavones
Ultra Soy Extract

SPECIAL PURPOSE FORMULA

Anti-Alcohol Antioxidants w/HepatoProtection Complex
Benzotiamine w/Thiamine
Breast Health Formula
Butterbur Extract w/Standardized Rosmarinic Acid
Chlorella
Chlorophyllin w/Zinc
CocoaGold w/Beta Glucan
Coriolus Versicolor Super Strength
Enhanced Cinsulin® w/Glucose Management Proprietary Blend
EDTA
(European Leg Solution) Diosmin95
Fem Dophilus
GlucoFit™
Lustre™
Maitake SX-Fraction™
Menopause Solutions
Migr-eze™
Natural Female Support
Pecta-Sol®
Potassium Iodide

Prelox® Natural Sex for Men®
Pyridoxamine
Rosmarinic Acid Extract

SPORTS PERFORMANCE

Enhanced Life Extension Protein
DMG (N, N-dimethylglycine)
Inosine
L-Glutamine Capsules
L-Glutamine Powder
Micronized Creatine Capsules
Micronized Creatine Powder
Octacosanol

VITAMINS

Ascorbic Acid Powder
Ascorbyl Palmitate Capsules
Ascorbyl Palmitate Powder
B1
B2
B12
Beta-Carotene
Biotin Capsules
Biotin Powder
Buffered Vitamin C Powder
Calcium Ascorbate Powder
Complete B Complex
Eligen® B12
Folic Acid + B12
Folic Acid powder
Gamma E Tocopherol w/Sesame Lignans
Gamma E Tocopherol/Tocotrienols
Inositol Capsules
Inositol Powder
Liquid Emulsified Vitamin A
Mega Lycopene Extract
Methylcobalamin
MK-7
No-Flush Niacin
Optimized Folate
PABA Capsules
PABA Powder
Super Ascorbate C Capsules
Super Ascorbate C Powder
Super K w/Advanced K2 Complex
Tocotrienols w/Sesame Lignans
Vitamin B1 Powder
Vitamin B2 Powder
Vitamin B3 (Niacin) Capsules
Vitamin B5 Powder
Vitamin B6
Vitamin B12 Powder
Vitamin C
Vitamin D
Vitamin D3
Vitamin D3 w/Sea-Iodine™
Vitamin E Succinate
Vitamin K1

WEIGHT MANAGEMENT

Alli® Refill Pack
Belly Slim & Tone
Chitosan
CitiChrome
7-Keto DHEA
DHEA Complete
Enhanced Fiber Food
Fucosanthin Slim™
HCA
Integra-Lean® Irvingia
Optimized Irvingia w/Phase 3™ Calorie Control Complex
Life Mix
Natural Appetite Control
SlimStyles® PGX Natural Weight Loss
Stevia Liquid Extract
Super CLA Blend w/Guarana and Sesame Lignans
Super CLA Blend w/Sesame Lignans
Udo's Choice Wholesome Fast Food Blend
WellBetX PGX® Soluble Fiber Blend

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No.		Retail	Member Price	Qty	Total
A					
00449	ACETYL-L-CARNITINE - 500 mg, 100 caps	\$56.00	\$42.00		
	ACETYL-L-CARNITINE - 4 bottles of 100 caps	186.64	139.98		
00788	ACETYL-L-CARNITINE ARGINATE - 100 caps	59.00	44.25		
	ACETYL-L-CARNITINE ARGINATE - 4 bottles of 100 caps	203.96	152.97		
01308	ADVANCED LIPID CONTROL - 60 veg. caps	30.00	22.50		
	ADVANCED LIPID CONTROL - 4 bottles of 60 veg. caps	108.00	81.00		
01300	ADVANCED ORAL HYGIENE - 60 veg. mint lozenges	20.00	15.00		
	ADVANCED ORAL HYGIENE - 4 bottles of 60 veg. mint lozenges	72.00	54.00		
01417	AGAVE DIGESTIVE-IMMUNE SUPPORT - 360 grams of powder	30.00	22.50		
	AGAVE DIGESTIVE-IMMUNE SUPPORT - 4 bottles of powder	108.00	81.00		
00681	AHCC - 500 mg, 30 caps	49.98	37.49		
	AHCC - 4 bottles of 30 caps	180.00	135.00		
46925	ALLI® REFILL PACK - 60 mg, 120 caps	69.95	58.00		
00261	ALL-IN-ONE WEIGHT LOSS FORMULA - 300 caps	59.95	44.96		
	ALL-IN-ONE WEIGHT LOSS FORMULA - 2 bottles of 300 caps	110.00	82.50		
	ALL-IN-ONE WEIGHT LOSS FORMULA - 4 bottles of 300 caps	174.00	130.50		
00457	(SUPER) ALPHA-LIPOIC ACID w/BIOTIN - 250 mg, 60 caps	37.00	27.75		
	(SUPER) ALPHA-LIPOIC ACID w/BIOTIN - 4 bottles of 60 caps	128.00	96.00		
01420	ANTI-ALCOHOL ANTIOXIDANTS W/HEPATOPRO - 100 caps	18.00	13.50		
	ANTI-ALCOHOL ANTIOXIDANTS W/HEPATOPRO - 4 bottles of 100 caps	64.00	48.00		
00105	APPLE PECTIN POWDER - 8 oz bottle of powder	14.75	11.06		
	APPLE PECTIN POWDER - 4 bottles of powder	55.00	41.25		
01012	ARGININE CAPS - 800 mg, 200 caps	26.50	19.88		
	ARGININE CAPS - 4 bottles of 200 caps	93.00	69.75		
01039	ARGININE/ORNITHINE CAPS - 500/250 mg, 100 caps	16.00	12.00		
	ARGININE/ORNITHINE CAPS - 4 bottles of 100 caps	58.00	43.50		
00038	ARGININE/ORNITHINE POWDER - 150 grams of powder	22.95	17.21		
	ARGININE/ORNITHINE POWDER - 4 bottles of powder	76.00	57.00		
01025	(L)-ARGININE FREE-FORM POWDER - 100 grams of powder	18.50	13.88		
	(L)-ARGININE FREE-FORM POWDER - 4 bottles of powder	67.00	50.25		
00040	(L)-ARGININE HCL - 1 kilogram of powder	118.00	88.50		
	(L)-ARGININE HCL - 4 bottles of powder	432.00	324.00		
01303	ARTHROMAX™ w/THEAFLAVINS - 120 veg. caps	44.00	33.00		
	ARTHROMAX™ w/THEAFLAVINS - 4 bottles of 120 veg. caps	160.00	120.00		
01404	ARTHRO-IMMUNE JOINT SUPPORT - 60 veg. caps	30.00	22.50		
	ARTHRO-IMMUNE JOINT SUPPORT - 4 bottles of 60 veg. caps	108.00	81.00		
00919	ARTICHOKE LEAF EXTRACT - 500 mg, 180 veg. caps	28.00	21.00		
	ARTICHOKE LEAF EXTRACT - 4 bottles of 180 veg. caps	101.52	76.14		
00086	(SUPER) ASCORBATE C POWDER - 8 oz of powder	24.95	18.71		
	(SUPER) ASCORBATE C POWDER - 4 bottles of powder	92.52	69.39		
00080	ASCORBIC ACID POWDER - 16 oz of powder vitamin C	38.00	28.50		
	ASCORBIC ACID POWDER - 4 bottles of powder vitamin C	139.72	104.79		
00082	ASCORBYL PALMITATE CAPSULES - 500 mg, 100 caps	22.50	16.88		
	ASCORBYL PALMITATE CAPSULES - 4 bottles of 100 caps	80.00	60.00		
00083	ASCORBYL PALMITATE POWDER - 100 grams of powder	32.00	24.00		
	ASCORBYL PALMITATE POWDER - 4 bottles of powder	120.00	90.00		
00888	(OPTIMIZED) ASHWAGANDHA EXTRACT - 60 veg. caps	10.00	7.50		
	(OPTIMIZED) ASHWAGANDHA EXTRACT - 4 bottles of 60 veg. caps	36.00	27.00		
00892	(LOW DOSE) ASPIRIN ENTERIC COATED - 81 mg, 500 tablets	10.00	7.50		
	(LOW DOSE) ASPIRIN ENTERIC COATED - 4 bottles of 500 tablets	36.00	27.00		
00708	ASTAXANTHIN - 2 mg, 30 softgels	9.98	7.49		

SUB-TOTAL OF COLUMN 1

FEBRUARY 2010

* These products are not 25% off retail price.

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN ADDITIONAL 10% ON ALL PRODUCTS DURING SUPER SALE

No.		Retail	Member Price	Qty	Total
B					
01040	BELLY SLIM & TONE - 6 fl oz	\$42.00	\$31.50		
	BELLY SLIM & TONE - 4 bottles of 6 fl oz	152.00	114.00		
00920	BENFOTIAMINE W/ THIAMINE - 100 mg, 120 caps	19.95	14.96		
	BENFOTIAMINE W/ THIAMINE - 4 bottles of 120 caps	74.40	55.80		
00925	(MEGA) BENFOTIAMINE - 250 mg, 120 veg. caps	30.00	22.50		
	(MEGA) BENFOTIAMINE - 4 bottles of 120 veg. caps	108.00	81.00		
01206	BERRY COMPLETE - 30 veg. caps	21.00	15.75		
	BERRY COMPLETE - 4 bottles of veg. caps	74.68	56.00		
00664	BETA-CAROTENE - 25,000 IU, 100 softgels	10.50	7.88		
	BETA-CAROTENE - 4 bottles of 100 softgels	38.00	28.50		
00088	BETA-CAROTENE POWDER - 100 grams of powder	27.00	20.25		
	BETA-CAROTENE POWDER - 4 bottles of powder	100.00	75.00		
00653	BETTERWOMAN - 40 caps	40.00	30.00		
	BETTERWOMAN - 4 bottles of 40 caps	152.00	114.00		
00497	BILBERRY EXTRACT - 100 mg, 100 caps	28.00	21.00		
	BILBERRY EXTRACT - 4 bottles of 100 caps	100.00	75.00		
*01006	BIOSIL™ - 5 mg, 30 veg. caps	18.95	15.16		
*01007	BIOSIL™ - 30 fl ml	29.99	23.99		
00102	BIOTIN CAPSULES - 600 mcg, 100 caps	7.50	5.63		
	BIOTIN CAPSULES - 4 bottles of 100 caps	26.00	19.50		
00145	BIOTIN POWDER - 30 grams of powder	12.50	9.38		
	BIOTIN POWDER - 4 bottles of powder	44.00	33.00		
01037	BLACKCURRANT FREEZE DRIED EXTRACT - 60 veg. caps	21.95	16.46		
01034	BLACKCURRANT JUICE CONCENTRATE - 12 fl. oz	17.95	13.46		
01008	BLAST™ - 600 grams of powder	26.95	20.21		
70000	BLOOD PRESSURE MONITOR-ARM CUFF (medium)	99.95	64.97		
70001	BLOOD PRESSURE MONITOR-WRIST (travel size)	69.95	45.47		
01214	BLUEBERRY EXTRACT - 60 veg. caps	22.50	16.88		
	BLUEBERRY EXTRACT - 4 bottles of 60 veg. caps	80.00	60.00		
01338	BLUEBERRY EXTRACT w/POMEGRANATE & COCOAGOLD™ - 60 veg. caps	30.00	22.50		
	BLUEBERRY EXTRACT w/POMEGRANATE & COCOAGOLD™ - 4 bottles of 60 veg. caps	108.00	81.00		
00811	BONE RESTORE - 150 caps	22.50	16.88		
	BONE RESTORE - 4 bottles of 150 caps	78.00	58.50		
01211	BONE STRENGTH FORMULA w/KOACT™ - 120 caps	42.00	31.50		
	BONE STRENGTH FORMULA w/KOACT™ - 4 bottles of 120 caps	152.00	114.00		
00313	BONE-UP® - 240 caps (hydroxyapatite complex)	24.95	18.71		
	BONE-UP® - 4 bottles of 240 caps	90.00	67.50		
01079	BOOSTER - 60 softgels	48.00	36.00		
	BOOSTER - 4 bottles of 60 softgels	176.00	132.00		
01380	(SUPER) BOOSTER w/ADVANCED K2 COMPLEX - 60 softgels	42.00	31.50		
	(SUPER) BOOSTER w/ADVANCED K2 COMPLEX - 4 bottles of 60 softgels	152.00	114.00		
00621	BORON CAPS - 3 mg, 100 caps	5.95	4.46		
	BORON CAPS - 4 bottles of 100 caps	21.00	15.75		
00202	BOSWELLA - 100 caps	38.00	28.50		
	BOSWELLA - 4 bottles of 100 caps	120.00	90.00		
00258	BOSWELLA TOPICAL CREAM - 4 oz jar	15.00	11.25		
	BOSWELLA TOPICAL CREAM - 4 jars	52.00	39.00		
00253	BRANCHED CHAIN AMINO ACIDS - 90 caps	19.50	14.63		
	BRANCHED CHAIN AMINO ACIDS - 4 bottles of 90 caps	68.00	51.00		
00999	BREAST HEALTH FORMULA - 60 veg. caps	34.00	25.50		
	BREAST HEALTH FORMULA - 4 bottles of 60 veg. caps	120.00	90.00		
00893	BRITE EYES III - 2 vials, 5 ml each	34.00	25.50		
	BRITE EYES III - 4 boxes	128.00	96.00		

SUB-TOTAL OF COLUMN 2

OFFER ENDS FEBRUARY 1, 2010

To order online visit www.lifeextension.com/SuperSale

No.		Retail	Member Price	Qty	Total
00136	BROMELAIN POWDER - 100 grams of powder	\$21.00	\$15.75		
	BROMELAIN POWDER - 4 bottles of powder	75.00	56.25		
01203	(SPECIALY COATED) BROMELAIN - 500 mg, 60 enteric coated tablets	21.00	15.75		
	(SPECIALY COATED) BROMELAIN - 4 bottles of 60 tablets	76.00	57.00		
00884	BUTTERBUR EXT. W/ STANDARDIZED ROSMARINIC ACID - 60 softgels	44.00	33.00		
	BUTTERBUR EXT. W/ STANDARDIZED ROSMARINIC ACID - 4 bottles of 60 softgels	158.40	118.80		
C					
00146	CALCIUM ASCORBATE POWDER - 300 grams of powder	\$24.00	\$18.00		
	CALCIUM ASCORBATE POWDER - 4 bottles of powder	88.00	66.00		
00058	CALCIUM CARBONATE POWDER - 1 kilogram of powder	13.95	10.46		
	CALCIUM CARBONATE POWDER - 4 bottles of powder	52.00	39.00		
01015	CALCIUM CITRATE - 300 caps	24.00	18.00		
	CALCIUM CITRATE - 4 bottles of 300 caps	85.00	63.75		
00535	CALCIUM D-GLUCARATE - 200 mg, 60 caps	18.00	13.50		
	CALCIUM D-GLUCARATE - 4 bottles of 60 caps	60.00	45.00		
00149	CALCIUM/MAGNESIUM POWDER - 1 kilogram of powder	21.00	15.75		
	CALCIUM/MAGNESIUM POWDER - 4 bottles of powder	75.00	56.25		
01419	CALORIE RESTRICTION MIMETIC FORMULA - 60 veg. caps	36.00	27.00		
	CALORIE RESTRICTION MIMETIC FORMULA - 4 bottles of 60 veg. caps	132.00	99.00		
00118	CAPSULE FILLER MACHINE FOR "00" CAPSULES	12.00	9.00		
	CAPSULE FILLER MACHINE FOR "00" CAPSULES - 4 machines	45.00	33.75		
00613	CAPSULE FILLER MACHINE FOR "0" CAPSULES	12.00	9.00		
	CAPSULE FILLER MACHINE FOR "0" CAPSULES - 4 machines	45.00	33.75		
00916	(OPTIMIZED) CARNITINE w/ GLYCOCARN® - 60 veg. caps	29.00	21.75		
	(OPTIMIZED) CARNITINE w/ GLYCOCARN® - 4 bottles of 60 veg. caps	104.40	78.30		
01041	L-CARNITINE CAPSULES - 500 mg, 30 caps	15.00	11.25		
	L-CARNITINE CAPSULES - 4 bottles of 30 caps	52.80	39.60		
00051	L-CARNITINE POWDER - 100 grams of powder	85.00	63.75		
	L-CARNITINE POWDER - 4 bottles of powder	295.00	221.25		
01258	CARNOSOOTHE w/ PICROPROTECT™ - 60 veg. caps	29.95	22.46		
	CARNOSOOTHE w/ PICROPROTECT™ - 4 bottles of 60 veg. caps	108.00	81.00		
01287	(SUPER) CARNOSINE CAPS - 500 mg, 90 caps	66.00	49.50		
	(SUPER) CARNOSINE CAPS - 4 bottles of 90 caps	240.00	180.00		
01011	CELLULITE SUPPRESS™ FORMULA - 6 fl. oz	58.00	43.50		
	CELLULITE SUPPRESS™ FORMULA - 4-6oz bottles	208.00	156.00		
00390	CDP CHOLINE CAPS - 250 mg, 60 caps	36.00	27.00		
	CDP CHOLINE CAPS - 4 bottles of 60 caps	136.00	102.00		
00998	CELL SENSOR-EMF DETECTION/MEASUREMENT METER	39.95	29.95		
01370	CHILDREN'S FORMULA MIX - 100 chewable tablets	18.00	13.50		
	CHILDREN'S FORMULA MIX - 4 bottles of 100 chewable tablets	64.00	48.00		
00216	CHITOSAN - 250 mg, 180 caps	28.00	21.00		
	CHITOSAN - 4 bottles of 180 caps	100.00	75.00		
	CHITOSAN - 10 bottles of 180 caps	230.00	172.50		
00369	CHITOSAN - 500 mg, 180 caps	55.00	41.25		
	CHITOSAN - 4 bottles of 180 caps	200.00	150.00		
00550	CHLORELLA - 500 mg, 200 tablets	23.50	17.63		
	CHLORELLA - 4 bottles of 200 tablets	84.00	63.00		
01048	CHLOROPHYLLIN w/ZINC - 100 mg, 100 caps	24.00	18.00		
	CHLOROPHYLLIN w/ZINC - 4 bottles of 100 caps	80.00	60.00		
01045	CHLOROPLEX - 100 caps	28.00	21.00		
	CHLOROPLEX - 4 bottles of 100 caps	98.00	73.50		
	CHLOROPLEX - 12 bottles of 100 caps	256.00	192.00		
00142	CHOLINE BITARTRATE - 300 grams of powder	17.75	13.31		
	CHOLINE BITARTRATE - 4 bottles of powder	64.00	48.00		

SUB-TOTAL OF COLUMN 3

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

Buyers Club Order Form

No.		Retail	Member Price	Qty	Total
00541	CHOLINE CHLORIDE - 16 oz liquid	\$14.95	\$11.21		
	CHOLINE CHLORIDE - 4 bottles of 16 oz liquid	56.00	42.00		
00364	CHONDROITIN SULFATE - 400 mg, 60 tablets	19.95	14.96		
	CHONDROITIN SULFATE - 4 bottles of 60 tablets	72.00	54.00		
00169	CHROMIUM - 200 mcg polynicotinate, 365 caps	28.00	21.00		
	CHROMIUM - 4 bottles of 365 caps	104.00	78.00		
01071	CHROMIUM ULTRA - 100 veg. caps	24.00	18.00		
	CHROMIUM ULTRA - 4 bottles of 100 veg. caps	84.00	63.00		
00680	CHRONOFORTE WITH LUTEOLIN - 180 caps	96.00	72.00		
	CHRONOFORTE WITH LUTEOLIN - 4 bottles of 180 caps	354.00	265.50		
00551	CILANTRO HERBAL EXTRACT - 1 oz	12.00	9.00		
	CILANTRO HERBAL EXTRACT - 4 bottles of 1oz	44.00	33.00		
01267	(ENHANCED) CINSULIN® W/ GLUCOSE MGMT PROP BLEND - 90 veg. caps	32.00	24.00		
	(ENHANCED) CINSULIN® W/ GLUCOSE MGMT PROP BLEND - 4 bottles of 90 veg. caps	120.00	90.00		
00223	CITRICHROME - 90 caps	22.95	17.21		
	CITRICHROME - 4 bottles of 90 caps	80.00	60.00		
00069	CITRUS BIOFLAVONOID - 100 caps	14.95	11.21		
	CITRUS BIOFLAVONOID - 4 bottles of 100 caps	56.00	42.00		
00818	(SUPER) CLA BLEND W/ SESAME LIGNANS - 1000 mg, 120 softgels	36.00	27.00		
	(SUPER) CLA BLEND w/ SESAME LIGNANS - 4 bottles of 120 softgels	132.00	99.00		
	(SUPER) CLA BLEND w/ SESAME LIGNANS - 10 bottles, 120 softgels	263.30	197.48		
00819	(SUPER) CLA BLEND w/ GUARANA & SESAME - 120 softgels	42.00	31.50		
	(SUPER) CLA BLEND w/ GUARANA & SESAME - 4 btl of 120 softgels	153.32	114.99		
00863	COCOA GOLD™ - 130 mg, 60 veg. caps	12.50	9.38		
	COCOA GOLD™ - 4 bottles of 60 veg. caps	44.00	33.00		
01285	COCOA GOLD™ w/ BETA-GLUCAN POWDER - 180 grams	20.00	15.00		
	COCOA GOLD™ w/ BETA-GLUCAN POWDER - 4 jars of 180 grams	72.00	54.00		
00050	COD LIVER OIL - 12 fl oz (355 ml) (Emulsified)	9.50	7.13		
	COD LIVER OIL - 4 bottles of 12 fl oz (355 ml)	32.00	24.00		
00293	COD LIVER OIL - 100 softgels (Emulsified)	9.95	7.46		
	COD LIVER OIL - 4 bottles of 100 softgels	37.52	28.14		
00922	COGNITEX w/ PREGNENOLONE & NEUROPROTECTION COMPLEX - 90 softgels	74.00	55.50		
	COGNITEX w/ PREGNENOLONE & NEUROPROTECTION COMPLEX - 4 btl of 90 softgels	266.40	199.80		
	COGNITEX w/ PREGNENOLONE & NEUROPROTECTION COMPLEX - 12 btl of 90 softgels	768.00	576.00		
00921	COGNITEX w/o PREGNENOLONE & NEUROPROTECTION COMPLEX - 90 softgels	72.00	54.00		
	COGNITEX w/o PREGNENOLONE & NEUROPROTECTION COMPLEX - 4 bottles of 90 softgels	256.00	192.00		
	COGNITEX w/o PREGNENOLONE & NEUROPROTECTION COMPLEX - 12 bottles of 90 softgels	744.00	558.00		
00110	COMPLETE B-COMPLEX - 180 caps	21.50	16.13		
	COMPLETE B-COMPLEX - 4 bottles of 180 caps	72.00	54.00		
01325	COMPREHENSIVE NUTRIENT PACK - 30-day supply	189.00	141.75		
	COMPREHENSIVE NUTRIENT PACK - 120-day supply	684.00	513.00		
00119	COPPER CAPSULES - 2 mg, 100 caps	8.95	6.71		
	COPPER CAPSULES - 4 bottles of 100 caps	32.36	24.27		
00208	(ENHANCED) COQ10 (with Brewer's Yeast) - 30 mg, 100 caps	32.00	24.00		
	(ENHANCED) COQ10 (with Brewer's Yeast) - 4 bottles of 100 caps	108.00	81.00		
	(ENHANCED) COQ10 (with Brewer's Yeast) - 10 bottles of 100 caps	245.00	183.75		
00949	(SUPER ABSORBABLE) COQ10™ w/ d-LIMONENE - 50 mg, 60 softgels	28.50	21.38		
	(SUPER ABSORBABLE) COQ10™ w/ d-LIMONENE - 4 bottles of 60 softgels	99.00	74.25		
	(SUPER ABSORBABLE) COQ10™ w/ d-LIMONENE - 10 bottles of 60 softgels	230.00	172.50		
00950	(SUPER ABSORBABLE) COQ10™ w/ d-LIMONENE - 100 mg, 100 softgels	73.00	54.75		
	(SUPER ABSORBABLE) COQ10™ w/ d-LIMONENE - 4 bottles of 100 softgels	260.00	195.00		
	(SUPER ABSORBABLE) COQ10™ W/ D-LIMONENE - 10 bottles of 100 softgels	620.00	465.00		

SUB-TOTAL OF COLUMN 4

FEBRUARY 2010

Buyers Club Order Form

SUPER SALE SAVINGS ON ALL PRODUCTS
To order call: 1-954-766-8433 or 1-800-544-4440

No.		Retail	Member Price	Qty	Total
C CONTINUED					
01226	(SUPER UBIQUINOL) COQ10 - 100 mg, 60 softgels	\$62.00	\$46.50		
	(SUPER UBIQUINOL) COQ10 - 4 bot of 60 softgels	224.00	168.00		
	(SUPER UBIQUINOL) COQ10 - 10 bot of 60 softgels	520.00	390.00		
01426	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 100 mg, 60 softgels	62.00	46.50		
	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 4 bot of 60 softgels	224.00	168.00		
	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 10 bot of 60 softgels	520.00	390.00		
01425	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 50 mg, 100 softgels	58.00	43.50		
	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 4 bot of 100 softgels	212.00	159.00		
	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 10 bot of 100 softgels	500.00	375.00		
01427	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 50 mg, 30 softgels	20.00	15.00		
	(SUPER UBIQUINOL) COQ10 w/ENH MITOCHONDRIAL SUPPORT™ - 4 bot of 30 softgels	72.00	54.00		
01053	CORIOLUS VERSICOLOR SUPER STRENGTH - 600 mg, 150 caps	99.95	74.96		
00857	COROMEGA CHILD BRAIN & BODY - (Lemon Lime) 30 packets	25.00	18.75		
	COROMEGA CHILD BRAIN & BODY - (Lemon Lime) 4 boxes of 30 packets	90.00	67.00		
80118	COSMESIS ANTI-AGING MASK - 2 oz bottle	66.60	49.95		
	COSMESIS ANTI-AGING MASK - 2-2 oz bottles	117.22	87.92		
80134	COSMESIS ANTI-GLYCATION SERUM - 1 oz bottle w/BLUEBERRY & POMEGRANATE EXTRACTS	24.00	18.00		
	COSMESIS ANTI-GLYCATION SERUM - 2-1 oz bottles w/BLUEBERRY & POMEGRANATE EXTRACTS	45.60	34.20		
80133	COSMESIS ANTI-OXIDANT FACIAL MIST - 2 oz bottle	29.95	22.46		
	COSMESIS ANTI-OXIDANT FACIAL MIST - 2-2 oz bottles	56.90	42.68		
80132	COSMESIS ANTI-OXIDANT FACIAL MIST - 4 oz bottle	39.95	29.96		
	COSMESIS ANTI-OXIDANT FACIAL MIST - 2-4 oz bottles	76.00	57.00		
80127	COSMESIS ANTI-OXIDANT REJUVENATING FOOT CREAM - 2 oz jar	39.95	29.96		
	COSMESIS ANTI-OXIDANT REJUVENATING FOOT CREAM - 2-2 oz jar	76.00	57.00		
80128	COSMESIS ANTI-OXIDANT REJUVENATING FOOT SCRUB - 2 oz jar	52.95	39.71		
	COSMESIS ANTI-OXIDANT REJUVENATING FOOT SCRUB - 2-2 oz jar	93.20	69.90		
80117	COSMESIS ANTI-OXIDANT REJUVENATING HAND CREAM - 2 oz jar	59.00	44.25		
	COSMESIS ANTI-OXIDANT REJUVENATING HAND CREAM - 2-2 oz jars	106.00	79.50		
80121	COSMESIS ANTI-OXIDANT REJUVENATING HAND SCRUB - 2 oz jar	52.95	39.71		
	COSMESIS ANTI-OXIDANT REJUVENATING HAND SCRUB - 2-2 oz jars	93.20	69.90		
80105	COSMESIS ANTI-REDNESS & BLEMISH LOTION - 1 oz	69.95	52.46		
	COSMESIS ANTI-REDNESS & BLEMISH LOTION - 2-1 oz bottles	123.12	92.34		
80120	COSMESIS CORRECTIVE CLEARING MASK - 2 oz jar	59.95	44.96		
	COSMESIS CORRECTIVE CLEARING MASK - 2-2 oz jars	105.52	79.14		
80108	COSMESIS ESSENTIAL PLANT LIPIDS REPARATIVE SERUM - 1 oz	71.60	53.70		
	COSMESIS ESSENTIAL PLANT LIPIDS REPARATIVE SERUM - 2-1 oz bottles	126.00	94.50		
80123	COSMESIS FACE REJUVENATING ANTIOXIDANT CREAM - 2 oz jar	65.00	48.75		
	COSMESIS FACE REJUVENATING ANTIOXIDANT CREAM - 2-2 oz jars	114.40	85.80		
80107	COSMESIS FINE LINE-LESS - 1 oz	69.95	52.46		
	COSMESIS FINE LINE-LESS - 2-1 oz bottles	123.12	92.34		
80131	COSMESIS HAIR SUPPRESS FORMULA - 4 oz bottle	53.75	40.31		
	COSMESIS HAIR SUPPRESS FORMULA - 2-4 oz bottles	94.60	70.95		
80115	COSMESIS HEALING MASK - 2 oz bottle	59.95	44.96		
	COSMESIS HEALING MASK - 2-2 oz bottles	105.52	79.14		
80109	COSMESIS HYALURONIC FACIAL MOISTURIZER - 1 oz	52.95	39.71		
	COSMESIS HYALURONIC FACIAL MOISTURIZER - 2 bottles of 1 oz	93.20	69.90		
80110	COSMESIS HYALURONIC OIL-FREE FACIAL MOISTURIZER - 1 oz	52.95	39.71		
	COSMESIS HYALURONIC OIL-FREE FACIAL MOISTURIZER - 2 btl of 1 oz	93.20	69.90		
80103	COSMESIS LIFTING & TIGHTENING COMPLEX - 1 oz	69.95	52.46		
	COSMESIS LIFTING & TIGHTENING COMPLEX - 2 bottles of 1 oz	123.12	92.34		
80114	COSMESIS MILD FACIAL CLEANSER - 8 oz	52.95	39.71		
	COSMESIS MILD FACIAL CLEANSER - 2 bottles of 8 oz	93.20	69.90		

SUB-TOTAL OF COLUMN 5

FEBRUARY 2010

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

No.		Retail	Member Price	Qty	Total
D					
01313	DERMA WHEY- 400 mg, 60 caps		\$65.00	\$48.75	
	DERMA WHEY- 4 bottles of 60 caps		236.00	177.00	
00856	(BARLEAN'S KID'S) DHA (Fruit Punch Flavor)- 8 fl. oz		22.00	16.50	
	(BARLEAN'S KID'S) DHA (Fruit Punch Flavor)- 4 bottles of 8 fl. oz		80.00	60.00	
00995	DHA 240 - 240 mg, 60 softgels		19.00	14.25	
	DHA 240 - 4 bottles of 60 softgels		68.00	51.00	
00658	7-KETO® DHEA METABOLITE - 25 mg, 100 caps		28.00	21.00	
	7-KETO® DHEA METABOLITE - 4 bottles of 100 caps		96.00	72.00	
01271	7-KETO® DHEA METABOLITE - 100 mg, 60 veg. caps		40.00	30.00	
	7-KETO® DHEA METABOLITE - 4 bottles of 60 veg. caps		144.00	108.00	
00607	DHEA - 25 mg dissolve in mouth 100 tablets pharmaceutical-grade		14.00	10.50	
	DHEA - 4 bottles of 100 tablets pharmaceutical-grade		47.00	35.25	
01250	DHEA COMPLETE - 60 veg. caps		48.00	36.00	
	DHEA COMPLETE - 4 bottles of 60 veg. caps		172.80	129.60	

SUB-TOTAL OF COLUMN 6

OFFER ENDS FEBRUARY 1, 2010

To order online visit www.lifeextension.com/SuperSale

No.		Retail	Member Price	Qty	Total
00335	DHEA - 25 mg, 100 caps	\$15.00	\$11.25		
	DHEA - 4 bottles of 100 caps	50.00	37.50		
00454	DHEA - 15 mg, 100 caps	12.00	9.00		
	DHEA - 4 bottles of 100 caps	40.00	30.00		
00882	DHEA - 50 mg, 60 caps	16.00	12.00		
	DHEA - 4 bottles of 60 caps	56.00	42.00		
00883	DHEA - 100 mg, 60 caps	22.50	16.88		
	DHEA - 4 bottles of 60 caps	76.00	57.00		
01058	DHEA SKIN CREME - 1.7 oz pump	29.99	22.49		
01358	DIGEST RC - 30 tablets	19.95	14.96		
	DIGEST RC - 4 boxes of 30 tablets	68.00	51.00		
01272	(ENHANCED SUPER) DIGESTIVE ENZYMES - 100 veg. caps	18.95	14.21		
	(ENHANCED SUPER) DIGESTIVE ENZYMES - 4 bottles of 100 veg. caps	64.00	48.00		
01273	(ENHANCED SUPER) DIGESTIVE ENZYMES w/PROBIOTICS - 100 veg. caps	24.00	18.00		
	(ENHANCED SUPER) DIGESTIVE ENZYMES w/PROBIOTICS - 4 bottles of 100 veg. caps	88.00	66.00		
00114	DILAURYL THIODIPROPIONATE POWDER - 100 grams of powder	22.00	16.50		
	DILAURYL THIODIPROPIONATE POWDER - 4 bottles of powder	80.00	60.00		
**01042	(EUROPEAN LEG SOLUTION) DIOSMIN95 - 600 mg, 30 veg. tabs	20.00	15.00		
	(EUROPEAN LEG SOLUTION) DIOSMIN95 - 4 bottles of 30 veg. tabs	72.00	54.00		
00034	D,L-PHENYLALANINE CAPSULES - 500 mg, 100 caps	18.75	14.06		
	D,L-PHENYLALANINE CAPSULES - 4 bottles of 100 caps	64.00	48.00		
00033	D,L-PHENYLALANINE PLUS POWDER - 100 grams of powder	26.00	19.50		
	D,L-PHENYLALANINE PLUS POWDER - 4 bottles of powder	86.00	64.50		
00257	DMAE BITARTRATE - 150 mg, 200 caps	14.00	10.50		
	DMAE BITARTRATE - 4 bottles of 200 caps	42.00	31.50		
00022	DMAE POWDER (37% DMAE) - 100 grams of powder	24.00	18.00		
	DMAE POWDER (37% DMAE) - 4 bottles of powder	80.00	60.00		
00197	DMAE-GINKGO CAPSULES - 100 caps	36.00	27.00		
	DMAE-GINKGO CAPSULES - 4 bottles of 100 caps	120.00	90.00		
	DMAE-GINKGO CAPSULES - 8 bottles of 100 caps	220.00	165.00		
00059	DMG - 125 mg dimethylglycine, 60 tablets	18.50	13.88		
	DMG - 4 boxes of 60 tablets	68.00	51.00		
01052	DNA PROTECTION FORMULA - 60 veg. caps	28.00	21.00		
	DNA PROTECTION FORMULA - 4 bottles of 60 veg. caps	100.80	75.60		
00321	DR. PROCTOR'S ADVANCED HAIR FORMULA - 2 oz	39.95	29.96		
	DR. PROCTOR'S ADVANCED HAIR FORMULA - 4 bottles	128.00	96.00		
00320	DR. PROCTOR'S HAIR FORMULA SHAMPOO - 8 oz bottle	24.95	18.71		
	DR. PROCTOR'S HAIR FORMULA SHAMPOO - 4 bottles	88.00	66.00		
00376	DR. TUNG'S TONGUE CLEANER	8.45	6.34		
	DR. TUNG'S TONGUE CLEANERS - 4 quantity	32.20	24.15		
00899	DUAL-ACTION MICRODERMABRASION ADV. EXFOLIATE - 2.4 net oz jar	39.95	29.96		
	DUAL-ACTION MICRODERMABRASION ADV. EXFOLIATE - 4 jars of 2.4 net oz	155.80	116.85		
00240	DUAL-C - 90 caps	12.00	9.00		
	DUAL-C - 4 bottles of 90 caps	42.00	31.50		
E					
00388	ECHINACEA - 250 mg, 60 caps	\$14.35	\$10.76		
	ECHINACEA - 4 bottles of 60 caps	50.00	37.50		
01024	EDTA - 500 mg, 100 caps	17.50	13.13		
01403	ELIGEN® B12 - 100 mcg 30 veg tablets	28.00	21.00		
	ELIGEN® B12 - 4 bottles of 30 veg tablets	100.00	75.00		
01297	ENDOTHELIAL DEFENSE™ w/GLISODIN® and COCOAGOLD™ - 60 veg. caps	54.00	40.50		
	ENDOTHELIAL DEFENSE™ w/GLISODIN® AND COCOAGOLD™ - 4 bottles of 60 veg. caps	192.00	144.00		

* Due to license restrictions, this product is not for sale to wholesalers outside of the United States of America and Canada.

** Can only be sold in the United States and cannot be sold into the health food retail store channel.

† Member pricing not valid on this item.

SUB-TOTAL OF COLUMN 7

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10% ON ALL PRODUCTS DURING SUPER SALE**

Buyers Club Order Form

No.		Retail	Member Price	Qty	Total
00625	(MEGA) EPA/DHA - 120 softgels	\$19.95	\$14.96		
	(MEGA) EPA/DHA - 4 bottles of 120 softgels	72.00	54.00		
F					
†01054	FACE MASTER® PLATINUM	\$199.00	\$199.00		
00965	FAST-ACTING JOINT FORMULA - 30 caps	39.00	29.25		
	FAST-ACTING JOINT FORMULA - 4 bottles of 30 caps	144.00	108.00		
20053	FEM DOPHILUS® - 30 caps	24.95	18.71		
20055	FEM DOPHILUS® - 60 caps	35.95	26.96		
*01311	(ENHANCED) FERNBLOCK® w/SENDARA™ - 30 veg. caps	39.00	29.25		
	(ENHANCED) FERNBLOCK® w/SENDARA™ - 4 bottles of 30 veg. caps	140.00	105.00		
00229	FIBER FOOD CAPS - 200 caps	15.00	11.25		
	FIBER FOOD CAPS - 4 bottles of 200 caps	50.00	37.50		
	FIBER FOOD CAPS - 10 bottles of 200 caps	119.00	89.25		
00228	FIBER FOOD - 300 grams of powder	15.00	11.25		
	FIBER FOOD - 4 bottles of powder	50.00	37.50		
	FIBER FOOD - 10 bottles of powder	119.00	89.25		
01307	(ENHANCED) FIBER FOOD POWDER - 453 grams of powder	34.00	25.50		
	(ENHANCED) FIBER FOOD POWDER - 4 jars of powder	120.00	90.00		
00718	FIBRINOGEN RESIST FORMULA (NATTOKINASE) - 30 veg. caps	49.00	36.75		
	FIBRINOGEN RESIST FORMULA (NATTOKINASE) - 4 bottles of 30 caps	176.00	132.00		
00873	FLAXSEED POWDER (HI-LIGNAN NUTRI-FLAX) - 16 oz powder	9.95	7.46		
01200	FLORASTOR® - 50 caps	55.15	41.36		
01312	(OPTIMIZED) FOLATE (5-METHYLTETRAHYDROFOLATE) - 100 veg. caps	28.00	21.00		
	(OPTIMIZED) FOLATE (5-METHYLTETRAHYDROFOLATE) - 4 bottles of 100 veg. caps	100.00	75.00		
00347	FOLIC ACID + B12 CAPSULES - 200 caps	10.50	7.88		
	FOLIC ACID + B12 CAPSULES - 4 bottles of 200 caps	38.00	28.50		
00148	FOLIC ACID POWDER - 30 grams of powder	16.00	12.00		
	FOLIC ACID POWDER - 4 bottles of powder	55.00	41.25		
00300	FORSKOLIN - 10 mg, 60 caps of elemental forskolin	15.00	11.25		
	FORSKOLIN - 4 bottles of 60 caps of elemental forskolin	50.00	37.50		
00993	FUCOXANTHIN-SLIM™ - 90 softgels	44.00	33.00		
	FUCOXANTHIN-SLIM™ - 4 bottles of 90 softgels	156.00	117.00		
G					
00127	GABA POWDER - 100 grams of powder	\$23.75	\$17.81		
	GABA POWDER - 4 bottles of powder	90.00	67.50		
00559	GAMMA E TOCOPHEROL/TOCOTRIENOLS - 60 softgels	42.00	31.50		
	GAMMA E TOCOPHEROL/TOCOTRIENOLS - 4 bottles of 60 softgels	148.00	111.00		
00759	GAMMA E TOCOPHEROL W/SESAME LIGNANS - 60 softgels	32.00	24.00		
	GAMMA E TOCOPHEROL W/SESAME LIGNANS - 4 bottles of 60 softgels	116.00	87.00		
00612	GELATIN CAPSULES - Set of 1000 empty caps size "0"	15.00	11.25		
	GELATIN CAPSULES - 4 Sets	56.00	42.00		
00117	GELATIN CAPSULES - Set of 1000 empty caps size "00"	19.00	14.25		
	GELATIN CAPSULES - 4 Sets	72.00	54.00		
01301	GH PITUITARY SUPPORT DAY FORMULA - 120 vegetarian tabs	48.00	36.00		
	GH PITUITARY SUPPORT DAY FORMULA - 4 bottles of 120 veg. tabs	176.00	132.00		
01302	GH PITUITARY SUPPORT NIGHT FORMULA - 120 veg. caps	25.00	18.75		
	GH PITUITARY SUPPORT NIGHT FORMULA - 4 bottles of 120 veg. caps	90.00	67.50		
00778	GINKGO BILOBA CERTIFIED EXTRACT™ - 120 mg, 365 caps	46.00	34.50		
	GINKGO BILOBA CERTIFIED EXTRACT™ - 2 bottles of 365 caps	87.00	65.25		
00504	(SUPER) GINKGO EXTRACT - 120 mg, 100 caps	29.00	21.75		
	(SUPER) GINKGO EXTRACT - 4 bottles of 100 caps	106.00	79.50		
01032	(PANAX RED) GINSENG - 60 veg. caps	29.99	22.49		
00756	(MEGA) GLA WITH SESAME LIGNANS - 60 softgels	19.50	14.63		
	(MEGA) GLA WITH SESAME LIGNANS - 4 bottles of 60 softgels	72.00	54.00		
00345	(L) GLUTAMINE CAPSULES - 500 mg, 100 caps	14.95	11.21		
	(L) GLUTAMINE CAPSULES - 4 bottles of 100 caps	54.00	40.50		
SUB-TOTAL OF COLUMN 8					

FEBRUARY 2010

Buyers Club Order Form

SUPER SALE SAVINGS ON ALL PRODUCTS
To order call: 1-954-766-8433 or 1-800-544-4440

No.		Retail	Member Price	Qty	Total
G CONTINUED					
00141	(L)-GLUTAMINE POWDER - 100 grams of powder	\$19.00	\$14.25		
	(L)-GLUTAMINE POWDER - 4 bottles of powder	67.52	50.64		
00522	GLUCOSAMINE/CHONDROITIN CAPSULES - 100 caps	38.00	28.50		
	GLUCOSAMINE/CHONDROITIN CAPSULES - 4 bottles of 100 caps	128.00	96.00		
00512	GLUCOFIT™ (COROSOLIC ACID) - 100 softgels	28.00	21.00		
	GLUCOFIT™ (COROSOLIC ACID) - 4 bottles of 100 softgels	100.00	75.00		
00113	GLUTATHIONE, C & CYSTEINE - 750 mg, 100 caps	18.00	13.50		
	GLUTATHIONE, C & CYSTEINE - 4 bottles of 100 caps	64.00	48.00		
00314	(MEGA) L-GLUTATHIONE - 250 mg, 60 caps	35.95	26.96		
	(MEGA) L-GLUTATHIONE - 4 bottles of 60 caps	129.44	97.08		
00795	GLYCINE CAPSULES - 1000 mg, 100 caps	12.00	9.00		
	GLYCINE CAPSULES - 4 bottles of 100 caps	43.20	32.40		
00128	GLYCINE POWDER - 300 grams of powder	18.00	13.50		
	GLYCINE POWDER - 4 bottles of powder	64.00	48.00		
01091	(WHOLE) GRAPE EXTRACT w/RESVERATROL - 60 veg. caps	36.00	27.00		
	(WHOLE) GRAPE EXTRACT w/RESVERATROL - 4 bottles of 60 veg. caps	136.00	102.00		
01411	GRAPE SEED EXTRACT w/RESVERATROL & PTEROSTILBENE-100 mg, 60 veg. caps	36.00	27.00		
	GRAPE SEED EXTRACT w/RESVERATROL & PTEROSTILBENE-4 bottles of 60 veg. caps	136.00	102.00		
00203	GREEN TEA LEAVES - 300 grams of powder	15.00	11.25		
	GREEN TEA LEAVES - 4 bottles of powder	50.00	37.50		
00953	(MEGA) GREEN TEA EXTRACT - lightly caffeinated - 100 veg. caps	28.00	21.00		
	(MEGA) GREEN TEA EXTRACT - 4 bottles of 100 veg. caps	106.00	79.50		
00954	(MEGA) GREEN TEA EXTRACT - decaffeinated - 100 veg. caps	28.00	21.00		
	(MEGA) GREEN TEA EXTRACT - 4 bottles of 100 veg. caps	106.00	79.50		
H					
00230	HCA - 90 caps	\$18.00	\$13.50		
	HCA - 4 bottles of 90 caps	62.00	46.50		
00656	HEPATOPRO - 900 mg, 60 softgel	49.50	37.13		
	HEPATOPRO - 4 bottles of 60 softgel	188.00	141.00		
00138	HESPERIDIN COMPLEX POWDER - 300 grams of powder	19.00	14.25		
	HESPERIDIN COMPLEX POWDER - 4 bottles of powder	65.00	48.75		
00873	HI-LIGNAN NUTRI-FLAX - 16 oz powder	9.95	7.46		
00800	HOMOCYSTEINE RESIST - 100 caps	24.00	18.00		
	HOMOCYSTEINE RESIST - 4 bottles of 100 caps	86.40	64.80		
00067	HOODIA GORDONII - 60 caps	14.00	10.50		
00627	HUPERZINE A W/NATURAL VITAMIN E - 50 mcg, 60 caps	27.95	20.96		
	HUPERZINE A W/NATURAL VITAMIN E - 4 bottles of 60 caps	96.00	72.00		
00661	HYDRODERM® - 1 oz bottle	79.95	59.96		
	HYDRODERM® - 2 - 1 oz bottles	130.66	98.00		
I					
00955	IMMUNE PROTECT W/PARACTIN® - 30 veg. caps	\$29.50	\$22.13		
	IMMUNE PROTECT W/PARACTIN® - 4 bottles of 30 veg. caps	106.20	79.65		
01004	INNERPOWER™ W/SWEET STEVIA- Tropical Flavor Powder	44.00	33.00		
00155	INOSINE CAPSULES - 500 mg, 60 caps	22.95	17.21		
	INOSINE CAPSULES - 4 bottles of 60 caps	84.00	63.00		
00140	INOSINE POWDER - 100 grams of powder	52.00	39.00		
	INOSINE POWDER - 4 bottles of powder	180.00	135.00		
00109	INOSITOL CAPSULES - 500 mg, 100 caps	13.50	10.13		
	INOSITOL CAPSULES - 4 bottles of 100 caps	50.00	37.50		
00410	INOSITOL CAPSULES - 1000 mg, 360 caps	48.00	36.00		
	INOSITOL CAPSULES - 4 bottles of 360 caps	175.00	131.25		
00108	INOSITOL POWDER - 100 grams of powder	18.00	13.50		
	INOSITOL POWDER - 4 bottles of powder	67.52	50.64		
01021	INTACT™ DIGEST - 8 fl oz	29.95	22.46		
	INTACT™ DIGEST - 4 - 8 fl oz bottles	112.00	84.00		
SUB-TOTAL OF COLUMN 9					

FEBRUARY 2010

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

No.		Retail	Member Price	Qty	Total
J, K					
01292	INTEGRA-LEAN® IRVINGIA - 150 mg, 60 veg. caps	\$56.00	\$42.00		
	INTEGRA-LEAN® IRVINGIA - 4 bottles of 60 veg. caps	192.00	144.00		
01002	IODORAL - 180 caps	50.00	37.50		
00563	IRON PROTEIN PLUS - 300 mg, 100 caps	24.00	18.00		
	IRON PROTEIN PLUS - 4 bottles of 100 caps	88.00	66.00		
01492	(OPTIMIZED) IRVINGIA W/PHASE 3™ - 120 veg. caps	78.00	58.50		
	(OPTIMIZED) IRVINGIA w/PHASE 3™ - 4 bottles of 120 veg. caps	288.00	216.00		
L					
00513	LACTOFERRIN (APOLACTOFERRIN) CAPS - 300 mg, 60 caps	\$40.00	\$30.00		
	LACTOFERRIN (APOLACTOFERRIN) CAPS - 4 bottles of 60 caps	144.00	108.00		
00017	LAVILIN UNDERARM DEODORANT - 12.5 grams of cream	15.00	11.25		
	LAVILIN UNDERARM DEODORANT - 4 jars of cream	55.00	41.25		
01003	LIFE EXTENSION CAT MIX - 100 grams of powder	15.00	11.25		
	LIFE EXTENSION CAT MIX - 4 bottles of powder	48.00	36.00		
00544	LIFE EXTENSION DOG MIX - 100 grams of powder	19.50	14.63		
	LIFE EXTENSION DOG MIX - 4 bottles of powder	64.00	48.00		
00010	LIFE EXTENSION HAIR CONDITIONER - 16 oz bottle	6.00	4.50		
	LIFE EXTENSION HAIR CONDITIONER - 4 bottles	18.00	13.50		
00020	LIFE EXTENSION LECITHIN GRANULES - 16 oz. jar	15.00	11.25		
	LIFE EXTENSION LECITHIN GRANULES - 4 jars	50.00	37.50		
00019	LIFE EXTENSION LECITHIN w/B5 AND BHA - 16 oz. jar	15.00	11.25		
	LIFE EXTENSION LECITHIN w/B5 AND BHA - 4 16 oz. 4 jars	50.00	37.50		
01455	LIFE EXTENSION MIX™ - 315 tablets	98.00	73.50		
	LIFE EXTENSION MIX™ - 4 bottles of 315 tablets	344.00	258.00		
	LIFE EXTENSION MIX™ - 10 bottles of 315 tablets	695.00	521.25		
01457	LIFE EXTENSION MIX™ w/EXTRA NIACIN - 315 tablets	98.00	73.50		
	LIFE EXTENSION MIX™ w/EXTRA NIACIN - 4 bottles of 315 tablets	344.00	258.00		
	LIFE EXTENSION MIX™ w/EXTRA NIACIN - 10 bottles of 315 tablets	695.00	521.25		
01454	LIFE EXTENSION MIX™ - 490 caps	110.00	82.50		
	LIFE EXTENSION MIX™ - 4 bottles of 490 caps	392.00	294.00		
	LIFE EXTENSION MIX™ - 10 bottles of 490 caps	850.00	637.50		
01456	LIFE EXTENSION MIX™ POWDER - 14.81 oz of powder	98.00	73.50		
	LIFE EXTENSION MIX™ POWDER - 4 bottles of powder	344.00	258.00		
	LIFE EXTENSION MIX™ POWDER - 10 bottles of powder	720.00	540.00		
01465	LIFE EXTENSION MIX™ - 315 tablets without copper	98.00	73.50		
	LIFE EXTENSION MIX™ - 4 bottles of 315 tablets without copper	344.00	258.00		
	LIFE EXTENSION MIX™ - 10 bottles of 315 tablets without copper	695.00	521.25		
01467	LIFE EXTENSION MIX™ w/EXTRA NIACIN 315 tablets w/o copper	98.00	73.50		
	LIFE EXTENSION MIX™ w/EXTRA NIACIN 4 bottles of 315 tablets w/o copper	344.00	258.00		
	LIFE EXTENSION MIX™ W/EXTRA NIACIN 10 bottles of 315 tablets w/o copper	695.00	521.25		
SUB-TOTAL OF COLUMN 10					

OFFER ENDS FEBRUARY 1, 2010

To order online visit www.lifeextension.com/SuperSale

No.		Retail	Member Price	Qty	Total
01464	LIFE EXTENSION MIX™ - 490 caps without copper	\$110.00	\$82.50		
	LIFE EXTENSION MIX™ - 4 bottles of 490 caps without copper	392.00	294.00		
	LIFE EXTENSION MIX™ - 10 bottles of 490 caps without copper	850.00	637.50		
01466	LIFE EXTENSION MIX™ POWDER - 14.81 oz of powder without copper	98.00	73.50		
	LIFE EXTENSION MIX™ POWDER - 4 bottles of powder without copper	344.00	258.00		
	LIFE EXTENSION MIX™ POWDER - 10 bottles of powder without copper	720.00	540.00		
01279	LIFE EXTENSION MOUTHWASH w/POMEGRANATE - 16 oz bottle	18.50	13.88		
	LIFE EXTENSION MOUTHWASH w/POMEGRANATE - 4 -16 oz bottle	68.00	51.00		
01401	LIFE EXTENSION ONE-PER-DAY - 60 veg. tablets	19.95	14.96		
	LIFE EXTENSION ONE-PER-DAY - 4 bottles of 60 veg. tablets	72.00	54.00		
00011	LIFE EXTENSION SHAMPOO - 16 oz bottle	6.00	4.50		
	LIFE EXTENSION SHAMPOO - 4 bottles	19.00	14.25		
01278	LIFE EXTENSION TOOTHPASTE MINT FLAVOR - 4 oz	9.50	7.13		
	LIFE EXTENSION TOOTHPASTE MINT FLAVOR - 4, 4 oz tubes	34.68	26.00		
01416	LIFE EXTENSION TWO-PER-DAY - 60 veg. tablets	9.95	7.46		
	LIFE EXTENSION TWO-PER-DAY - 4 bottles of 60 veg. tablets	36.00	27.00		
01415	LIFE EXTENSION TWO-PER-DAY - 120 veg. tablets	18.95	14.21		
	LIFE EXTENSION TWO-PER-DAY - 4 bottles of 120 veg. tablets	68.00	51.00		
00263	LIFE FLORA™ - 300 mg, 120 caps	20.50	15.38		
	LIFE FLORA™ - 4 bottles of 120 caps	75.00	56.25		
00032	LIFE MIX - 1 lb can of powder	25.00	18.75		
	LIFE MIX - 4 cans of powder	90.00	67.50		
00294	LIQUID EMULSIFIED VITAMIN A DROPS - 20,000 IU per drop	27.00	20.25		
	LIQUID EMULSIFIED VITAMIN A DROPS - 4 bottles	96.00	72.00		
00939	5-LOXIN® - 75 mg, 100 veg. caps	19.50	14.63		
	5-LOXIN® - 4 bottles of 100 veg. caps	69.00	51.75		
01013	L-LYSINE CAPSULES - 620 mg, 100 caps	9.00	6.75		
	L-LYSINE CAPSULES - 4 bottles of 100 caps	32.00	24.00		
00129	L-LYSINE POWDER - 300 grams of powder	16.00	12.00		
	L-LYSINE POWDER - 4 bottles of powder	55.00	41.25		
01017	LUSTRE™ - 90 tablets	17.00	12.75		
	LUSTRE™ - 4 bottles of 90 tablets	61.00	45.75		
01014	LUTEIN PLUS - 150 grams of powder	25.00	18.75		
	LUTEIN PLUS - 4 bottles of powder	88.00	66.00		
01028	LYCOPENE - 10 mg, 60 softgels	24.98	18.74		
00455	(MEGA) LYCOPENE EXTRACT - 15 mg, 90 softgels	35.00	26.25		
	(MEGA) LYCOPENE EXTRACT - 4 bottles of 90 softgels	120.00	90.00		
M					
01044	MAGNESIUM CAPS - 500 mg, 100 caps	\$9.50	\$7.13		
	MAGNESIUM CAPS - 4 bottles of 100 caps	32.00	24.00		
00502	MAGNESIUM CITRATE CAPS - 160 mg, 100 caps	9.00	6.75		
	MAGNESIUM CITRATE CAPS - 4 bottles of 100 caps	30.00	22.50		
00057	MAGNESIUM OXIDE POWDER - 1 kilogram of powder	14.00	10.50		
	MAGNESIUM OXIDE POWDER - 4 bottles of powder	52.52	39.39		
00704	MAITAKE™ SX-FRACTION - 90 tablets	49.95	37.46		
00048	(SUPER) MAX EPA - 12 oz liquid concentrate	18.50	13.88		
	(SUPER) MAX EPA - 4 bottles of concentrate	66.00	49.50		
00547	MELATONIN - 300 mcg, 100 caps	5.75	4.31		
	MELATONIN - 4 bottles of 100 caps	20.00	15.00		
00328	MELATONIN CAPSULES - 500 mcg, 200 caps	16.00	12.00		
	MELATONIN CAPSULES - 4 bottles of 200 caps	56.00	42.00		
00329	MELATONIN CAPSULES - 1 mg, 60 caps	5.00	3.75		
	MELATONIN CAPSULES - 4 bottles of 60 caps	18.52	13.89		

SUB-TOTAL OF COLUMN 11

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

Buyers Club Order Form

No.		Retail	Member Price	Qty	Total
00330	MELATONIN CAPSULES - 3 mg, 60 caps	\$8.00	\$6.00		
	MELATONIN CAPSULES - 4 bottles of 60 caps	27.52	20.64		
00333	MELATONIN CAPSULES - 3 mg, 60 time-release caps	10.50	7.88		
	MELATONIN CAPSULES - 4 bottles of 60 time-release caps	37.52	28.14		
00331	MELATONIN CAPSULES - 10 mg, 60 caps	28.00	21.00		
	MELATONIN CAPSULES - 4 bottles of 60 caps	96.00	72.00		
00332	MELATONIN LOZENGES - 3 mg, 60 dissolve-in-mouth lozenges	8.00	6.00		
	MELATONIN LOZENGES - 4 bottles of 60 dissolve-in-mouth lozenges	27.52	20.64		
00546	MELATONIN TIME RELEASE - 300 mcg, 100 caps	6.25	4.69		
	MELATONIN TIME RELEASE - 4 bottles of 100 caps	22.52	16.89		
00540	MELATONIN TIME RELEASE - 750 mcg, 60 caps	5.75	4.31		
	MELATONIN TIME RELEASE - 4 bottles of 60 caps	20.00	15.00		
01009	MEMORY UPGRADE - 600 grams powder	26.95	20.21		
01047	MENOPAUSE SOLUTIONS - 120 tablets	49.95	37.46		
00536	METHYLCOBALAMIN - 1 mg, 60 dissolve-in-mouth lozenges (vanilla)	9.95	7.46		
	METHYLCOBALAMIN - 4 bottles of 60 dissolve-in-mouth lozenges	32.00	24.00		
00537	METHYLCOBALAMIN - 5 mg, 60 dissolve-in-mouth lozenges (vanilla)	32.00	24.00		
	METHYLCOBALAMIN - 4 bottles of 60 dissolve-in-mouth lozenges	100.00	75.00		
	METHYLCOBALAMIN - 10 bottles of 60 dissolve-in-mouth lozenges	230.00	172.50		
00130	L-METHIONINE POWDER - 100 grams of powder	22.00	16.50		
	L-METHIONINE POWDER - 4 bottles of powder	75.00	56.25		
00709	MIGRA-EEZE (BUTTERBUR) - 60 softgels	29.50	22.13		
	MIGRA-EEZE (BUTTERBUR) - 4 bottles of 60 softgels	105.32	79.00		
00623	MINERAL FORMULA FOR MEN - 100 caps	15.00	11.25		
	MINERAL FORMULA FOR MEN - 4 bottles of 100 caps	50.00	37.50		
	MINERAL FORMULA FOR MEN - 12 bottles of 100 caps	130.00	97.50		
00624	MINERAL FORMULA FOR WOMEN - 100 caps	15.00	11.25		
	MINERAL FORMULA FOR WOMEN - 4 bottles of 100 caps	50.00	37.50		
	MINERAL FORMULA FOR WOMEN - 12 bottles of 100 caps	130.00	97.50		
01315	(SUPER) MIRAFORTE w/STANDARDIZED LIGNANS - 120 caps	62.00	46.50		
	(SUPER) MIRAFORTE w/STANDARDIZED LIGNANS - 4 bottles of 120 caps	224.00	168.00		
01368	MITOCHONDRIAL ENERGY OPTIMIZER - 120 caps	86.00	64.50		
	MITOCHONDRIAL ENERGY OPTIMIZER - 4 bottles of 120 caps	312.00	234.00		
00803	MISTORAL III w/COQ10 - 2 fl oz bottle	28.00	21.00		
	MISTORAL III w/COQ10 - 4 bottles	101.32	75.99		
00065	MK-7 - 90 mcg, 60 softgels	28.00	21.00		
	MK-7 - 4 bottles of 60 softgels	100.00	75.00		
00451	MSM (METHYL-SULFONYL-METHANE) - 1000 mg, 100 caps	14.00	10.50		
	MSM (METHYL-SULFONYL-METHANE) - 4 bottles of 100 caps	47.80	35.85		
N					
00215	N-ACETYL CYSTEINE CAPSULES - 600 mg, 60 caps	\$14.00	\$10.50		
	N-ACETYL CYSTEINE CAPSULES - 4 bottles of 60 caps	54.00	40.50		
00168	NAPCA W/ALOE VERA - 8 fl oz bottle	9.50	7.13		
	NAPCA W/ALOE VERA - 4 bottles	35.00	26.25		
00066	NATTOKINASE - 60 softgels	34.98	26.24		
00718	NATTOKINASE (FIBRINOPEN RESIST FORMULA) - 30 caps	49.00	36.75		
	NATTOKINASE (FIBRINOPEN RESIST FORMULA) - 4 bottles of 30 caps	176.00	132.00		
00891	NATURAL APPETITE CONTROL - 90 softgels	28.00	21.00		
	NATURAL APPETITE CONTROL - 4 bottles of 90 softgels	100.80	75.60		
00984	NATURAL BP MANAGEMENT - 60 tablets	42.00	31.50		
	NATURAL BP MANAGEMENT - 4 bottles of 60 tablets	151.20	113.40		
00913	NATURAL ESOPHAGUARD - 10 softgels	19.95	14.96		
	NATURAL ESOPHAGUARD - 2 boxes of 10 softgels	38.00	28.50		

SUB-TOTAL OF COLUMN 12

FEBRUARY 2010

Buyers Club Order Form

SUPER SALE SAVINGS ON ALL PRODUCTS
To order call: 1-954-766-8433 or 1-800-544-4440

No.		Retail	Member Price	Qty	Total
N CONTINUED					
00712	NATURAL ESTROGEN w/POMEGRANATE EXTRACT - 60 caplets	\$38.00	\$28.50		
	NATURAL ESTROGEN w/POMEGRANATE EXTRACT - 4 bottles of 60 caplets	132.00	99.00		
01221	NATURAL FEMALE SUPPORT - 30 veg. caps	28.00	21.00		
	NATURAL FEMALE SUPPORT - 4 bottles of 30 veg. caps	96.00	72.00		
00698	NATURAL RELIEF 1222™ - 2 oz tube	28.00	21.00		
	NATURAL RELIEF 1222™ - 4, 2 oz tubes	100.00	75.00		
00990	NATURAL SLEEP - 3 mg, 60 caps	12.99	9.74		
	NATURAL SLEEP - 4 bottles of 60 caps	40.00	30.00		
00277	NATURAL SLEEP MELATONIN - 5 mg, 60 caps	18.00	13.50		
	NATURAL SLEEP MELATONIN - 4 bottles of 60 caps	64.00	48.00		
00987	NATURAL STRESS RELIEF - 30 veg. caps	28.00	21.00		
	NATURAL STRESS RELIEF - 4 bottles of 30 veg. caps	96.00	72.00		
01000	NEW FACE SOLUTION - 1.7 oz bottle	54.00	40.50		
	NEW FACE SOLUTION - 4 bottles of 1.7 oz	196.00	147.00		
00096	NIACIN (B-3) CAPSULES - 1000 mg, 100 caps	12.75	9.56		
	NIACIN (B-3) CAPSULES - 4 bottles of 100 caps	48.00	36.00		
00372	NIACIN (B-3) CAPSULES - 500 mg, 100 caps	7.65	5.74		
	NIACIN (B-3) CAPSULES - 4 bottles of 100 caps	26.60	19.95		
00373	NO-FLUSH NIACIN - 800 mg, 100 caps	19.00	14.25		
	NO-FLUSH NIACIN - 4 bottles of 100 caps	68.00	51.00		
01050	NKO KRILL OIL - 60 softgels	33.95	25.46		
00262	NUTRAFLORA - 500 grams of powder	46.00	34.50		
	NUTRAFLORA - 4 bottles of powder	160.00	120.00		
01035	NUTRIM POWDER - 170 grams powder	25.95	19.46		
00523	N-ZIMES® - 270 caps	23.90	17.93		
	N-ZIMES® - 4 bottles of 270 caps	88.00	66.00		
O					
00104	OCTACOSANOL - 8000 mcg, 60 caps	\$15.75	\$11.81		
	OCTACOSANOL - 4 bottles of 60 caps	56.00	42.00		
01483	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT - 60 softgels	18.00	13.50		
	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT - 4 bottles of 60 softgels	64.00	48.00		
01482	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT - 120 softgels	32.00	24.00		
	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT - 4 bottles of 120 softgels	112.00	84.00		
	(SUPER) OMEGA 3 EPA/DHA W/SESAME LIGNANS & OLIVE FRUIT EXTRACT - 10 bottles of 120 softgels	249.00	186.75		
01484	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT (ENTERIC COATED) - 120 softgels	34.00	25.50		
	(SUPER) OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE FRUIT EXTRACT (ENTERIC COATED) - 4 bottles of 120 softgels	124.00	93.00		
	(SUPER) OMEGA 3 EPA/DHA W/SESAME LIGNANS & OLIVE FRUIT EXTRACT (ENTERIC COATED) - 10 bottles of 120 softgels	280.00	210.00		
00717	ONLY TRACE MINERALS - 90 caps	15.00	11.25		
	ONLY TRACE MINERALS - 4 bottles of 90 caps	50.00	37.50		
00915	OPTIZINC® - 30 mg, 90 veg. caps	5.95	4.46		
	OPTIZINC® - 4 bottles of 90 veg. caps	20.00	15.00		
00043	L-ORNITHINE CAPSULES - 500 mg, 100 caps	24.95	18.71		
	L-ORNITHINE CAPSULES - 4 bottles of 100 caps	80.00	60.00		
P					
00107	PABA CAPS - 500 mg, 100 caps	\$10.50	\$7.88		
	PABA CAPS - 4 bottles of 100 caps	37.50	28.12		
00106	PABA POWDER - 100 grams of powder	15.00	11.25		
	PABA POWDER - 4 bottles of powder	55.00	41.25		

SUB-TOTAL OF COLUMN 13

FEBRUARY 2010

* Product cannot be sold outside the USA.

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

No.		Retail	Member Price	Qty	Total
00073	PANCREATIN - 500 mg, 50 caps		\$9.75	\$7.31	
	PANCREATIN - 4 bottles of 50 caps		36.00	27.00	
00137	PAPAIN POWDER - 100 grams of powder		24.00	18.00	
	PAPAIN POWDER - 4 bottles of powder		80.00	60.00	
00923	PEAK ATP® WITH GLYCOCARN® - 60 tablets		59.00	44.25	
	PEAK ATP® WITH GLYCOCARN® - 4 bottles of 60 tablets		224.00	168.00	
00342	PECTA-SOL - One-month supply modified citrus pectin		99.95	74.96	
	PECTA-SOL - Four-month supply modified citrus pectin		382.80	287.10	
00872	PGX FIBER POWDER (SLIMSTYLES) - 6.4 oz powder		22.95	17.21	
00673	(WELLBETX) PGX SOLUBLE FIBER BLEND -180 caps		34.95	26.21	
00865	PHARMAGABA™ - 60 chewable tablets		29.95	22.46	
	PHARMAGABA™ - 4 bottles of 60 chewable tablets		108.00	81.00	
00131	L-PHENYLALANINE POWDER - 100 grams of powder		28.00	21.00	
	L-PHENYLALANINE POWDER - 4 bottles of powder		100.00	75.00	
00368	PHOSPHATIDYLSERINE CAPS - 100 mg, 100 caps		54.00	40.50	
	PHOSPHATIDYLSERINE CAPS - 4 bottles of 100 caps		192.00	144.00	
01016	PHYTO-FOOD - 80 grams of powder		46.00	34.50	
	PHYTO-FOOD - 4 bottles of powder		172.00	129.00	
00614	PILL CUTTER/GRINDER		11.95	8.96	
00561	POLICOSANOL - 10 mg, 60 tablets		24.00	18.00	
	POLICOSANOL - 6 bottles of 60 tablets		108.00	81.00	
00994	(SUPER) POLYPHENOL EXTRACT w/ COCOAGOLD™ - 30 veg caps		24.00	18.00	
	(SUPER) POLYPHENOL EXTRACT w/ COCOAGOLD™ - 4 bottles of 30 veg caps		84.00	63.00	
00956	POMEGRANATE EXTRACT - 30 veg. caps		19.50	14.63	
	POMEGRANATE EXTRACT - 4 bottles of 30 veg. caps		70.20	52.64	
01256	POMEGRANATE EXTRACT w/ COCOAGOLD™ - 30 veg caps		24.00	18.00	
	POMEGRANATE EXTRACT w/ COCOAGOLD™ - 4 bottles of 30 veg caps		84.00	63.00	
00957	POMEGRANATE JUICE CONCENTRATE - 16 oz. liquid		25.95	19.46	
	POMEGRANATE JUICE CONCENTRATE - 4 bottles of 16 oz liquid		96.00	72.00	
00577	POTASSIUM IODIDE - 1 box, 14 tablets		6.95	5.21	
	POTASSIUM IODIDE - 4 boxes, 14 tablets		21.00	15.75	
00302	PREGNENOLONE CAPSULES - 50 mg, 100 caps		24.00	18.00	
	PREGNENOLONE CAPSULES - 4 bottles of 100 caps		76.00	57.00	
00700	PREGNENOLONE CAPSULES - 100 mg, 100 caps		26.00	19.50	
	PREGNENOLONE CAPSULES - 4 bottles of 100 caps		90.00	67.50	
*01373	PRELOX® NATURAL SEX FOR MEN® - 60 tablets		50.00	37.50	
	PRELOX® NATURAL SEX FOR MEN® - 4 bottles of 60 tablets		180.00	135.00	
00571	PRIMAL DEFENSE® - 900 mg, 90 caplets		49.95	37.46	
	PRIMAL DEFENSE® - 4 bottles of 90 caplets		194.00	145.50	
01019	PRIMAL DEFENSE® - 180 caplets		78.95	59.21	
00525	PROBOOST THYMIC PROTEIN A™ - 4 mcg, 30 packets		72.00	54.00	
	PROBOOST THYMIC PROTEIN A™ - 4 boxes		280.00	210.00	
00869	PRO FEM CREAM - 2 oz jar, Progesterone Cream		26.00	19.50	
	PRO FEM CREAM - 4, 2 oz jars, Progesterone Cream		96.00	72.00	
01020	PROGREENS® - 15 stick pack		22.50	16.88	
00396	PRO M - 500 mg, 100 caps		28.00	21.00	
	PRO M - 4 bottles of 100 caps		104.00	78.00	
*01475	(ULTRA NAT) PROSTATE FORMULA w/ 5-LOXIN® STAND. LIGNANS-60 softgels		38.00	28.50	
	(ULTRA NAT) PROSTATE FORMULA w/ 5-LOXIN® STAND. LIGNANS-4 bot of 60 softgels		140.00	105.00	
	(ULTRA NAT) PROSTATE FORMULA w/ 5-LOXIN® STAND. LIGNANS-12 bot of 60 softgels		384.00	288.00	
01092	(ENHANCED LIFE EXTENSION) PROTEIN (VANILLA) - 1 kg. of powder		38.00	28.50	
	(ENHANCED LIFE EXTENSION) PROTEIN (VANILLA) - 4 bot of powder		144.00	108.00	
01093	(ENHANCED LIFE EXTENSION) PROTEIN (CHOCOLATE) - 1 kg of powder		38.00	28.50	
	(ENHANCED LIFE EXTENSION) PROTEIN (CHOCOLATE) - 4 bot of powder		144.00	108.00	

SUB-TOTAL OF COLUMN 14

OFFER ENDS FEBRUARY 1, 2010

To order online visit www.lifeextension.com/SuperSale

No.		Retail	Member Price	Qty	Total
01094	(ENHANCED LIFE EXTENSION) PROTEIN (NATURAL) - 1 kg of powder	\$38.00	\$28.50		
	(ENHANCED LIFE EXTENSION) PROTEIN (NATURAL) - 4 bot of powder	144.00	108.00		
01095	(ENHANCED LIFE EXTENSION) PROTEIN (BERRY) - 1 kg of powder	38.00	28.50		
	(ENHANCED LIFE EXTENSION) PROTEIN (BERRY) - 4 bot of powder	144.00	108.00		
00290	(DESIGNER WHEY) PROTEIN - Vanilla 2 lbs of powder	41.95	31.46		
	(DESIGNER WHEY) PROTEIN - 4 bottles of Vanilla powder	152.00	114.00		
00282	(DESIGNER WHEY) PROTEIN - Chocolate 2 lbs of powder	41.95	31.46		
	(DESIGNER WHEY) PROTEIN - 4 bottles of Chocolate powder	152.00	114.00		
00283	(DESIGNER WHEY) PROTEIN - Natural 2 lbs of powder	41.95	31.46		
	(DESIGNER WHEY) PROTEIN - 4 bottles of Natural powder	152.00	114.00		
01056	PURE-GAR® - 200 caps	24.95	18.71		
	PURE-GAR® - 4 bottles of 200 caps	84.00	63.00		
01209	(WATER-SOLUBLE) PUMPKIN SEED EXTRACT - 60 veg. caps	20.00	15.00		
	(WATER-SOLUBLE) PUMPKIN SEED EXTRACT - 4 bottles 60 veg. caps	72.00	54.00		
01210	(WATER-SOLUBLE) PUMPKIN SEED EXT w/SOY ISOFLAVONES - 60 veg. caps	22.00	16.50		
	(WATER-SOLUBLE) PUMPKIN SEED EXT w/SOY ISOFLAVONES - 4 bottles 60 veg.	80.00	60.00		
01031	PYCNOGENOL® - 60 veg. caps	86.95	65.21		
01207	PYRIDOXAMINE CAPS - 50 mg, 60 veg. caps	22.00	16.50		
	PYRIDOXAMINE CAPS - 4 bottles of 60 veg. caps	79.20	59.40		
Q, R					
01309	(OPTIMIZED) QUERCETIN - 250 mg, 60 veg. caps	\$18.00	\$13.50		
	(OPTIMIZED) QUERCETIN - 4 bottles of 60 veg. caps	68.00	51.00		
01030	RED YEAST RICE (Bluebonnet) - 600 mg, 60 veg. caps	15.95	11.96		
00979	RED YEAST RICE (Nature's Plus) - 600 mg, 60 veg. caps	22.75	17.06		
00060	RED YEAST RICE EXTENDED RELEASE (Nature's Plus) - 30 veg. tablets	20.55	15.41		
00605	REGIMINT® - 60 enteric-coated caps	19.95	14.96		
	REGIMINT® - 4 bottles of 60 caps	74.68	56.01		
00448	REJUVENEX® BODY LOTION - 6 oz tube	24.00	18.00		
	REJUVENEX® BODY LOTION - 4 tubes	79.20	59.40		
	REJUVENEX® BODY LOTION - 8 tubes	136.00	102.00		
00918	REJUVENEX® FACTOR - 1.7 oz Airless pump bottle	65.00	48.75		
	REJUVENEX® FACTOR - 2 bottles	120.00	90.00		
	REJUVENEX® FACTOR - 4 bottles	220.00	165.00		
	REJUVENEX® FACTOR - 8 bottles	399.28	299.46		
01220	(ULTRA) REJUVENEX® - 2 oz jar	52.00	39.00		
	(ULTRA) REJUVENEX® - 2 jars	96.00	72.00		
	(ULTRA) REJUVENEX® - 4 jars	176.00	132.00		
	(ULTRA) REJUVENEX® - 8 jars	319.44	239.38		
00676	(ULTRA) REJUVENIGHT (DREAM CREAM) - 2 oz jar	39.95	29.96		
	(ULTRA) REJUVENIGHT (DREAM CREAM) - 4 jars	144.00	108.00		
00706	(ULTRA) REJUVENIGHT w/PROGESTERONE - 2 oz jar	42.00	31.50		
	(ULTRA) REJUVENIGHT w/PROGESTERONE - 4 jars	152.00	114.00		
01413	RESVERATROL w/PTEROSTILBENE - 20 mg, 60 veg. caps	24.00	18.00		
	RESVERATROL w/PTEROSTILBENE - 4 bottles of 60 veg. caps	88.00	66.00		
01410	RESVERATROL w/PTEROSTILBENE - 100 mg, 60 veg. caps	36.00	27.00		
	RESVERATROL w/PTEROSTILBENE - 4 bottles of 60 veg. caps	128.00	96.00		
01409	(OPTIMIZED) RESVERATROL w/PTEROSTILBENE - 250 mg, 60 veg. caps	46.00	34.50		
	(OPTIMIZED) RESVERATROL w/PTEROSTILBENE - 4 bottles of 60 veg. caps	165.32	124.00		
00889	RHODIOLA EXTRACT - 250 mg, 60 veg. caps	11.75	8.81		
	RHODIOLA EXTRACT - 4 bottles of 60 veg. caps	42.32	31.74		
00972	(D) RIBOSE POWDER - 150 grams of powder	27.50	20.63		
	(D) RIBOSE POWDER - 4 jars of 150 grams	99.00	74.25		
00973	(D) RIBOSE CAPS - 120 veg. caps	32.00	24.00		
	(D) RIBOSE CAPS - 4 bottles of 120 veg. caps	115.20	86.40		

SUB-TOTAL OF COLUMN 15

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10% ON ALL PRODUCTS DURING SUPER SALE**

Buyers Club Order Form

No.		Retail	Member Price	Qty	Total
01208	(SUPER) R-LIPOIC ACID - 300 mg, 60 veg. caps	\$49.00	\$36.75		
	(SUPER) R-LIPOIC ACID - 4 bottles of 60 veg. caps	180.00	135.00		
00070	RNA CAPSULES - 500 mg, 100 caps	17.95	13.46		
	RNA CAPSULES - 4 bottles of 100 caps	64.64	48.48		
00143	RNA POWDER - 100 grams of powder	24.00	18.00		
	RNA POWDER - 4 bottles of powder	80.00	60.00		
00917	ROSMARINIC ACID EXTRACT - 60 veg. caps	38.00	28.50		
	ROSMARINIC ACID EXTRACT - 4 bottles of 60 veg. caps	144.00	108.00		
00139	RUTIN POWDER - 100 grams of powder	15.50	11.63		
	RUTIN POWDER - 4 bottles of powder	55.00	41.25		
S					
* 00573	SAMBU GUARD - 175 ml		\$17.49	\$13.99	
00358	SAME (S-ADENOSYL-METHIONINE) - 200 mg, 20 enteric coated tablets	18.95	14.21		
	SAME (S-ADENOSYL-METHIONINE) - 8 boxes of 20 enteric coated tablets	132.80	99.60		
00453	SAME (S-ADENOSYL-METHIONINE) - 200 mg, 50 enteric coated tablets	45.00	33.75		
	SAME (S-ADENOSYL-METHIONINE) - 4 bottles of 50 enteric coated tablets	156.00	117.00		
00557	SAME (S-ADENOSYL-METHIONINE) - 400 mg, 20 enteric coated tablets	35.00	26.25		
	SAME (S-ADENOSYL-METHIONINE) - 6 boxes of 20 enteric coated tablets	168.00	126.00		
01043	SEA-IODINE™ - 1000 mcg, 60 veg. caps	8.00	6.00		
	SEA-IODINE™ - 4 bottles of 60 veg. caps	28.80	21.60		
00046	SELENIUM - 2 oz dropper bottle	9.45	7.09		
	SELENIUM - 4 bottles of 2 oz bottle	36.00	27.00		
00567	SE-METHYLELENOCYSTEINE - 200 mcg, 100 caps	12.00	9.00		
	SE-METHYLELENOCYSTEINE - 4 bottles of 100 caps	44.00	33.00		
00318	SERRAFLAZYME - 100 tablets	18.00	13.50		
	SERRAFLAZYME - 4 bottles of 100 tablets	64.00	48.00		
00339	SHARK-A CARTILAGE - 750 mg, 270 caps	68.00	51.00		
	SHARK-A CARTILAGE - 4 bottles of 270 caps	255.00	191.25		
00284	SHARK LIVER OIL (NORWEGIAN) - 1000 mg, 30 softgels	18.00	13.50		
	SHARK LIVER OIL (NORWEGIAN) - 4 bottles of 30 softgels	64.00	48.00		
00503	SILIBININ PLUS - 90 veg. caps	36.00	27.00		
	SILIBININ PLUS - 4 bottles of 90 veg. caps	128.00	96.00		
00184	SILYMARIN - 100 mg, 50 caps	9.25	6.94		
	SILYMARIN - 4 bottles of 50 caps	33.00	24.75		
00702	(MEGA) SILYMARIN WITH ISOSILYBIN B - 100 caps	38.00	28.50		
	(MEGA) SILYMARIN WITH ISOSILYBIN B - 4 bottles of 100 caps	136.80	102.60		
00872	SLIMSTYLES PGX® - 6.4 oz powder	22.95	17.21		
00961	SODZYME® W/GLISODIN® AND WOLFBERRY - 90 veg. caps	28.00	21.00		
	SODZYME® W/GLISODIN® AND WOLFBERRY - 4 bottles of 90 veg. caps	96.00	72.00		
00657	SOLARSHIELD SUNGLASSES - 1 pair smoke color	12.99	9.74		
	SOLARSHIELD SUNGLASSES - 2 pair smoke color	23.00	17.25		
00785	(ULTRA) SOY EXTRACT - 675 mg, 600 caps	349.00	261.75		
	(ULTRA) SOY EXTRACT - 4 bottles of 600 caps	1260.00	945.00		
00304	SOY POWER POWDER - 300 grams of powder	31.50	23.63		
	SOY POWER POWDER - 4 bottles of powder	114.00	85.50		
00286	SOY PROTEIN POWDER (concentrated soy protein) - 16 oz of powder	12.00	9.00		
	SOY PROTEIN POWDER - 4 bottles of powder	42.50	31.87		
00432	STEVIA EXTRACT - 100 packets, 1 gram each	9.95	7.46		
	STEVIA EXTRACT - 4 boxes of 100 packets	36.00	27.00		
00351	ST. JOHN'S WORT EXTRACT - 300 mg, 60 caps	10.98	8.24		
	ST. JOHN'S WORT EXTRACT - 4 bottles of 60 caps	40.00	30.00		
00327	ST. JOHN'S WORT EXTRACT (PLANETARY) - 600 mg, 60 tablets	24.95	18.71		
	ST. JOHN'S WORT EXTRACT (PLANETARY) - 4 bottles of 60 tablets	85.00	63.75		
00971	SUN PROTECTION SPRAY W/BETA GLUCAN - SPF30 - 6 oz bottle	14.95	11.21		
	SUN PROTECTION SPRAY W/BETA GLUCAN - SPF30 - 4 bottles, 6 oz	54.00	40.50		

SUB-TOTAL OF COLUMN 16

* These products are not 25% off retail price.

FEBRUARY 2010

Buyers Club Order Form

SUPER SALE SAVINGS ON ALL PRODUCTS
To order call: 1-954-766-8433 or 1-800-544-4440

No.		Retail	Member Price	Qty	Total
00747	(OVERCAST POLARIZED) SUNGLASSES - gray color, large	\$27.00	\$20.25		
	(OVERCAST POLARIZED) SUNGLASSES - 2 pairs, gray color	42.00	31.50		
00748	(OVERCAST POLARIZED) SUNGLASSES - gray color, medium	27.00	20.25		
	(OVERCAST POLARIZED) SUNGLASSES - 2 pairs, gray color	42.00	31.50		
00758	SUPER ABSORBABLE SOY ISOFLAVONES - 60 caps	28.00	21.00		
	SUPER ABSORBABLE SOY ISOFLAVONES - 4 bottles of 60 caps	100.00	75.00		
01408	SUPER SAW PALMETTO/NETTLE ROOT W/BETA-SITOSTEROL 60 softgels	28.00	21.00		
	SUPER SAW PALMETTO/NETTLE ROOT w/BETA-SITOSTEROL 4 bottles of 60 softgels	104.00	78.00		
	SUPER SAW PALMETTO/NETTLE ROOT w/BETA-SITOSTEROL One-year supply	288.00	216.00		
01407	SUPER SAW PALMETTO W/BETA-SITOSTEROL - 30 softgels	15.00	11.25		
	SUPER SAW PALMETTO W/BETA-SITOSTEROL - 12 bottles of 30 softgels	144.00	108.00		
00578	SUPER SELENIUM COMPLEX - 200 mcg, 100 caps	12.00	9.00		
	SUPER SELENIUM COMPLEX - 4 bottles of 100 caps	40.00	30.00		
	SUPER SELENIUM COMPLEX - 12 bottles of 100 caps	108.00	81.00		
00674	SYTRINOL™ - 60 softgels	32.00	24.00		
	SYTRINOL™ - 4 bottles of 60 softgels	112.00	84.00		
T					
	TAL • SHI™ COSMETICS • For information please call 1-800-544-4440 or visit www.lifeextension.com				
00199	TAURINE CAPSULES - 1000 mg, 50 caps	\$8.95	\$6.71		
	TAURINE CAPSULES - 4 bottles of 50 caps	32.00	24.00		
00133	TAURINE POWDER - 300 grams, powder	20.00	15.00		
	TAURINE POWDER - 4 bottles of powder	67.52	50.64		
01304	THEAFLAVIN STANDARDIZED EXTRACT - 30 veg. caps	18.00	13.50		
	THEAFLAVIN STANDARDIZED EXTRACT - 4 bottles of 30 veg. caps	64.00	48.00		
00555	(L) THEANINE - 100 mg, 60 caps	24.00	18.00		
	(L) THEANINE - 4 bottles of 60 caps	82.00	61.50		
01038	ATHERALAC - 30 caps	44.95	33.71		
00224	THYMIC IMMUNE FACTORS - 100 caps	18.00	13.50		
	THYMIC IMMUNE FACTORS - 4 bottles of 100 caps	56.00	42.00		
00668	(METABOLIC ADVANTAGE) THYROID FORMULA - 100 caps	17.95	13.46		
00349	TMG - 50 grams of powder	14.00	10.50		
	TMG - 4 bottles of powder	44.00	33.00		
00359	TMG TABLETS - 500 mg, 180 tablets	18.00	13.50		
	TMG TABLETS - 4 bottles of 180 tablets	62.00	46.50		
00366	TOCOTRIENOLS - 50 mg, 60 softgel caps	39.95	29.96		
	TOCOTRIENOLS - 4 bottles of 60 softgel caps	144.00	108.00		
00781	TOCOTRIENOLS WITH SESAME LIGNANS - 60 softgels	38.00	28.50		
	TOCOTRIENOLS WITH SESAME LIGNANS - 4 bottles of 60 softgels	144.00	108.00		
01400	(SUPER-ABSORBABLE) TOCOTRIENOLS - 60 softgels	30.00	22.50		
	(SUPER-ABSORBABLE) TOCOTRIENOLS - 4 bottles of 60 softgels	112.00	84.00		
01274	TOTAL SUN PROTECTION CREAM W/BETAGLUCANS - SPF 30, 4 oz Tube & PHOTO-AGING RECOVERY COMPLEX	28.00	21.00		
	TOTAL SUN PROTECTION CREAM W/BETAGLUCANS - 4-4 oz Tubes & PHOTO-AGING RECOVERY COMPLEX	100.80	75.60		
01468	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT- 60 veg. caps	24.00	18.00		
	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT-4 bot 60 veg. caps	88.00	66.00		
01469	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT w/RESVERATROL - 60 veg. caps	32.00	24.00		
	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT w/RESVERATROL - 4 bottles	118.40	88.80		
01046	TRIPLE ACTION UNDER EYE REJUVENATOR - 0.5 oz pump	24.00	18.00		
	TRIPLE ACTION UNDER EYE REJUVENATOR - 4 boxes	88.00	66.00		
00866	TRYPTOPURE® L-TRYPTOPHAN - 500 mg, 90 veg. caps	38.00	28.50		
	TRYPTOPURE® L-TRYPTOPHAN - 4 bottles of 90 veg. caps	136.00	102.00		

SUB-TOTAL OF COLUMN 17

FEBRUARY 2010

* These products are not 25% off retail price.

LIFE EXTENSION MEMBERS RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

DEDUCT AN **ADDITIONAL 10%** ON ALL PRODUCTS DURING SUPER SALE

No.		Retail	Member Price	Qty	Total
01202	(OPTIMIZED) TRYPTOPURE® PLUS - 90 veg. caps	\$40.00	\$30.00		
	(OPTIMIZED) TRYPTOPURE® PLUS - 4 bottles of 90 veg. caps	144.00	108.00		
00036	L-TYROSINE POWDER - 100 grams of powder	22.50	16.88		
	L-TYROSINE POWDER - 4 bottles of powder	80.00	60.00		
00326	L-TYROSINE TABLETS - 500 mg, 100 tablets	16.98	12.74		
	L-TYROSINE TABLETS - 4 bottles of 100 tablets	62.52	46.89		
00035	L-TYROSINE PLUS POWDER - 100 grams of powder	28.00	21.00		
	L-TYROSINE PLUS POWDER - 4 bottles of powder	98.00	73.50		
U					
* 00310	UDO'S CHOICE® - 17 oz liquid (omega fatty acids)	\$27.49	\$21.99		
* 00311	UDO'S CHOICE® - 1000 mg, 180 caps	33.79	27.03		
* 00322	UDO'S CHOICE® WHOLESOME FAST FOOD - 1 lb bottle	27.49	21.99		
V					
00213	VANADYL SULFATE - 7.5 mg, 100 tablets	\$15.00	\$11.25		
	VANADYL SULFATE - 4 bottles of 100 tablets	50.00	37.50		
01018	VASCULAR PROTECT - 120 vegetarian caps	48.00	36.00		
	VASCULAR PROTECT - 4 bottles of 120 vegetarian caps	180.00	135.00		
00252	VELVET DEER ANTLER - 250 mg, 30 caps	36.00	27.00		
	VELVET DEER ANTLER - 4 bottles of 30 caps	136.00	102.00		
00408	VENOTONE - 60 caps	18.95	14.21		
	VENOTONE - 4 bottles of 60 caps	64.00	48.00		
00427	VINPOCETINE - 5 mg, 100 tablets	18.00	13.50		
	VINPOCETINE - 4 bottles of 100 tablets	56.00	42.00		
01033	VISION OPTIMIZER - 90 caps	29.95	22.46		
01098	VITAL GREENS MIX - 319.5 grams of powder	48.00	36.00		
	VITAL GREENS MIX - 4 bottles of powder	176.00	132.00		
00091	VITAMIN B1 CAPS - 500 mg, 100 caps	15.50	11.63		
	VITAMIN B1 CAPS - 4 bottles of 100 caps	56.00	42.00		
00090	VITAMIN B1 POWDER - 100 grams of powder (Thiamin)	18.00	13.50		
	VITAMIN B1 POWDER - 4 bottles of powder	67.52	50.64		
00093	VITAMIN B2 CAPS - 100 mg, 100 caps	9.50	7.13		
	VITAMIN B2 CAPS - 4 bottles of 100 caps	35.00	26.25		
00092	VITAMIN B2 POWDER - 30 grams of powder	10.50	7.88		
	VITAMIN B2 POWDER - 4 bottles of powder	37.52	28.14		
00096	VITAMIN B3 (NIACIN) - 1000 mg, 100 caps	12.75	9.56		
	VITAMIN B3 (NIACIN) - 4 bottles of 100 caps	48.00	36.00		
00372	VITAMIN B3 (NIACIN) - 500 mg, 100 caps	7.65	5.74		
	VITAMIN B3 (NIACIN) - 4 bottles of 100 caps	26.60	19.95		
00094	VITAMIN B3 POWDER - 300 grams of powder	15.00	11.25		
	VITAMIN B3 POWDER - 4 bottles of powder	56.00	42.00		
00098	VITAMIN B5 CAPS - 500 mg, 100 caps (Pantothenic Acid)	10.50	7.88		
	VITAMIN B5 CAPS - 4 bottles of 100 caps	37.52	28.14		
00097	VITAMIN B5 POWDER - 100 grams of powder	15.25	11.44		
	VITAMIN B5 POWDER - 4 bottles of powder	55.00	41.25		
00556	VITAMIN B6 CAPS - 250 mg, 100 caps	12.50	9.38		
	VITAMIN B6 CAPS - 4 bottles of 100 caps	44.00	33.00		
00099	VITAMIN B6 POWDER - 100 grams of powder	24.50	18.38		
	VITAMIN B6 POWDER - 4 bottles of powder	88.00	66.00		
00361	VITAMIN B12 TABLETS - 500 mcg, 100 dissolve-in-mouth tablets	8.75	6.56		
	VITAMIN B12 TABLETS - 4 bottles of 100 dissolve-in-mouth tablets	29.00	21.75		
00144	VITAMIN B12 POWDER - 100 grams of powder	12.00	9.00		
	VITAMIN B12 POWDER - 4 bottles of powder	42.52	31.89		
00927	VITAMIN C WITH DIHYDROQUERCETIN - 1000 mg, 250 tablets	25.50	19.13		
	VITAMIN C WITH DIHYDROQUERCETIN - 4 bottles of 250 tablets	93.00	69.75		

SUB-TOTAL OF COLUMN 18

OFFER ENDS FEBRUARY 1, 2010

To order online visit www.lifeextension.com/SuperSale

No.		Retail	Member Price	Qty	Total
00084	(BUFFERED) VITAMIN C POWDER - 454.6 grams of powder	\$23.95	\$17.96		
	(BUFFERED) VITAMIN C POWDER - 4 bottles of powder	88.00	66.00		
00864	VITAMIN D - 2000 IU, 1 fl oz	28.00	21.00		
	VITAMIN D - 4 bottles of 1 fl oz	100.00	75.00		
00251	VITAMIN D3 CAPS - 1000 IU, 250 caps	12.50	9.38		
	VITAMIN D3 CAPS - 4 bottles of 250 caps	45.00	33.75		
00713	VITAMIN D3 CAPS - 5000 IU, 60 caps	11.00	8.25		
	VITAMIN D3 CAPS - 4 bottles of 60 caps	39.60	29.70		
01418	VITAMIN D3 CAPS - 7000 IU, 60 caps	14.00	10.50		
	VITAMIN D3 CAPS - 4 bottles of 60 caps	50.40	37.80		
01371	VITAMIN D3 w/SEA-IODINE - 1000 IU, 250 veg. caps	22.00	16.50		
	VITAMIN D3 w/SEA-IODINE - 4 bottles of 250 veg. caps	80.00	60.00		
01372	VITAMIN D3 w/SEA-IODINE - 5000 IU, 60 veg. caps	14.00	10.50		
	VITAMIN D3 w/SEA-IODINE - 4 bottles of 60 veg. caps	50.00	37.50		
00063	VITAMIN E CAPS (NATURAL) - 400 IU, 100 caps	18.75	14.06		
	VITAMIN E CAPS (NATURAL) - 4 bottles of 100 caps	69.00	51.75		
	VITAMIN E CAPS (NATURAL) - 10 bottles of 100 caps	150.00	112.50		
00062	VITAMIN E POWDER (SYNTHETIC) - 300 grams of powder	28.95	21.71		
	VITAMIN E POWDER (SYNTHETIC) - 4 bottles of powder	100.00	75.00		
01225	(LOW-DOSE) VITAMIN K2 (MK-7) - 90 softgels	18.00	13.50		
	(LOW-DOSE) VITAMIN K2 (MK-7) - 4 bottles of 90 softgels	64.00	48.00		
Z					
00986	(SUPER) ZEAXANTHIN W/LUTEIN & MESO-ZEAXANTHIN - 60 softgels	\$22.00	\$16.50		
	(SUPER) ZEAXANTHIN W/LUTEIN & MESO-ZEAXANTHIN - 4 bottles	79.20	59.40		
01286	(SUPER) ZEAXANTHIN W/LUTEIN & MESO-ZEAXANTHIN - 60 softgels PLUS ASTAXANTHIN	42.00	31.50		
	(SUPER) ZEAXANTHIN W/LUTEIN & MESO-ZEAXANTHIN PLUS ASTAXANTHIN - 4 bottles of 60 softgels	152.00	114.00		
00152	ZINC GLUCONATE POWDER - 100 grams of powder	8.00	6.00		
	ZINC GLUCONATE POWDER - 4 bottles of powder	27.50	20.62		
00061	ZINC LOZENGES WITH VITAMIN C - 75 lozenges	9.50	7.13		
	ZINC LOZENGES WITH VITAMIN C - 4 bottles of 75 lozenges	27.00	20.25		
01051	ZYFLAMEND - 120 softgels	60.95	45.71		
01029	ZYFLAMEND EASYCAPS - 180 caps	31.95	23.96		

SUB-TOTAL OF COLUMN 19

**GIVE THE GIFT of HEALTH, with a
LIFE EXTENSION GIFT CARD!**



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\$50—ITEM # GC050
\$100—ITEM # GC100

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As a member of the Life Extension Foundation®, you have the opportunity to participate in a great scientific endeavor. We are the world's premier organization dedicated to stopping and reversing aging.

Our 29-year track record shows that we have been five to ten years ahead of conventional and alternative medicine in making new life-saving therapies available to our members.

When you join the Life Extension Foundation®, we update you on the latest published medical research by sending you FREE books. Our most impressive publication is the 1,666-page *Disease Prevention and Treatment* protocol book that contains novel therapies to treat 133 common diseases of aging. *Disease Prevention and Treatment* is the only book that combines conventional and alternative therapies in order to implement a treatment regimen for fighting the multiple processes involved in degenerative disease.

Each month, Life Extension Foundation® members receive a magazine packed with the latest medical findings from around the world. Members also can call a toll-free phone number to talk to our knowledgeable health advisors about their health issues.

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33599	YOUNGER YOU • by Eric Braverman, MD	—	\$24.95	\$15.75	
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33666	ULTRAMETABOLISM: THE SIMPLE PLAN FOR AUTOMATIC WEIGHT LOSS • by Mark Hyman, MD	2006	\$25.00	\$15.35	
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33677	DANGEROUS DOSES • by Katherine Eban	2005	\$25.00	\$16.25	
33600	DISEASE PREVENTION AND TREATMENT, EXPANDED FOURTH EDITION (hardcover)	2003	\$49.95	\$37.46	
	Price per book when 4 books purchased	—	\$44.00	\$33.00	
33594	THE EDGE EFFECT • by Eric Braverman, MD (paperback)	2004	\$12.95	\$9.71	
33700	ENDING AGING • by Aubrey DeGrey with Michael Rae	2007	\$28.94	\$17.75	
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33569	OVERDOSE • by Jay Cohen, MD	—	\$24.95	\$16.95	
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Brand New! RESVERATROL PLUS PTEROSTILBENE IN ONE FORMULA!

In 2003, the **Life Extension Foundation**® introduced a purified **resveratrol** supplement that was later documented to favorably alter some of the changes in gene expression that cause us to age.

On **January 25, 2009**, CBS News **60 Minutes** featured an in-depth report on the multiple benefits that resveratrol may confer in slowing and even reversing certain aspects of aging.

What differentiates **Life Extension**®'s resveratrol: It provides 100% **standardized trans-resveratrol** plus a full spectrum of natural compounds from the grape that have demonstrated remarkable biological properties. Most products currently on the market contain varying quantities of **trans-** and **cis-resveratrol**, and fail to provide enough **trans-resveratrol** for optimal results. In addition, **Life Extension** has added **pterostilbene**, which researchers have found works in a **synergistic** fashion with resveratrol to activate one's "longevity genes."

Research funded by **Life Extension** has shown that a combination of low-dose (20 mg) **resveratrol** plus **grapeseed extract** induced many of the favorable **gene expression** changes seen in **calorie-restricted** animals. Other studies, however, have indicated that **higher doses** may be needed to obtain all of resveratrol's positive benefits including:

- **Improved Insulin Sensitivity**
- **Enhanced Mitochondrial Function**
- **Reduced Expression of Inflammatory Factors**
- **Protection Against the Toxic Effects of a High-Fat Diet**

CBS News and its affiliates are not connected with **Life Extension**® in any way, and do not endorse or sponsor its products.

Life Extension has meticulously evaluated published studies on **resveratrol** and **pterostilbene** to establish doses people might need to take to duplicate these remarkable laboratory findings. The results of **Life Extension**'s analysis yield a wide range of potentially effective doses of resveratrol and pterostilbene.

Highest Potency Resveratrol

The resveratrol *potency* you choose may be based on your current state of health and/or your desire to personally reach the upper limits of a healthy human life span. **Life Extension** offers a wide range of **standardized trans-resveratrol** potencies in combination with **pterostilbene** (which also favorably alters gene expression), **quercetin**, and other grape phytonutrients.

Optimized Resveratrol with Pterostilbene and the **new Calorie Restriction Mimetic Formula** provide the highest dosage of **trans-resveratrol** as well as a broad array of grape polyphenols, quercetin and **pterostilbene**.

All these Resveratrol Products are fortified with Pterostilbene!



Item #01413

Resveratrol with Pterostilbene

Provides **20 mg** of **trans-resveratrol** plus **120 mg** of **quercetin** to facilitate resveratrol absorption. **Pterostilbene** content (125 mcg) equivalent to over **5 cups of blueberries**. Recommended dosage is 1 capsule daily with or without food. Each bottle contains 60 vegetarian capsules.

Retail Price: \$24

**FOUR-BOTTLE SUPER SALE MEMBER PRICE:
>> \$14.85 PER BOTTLE**

(See each product description for amount of pterostilbene.)



Item #01411

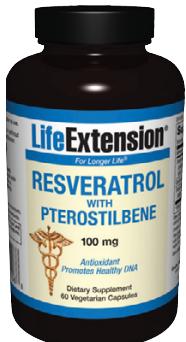
GrapeSeed Extract with Resveratrol & Pterostilbene

Provides 20 mg of *trans*-resveratrol plus 100 mg of grapeseed extract. Pterostilbene content (125 mcg) equivalent to over 5 cups of blueberries. Recommended dosage is 1 capsule daily with or without food. Each bottle contains 60 vegetarian capsules.

Retail Price: \$36

FOUR-BOTTLE SUPER SALE MEMBER PRICE:

>> \$22.95 PER BOTTLE



Item #01410

Resveratrol with Pterostilbene

Provides 100 mg of *trans*-resveratrol plus 120 mg of quercetin to facilitate resveratrol absorption. Pterostilbene content (250 mcg) equivalent to over 10 cups of blueberries. Recommended dosage is 1 capsule daily with or without food. Each bottle contains 60 vegetarian capsules.

Retail Price: \$36

FOUR-BOTTLE SUPER SALE MEMBER PRICE:

>> \$21.60 PER BOTTLE



Item #01409

Optimized Resveratrol with Pterostilbene

Provides in one capsule 250 mg of *trans*-resveratrol, 85 mg of grape plant polyphenols plus quercetin. Pterostilbene content (500 mcg) equivalent to over 20 cups of blueberries. Recommended dosage is 1 capsule daily with or without food. Each bottle contains 60 vegetarian capsules.

Retail Price: \$46

FOUR-BOTTLE SUPER SALE MEMBER PRICE:

>> \$27.90 PER BOTTLE

The Most Potent RESVERATROL-PTEROSTILBENE Formula

Among the most promising of caloric restriction mimics and enhancers are resveratrol, pterostilbene, quercetin, and grape seed polyphenols, along with black tea extract. These nutrients have been shown to generate many of the same effects in the body as caloric restriction, without significant dietary modification.¹⁻⁸ In particular, they help "mimic" caloric restriction's favorable impact on genes that influence the aging process.¹⁻⁸

A new Calorie Restriction Mimetic Formula includes resveratrol, higher levels of pterostilbene and quercetin plus, grape seed polyphenols, and black tea extract to provide even broader-spectrum gene expression support in one nutritional compound.

Two capsules of Calorie Restriction Mimetic Formula provide:

Trans-Resveratrol	250 mg
Trans-Pterostilbene	3 mg
Quercetin	150 mg
Black tea extract	300 mg
Grape seed polyphenols	50 mg



Item #01419

Each bottle contains 60 vegetarian capsules of the new Calorie Restriction Mimetic Formula.

Retail Price: \$36

FOUR-BOTTLE SUPER SALE MEMBER PRICE:

>> \$22.28 PER BOTTLE

1. Cell. 2006 Dec 15;127(6):1109-22.
2. Endocrinology. 2008 Jan;149(1):84-92.
3. Crit Care Med. 2004 Oct;32(10):2097-103.
4. J Agric Food Chem. 1999 Apr;47(4):1416-21.
5. Arch Pharm Res. 2002 Oct;25(5):561-71.
6. Nutr Cancer. 1999;35(1):80-6.
7. Anticancer Agents Med Chem. 2006 Sep;6(5):389-406.
8. Nature. 2006 Nov 16;444(7117):337-42.

To order these **Resveratrol products** or the new **Calorie Restriction Mimetic Formula**, call **1-800-544-4440** or visit www.LifeExtension.com



High Performance Supplement
Supports Thicker, Healthier

Hair

The adult scalp **loses** about **100 hairs every day**. Starting at the age of 40, hair follicles **shrink**, causing hair to grow back **thinner**, or worse, **not at all**. This **devastating** hair loss is experienced by **half** the population. While the prevention and treatment is complex, using a supplement with **tocotrienols** can be pivotal in addressing the various physiological mechanisms that lead to thinning hair.¹

Super Absorbable Tocotrienols combats the underlying causes of hair loss, supporting otherwise healthy hair follicles that genetic disposition, advancing age, high concentrations of DHT (the male hormone dihydrotestosterone), and sebum build-up may cause to weaken, degrade or shrink. **Super Absorbable Tocotrienols**, a natural, orally administered, bio-enhanced tocotrienol complex, can be taken to support **youthful hair thickness and growth**.

The Most Absorbable Tocotrienol Supplement Available Today

Through a revolutionary, patented delivery system, **Super Absorbable Tocotrienols** ensures optimal absorption²⁻⁷ and efficiently delivers the benefits of a unique proprietary complex of **phytonutrients** (squalene, phytosterols and trace

amounts of mixed carotene) naturally extracted together with tocotrienols to **support youthful hair growth, hair health and hair density**. In a study involving 30 volunteers who took tocotrienol supplements for eight months, nearly all the subjects showed significant improvement in hair thickness and density.¹

Those who want to obtain the benefits of this potent hair formula can order **Life Extension®'s Super Absorbable Tocotrienols**. The recommended dosage is one softgel taken two times daily with food. The retail price for a bottle of **Super Absorbable Tocotrienols** containing 60 softgels is \$30, but if a member buys four bottles during **Super Sale**, the price is reduced to **\$18.90 per bottle**.

To order **Super Absorbable Tocotrienols**,
call **1-800-544-4440**
or visit **www.LifeExtension.com**

Reference:

1. Randomized Clinical Trial of Tocotrienols. Supplementation vs. Placebo for Androgenetic Alopecia. Professor Yuen Kah Hay, B., School of Pharmaceutical Sciences, Universiti Sains, Malaysia. Submitted for publication 2009.
2. Antioxid Redox Signal. 2006 May-Jun;8(5-6):1059-68.
3. Mol Aspects Med. 2007 Oct-Dec;28(5-6):692-728.
4. J. of Pharmacy and Pharmacology. 200,53:67-71.
5. J. of Pharmacy and Pharmacology. 2003;55:53-58.
6. Intl. J. of Pharmaceutics. 2004,(281): 67-78.
7. US Patent No:US 6,596,306.



Item #01400

PROTECT EYE HEALTH AND FIGHT TIRED EYES



By absorbing blue-light, the **macula** protects delicate **photo-receptor cells** in the retina from light damage. The *density* of your macular pigment (composed of lutein, zeaxanthin, and meso-zeaxanthin) is essential to proper vision. Unfortunately, this density declines naturally over time. Some aging people also lose their ability to convert lutein into **meso-zeaxanthin** inside their macula.

Eating lots of lutein- and zeaxanthin-containing vegetables can help maintain the structural integrity of their macula. However, since **meso-zeaxanthin** is not part of the typical diet, it cannot be replaced except in supplement form.

Super Zeaxanthin with Lutein & Meso-zeaxanthin Plus Astaxanthin gives you therapeutic doses of **all three carotenoids** to help protect your precious eyesight plus a special ingredient that's scientifically proven to help fight tired eyes.

Fight Tired Eyes With Astaxanthin

If you're one of the more than 72 million people in America who use a computer daily at work,¹ you're probably familiar with eye fatigue. Staring at a fixed-distance object such as a computer screen for a long period of time can cause the muscles that focus your eyes (called the ciliary body) to tire or go into spasm. This can result in physical symptoms such as head discomfort, sensitivity to glare, tiredness, soreness, dryness, and blurry vision.



Item #01286

Super Zeaxanthin with Lutein & Meso-zeaxanthin Plus Astaxanthin contains a potent dose of **astaxanthin**, a carotenoid found in a red algae called *Haematococcus pluvialis*. Studies show that taking **astaxanthin** with other carotenoids protects against free-radical induced DNA damage, repairs UVA-irradiated cells, and inhibits inflammatory cell infiltration.²⁻⁵ Astaxanthin also helps support vascular health within the eye and improves visual acuity.⁴ Its fat-soluble nature offers protection to sensitive cells inside the eye.⁶

With its special combination of ingredients that targets eyestrain and protects visual acuity, **Super Zeaxanthin with Lutein & Meso-zeaxanthin Plus Astaxanthin** is truly an exciting breakthrough in eye care.

The retail price for a bottle containing 60 softgels of **Super Zeaxanthin with Lutein & Meso-zeaxanthin Plus Astaxanthin** is \$42. If a member buys four bottles during **Super Sale**, the price is reduced to just **\$25.65** per bottle.

**To order Super Zeaxanthin with Lutein & Meso-zeaxanthin Plus Astaxanthin,
call 1-800-544-4440 or visit www.LifeExtension.com**

References:

1. Report by the National Telecommunications and Information Administration of the US Department of Commerce. 2001 Sep.
2. *J Photochem Photobiol B*. 2007 Jul 27;88(1):1-10.
3. *J Photochem Photobiol B*. 2006 Dec 1;85(3):205-7.
4. *Ophthalmology*. 2008 Feb;115(2):324-333.e2.
5. *Invest Ophthalmol Vis Sci*. 2008 Apr;49(4):1679-85.
6. *Biochimica et Biophysica Acta*. 2001;1512: 251-8.

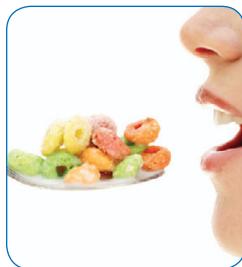
OptiLut® is a registered trademark of NutriScience Innovations, LLC. Lutein Plus® and MZ® are registered trademarks of Quantum Nutraceuticals. Licensed under U.S. Patents 5,523,494; 6,218,436; 6,329,432; and 6,504,067.



WHAT'S INSIDE

LifeExtension® Magazine

BLOCK ABSORPTION OF KILLER CARBOHYDRATES



An estimated **60 million** Americans are now pre-diabetic, the result of an impaired ability to process carbohydrates. A novel set of carbohydrate **enzyme-inhibitors** helps bring glucose levels into balance and improves age-related markers of health.

NEUTRALIZE LETHAL EFFECTS OF EXCESS CALORIE INGESTION



The most effective way to prevent disease and extend life span is to reduce calorie intake. In this groundbreaking issue of **Life Extension Magazine®** three methods are described to achieve the benefits of **calorie restriction** without having to endure strict dietary deprivation.

REVERSE MITOCHONDRIAL DYSFUNCTION



Age-related deterioration of the **mitochondria** in our cells is associated with an array of deadly conditions, ranging from senility to diabetes and heart failure. Cellular energizers have been identified that can help arrest and *reverse* progressive mitochondrial damage.

CUTTING-EDGE PROSTATE PROTECTION



Half of all men will experience debilitating **prostate** disorders sometime in their life. An advanced extraction technology now delivers even higher concentrations of the active ingredients in saw palmetto. Discover the nine natural compounds that provide maximum prostate support.

PLUS

Visit us at www.LifeExtension.com

HIGH HOMOCYSTEINE ASSOCIATED WITH SENILITY IN WOMEN
SOY SHOWS PROMISE FOR COLON CANCER PREVENTION
VITAMIN E BEATS DRUG TREATMENT FOR LIVER DISEASE
GREEN TEA HELPS PROTECT AGAINST ORAL CANCER