

Easy Ways to Prevent Colon Cancer

LifeExtension[®]

LifeExtension.com

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

August 2016

An Overlooked Step to Reverse Depression

Exclusive Report From Symposium on Sleep

Human Age Reversal Medical Conference

Removing Dead Cells On Your Skin Surface

Colon Cancer Rates Surge in the Elderly



PLUS—
Neutralize Damage Inflicted by Chronic Stress
Lutein Protects against Vascular Disease
Whey Protein May Help Manage Diabetes

THE MOST COMPREHENSIVE **MULTIVITAMIN** EVER FORMULATED!

LifeExtension Mix™

A unique, science-based formula with more than **80** bioactive ingredients including vitamins, minerals, and plant extracts for broad-spectrum health!



Life Extension Mix™ Tablets

Item #02055 • 315 tablets

	Retail Price	Your Price
1 bottle	\$80	\$60
4 bottles		\$52 each



Life Extension Mix™ Capsules

Item #02054 • 490 capsules

	Retail Price	Your Price
1 bottle	\$90	\$67.50
4 bottles		\$58 each



Life Extension Mix™ Powder

Item #02056 • 14.81 oz powder

	Retail Price	Your Price
1 bottle	\$80	\$60
4 bottles		\$52 each

PROVIDES:

- **VEGETABLE-FRUIT COMPLEX**
with 13 natural extracts
- **WATER-SOLUBLE VITAMINS**
and Enzymatic Activators with 11 ingredients including 5-MTHF
- **FAT-SOLUBLE VITAMINS**
with a natural mix of tocopherols
- **AMINO ACID COMPLEX**
including taurine and N-acetyl-L-cysteine
- **MINERAL COMPLEX**
composed of 12 key minerals from boron and selenium to chromium and zinc
- **CHOLINERGIC COMPLEX**
with choline, phosphatidylcholine, and inositol
- **FATTY ACID NUTRITION**
composed of medium-chain triglycerides
- **EACH BOTTLE PROVIDES A FIVE WEEK SUPPLY**

To order **Life Extension Mix™**, call toll-free **1-800-544-4440** or visit **www.LifeExtension.com**

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.

REPORTS



34 PROTECT AGAINST COLORECTAL CANCER

Colorectal cancer is the second leading cause of cancer death and the most preventable. Published studies demonstrate that healthy dietary practices and specific nutrients can help protect against colon cancer development.



46 YOUNGER-LOOKING SKIN

Microdermabrasion ranks among the most popular ways to restore youthful skin. A new **at-home** self-administered formula harnesses the unique properties of microdermabrasion utilizing ultra-fine **crystals**.



54 AGE-REVERSAL CONFERENCE

Human age reversal is evolving into scientific reality. **Life Extension Foundation**[®] is cosponsoring the **Revolution against Aging and Death** conference in San Diego on August 4-7, 2016. The focus is how age reversal can be accelerated through group collaboration.



64 ADAPTOGENS PROTECT AGAINST STRESS AND SUPPORT ADRENALS

Stress accelerates aging by disrupting cellular function. Natural extracts known as **adaptogens** protect the body against the assault of both internal and external stressors. Research has shown that four natural **adaptogens** can help neutralize damage caused by daily stress.



74 REPORT: SLEEP MEDICINE CONFERENCE

At the recent annual meeting of the **American Academy of Sleep Medicine and the Sleep Research Society**, scientists reported on new findings related to the detrimental impact of sleep deprivation. These include cognitive dysfunction, sleep apnea, insomnia, and hormone imbalance.

24 OVERLOOKED CAUSE OF DEPRESSION

A little known cause of depression are high levels of **homocysteine**. A controlled clinical study found that **5-MTHF**, a metabolically active form of **folate**, lowered homocysteine levels and significantly improved depression.



DEPARTMENTS



7 AS WE SEE IT: GOVERNMENT-INFLICTED COLON CANCER

Life Extension Magazine[®] publishes articles that sometimes contain material your government prefers you not read. They don't want this knowledge circulated because it increases their short term medical costs. Government groups are now recommending *against* colonoscopies for seniors. The tragic result will be colorectal cancers that go undetected and metastasize. Find out common-sense approaches to diagnosing early-stage colon lesions that can be removed before they transform into malignancy.



17 IN THE NEWS

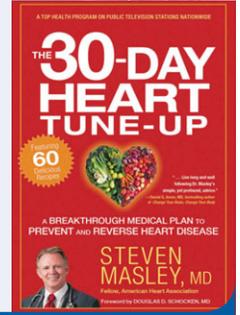
\$250 million cancer grant; supplements inhibit deafness; omega-3 lowers inflammation; ginseng improves diabetes; vitamin D reduces cardiovascular risk; and more.

87 RESEARCH UPDATE: NEWLY DISCOVERED LUTEIN BENEFITS

Lutein is well-known for protecting vision. New research reveals that lutein can also help prevent *cardio-metabolic* disorders such as heart disease, stroke, and diabetes.

93 AUTHOR INTERVIEW: THE 30-DAY HEART TUNE-UP

In his new book *The 30-Day Heart Tune-Up*, Steven Masley, MD, explains how you can reduce your risk factors for heart disease and stroke in just one month. He recently provided **Life Extension Magazine** with potentially lifesaving tips in an exclusive interview.





PUBLISHER • LE Publications, Inc.

CONNECT WITH LIFE EXTENSION ON THE WEB!



[Facebook.com/LifeExtension](https://www.facebook.com/LifeExtension)

For instant access to special offers and promotions, product news, and exclusive health and wellness information.



[Twitter.com/LifeExtension](https://twitter.com/LifeExtension)

For up-to-the-minute health tips, breaking industry news, and the latest updates in medical research.

Customer care is available to take your calls
24 hours a day, 7 days a week.

Visit the Life Extension Nutrition Center Store

- The Most Complete Line of Life Extension Supplements
- Blood Testing and Analysis
- Personal Consultation with Life Extension Product/Wellness Specialist



Nutrition Center of Florida, Inc.
5990 North Federal Highway,
Fort Lauderdale, FL 33308-2633 • 954-766-8144

Monday-Friday 9 am-8 pm,
Saturday 9 am-6 pm, Sunday 11 am-5 pm

EDITORIAL

Editor-in-Chief • Philip Smith
Executive Managing Editor • Renee Price
Medical Editor • Hernando Latorre, MD, MSc
Senior Copy Editor • Laurie Mathena
Senior Staff Writer • Michael Downey
Associate Editor • Garry Messick
Creative Director • Robert Vergara
Art Director • Alexandra Maldonado

CHIEF MEDICAL OFFICER

Steven Joyal, MD

VICE PRESIDENT OF PRODUCT INNOVATION & SCIENTIFIC DEVELOPMENT

Luke Huber, ND, MBA

SCIENTIFIC ADVISORY BOARD

Örn Adalsteinsson, PhD • John Boik, PhD • Aubrey de Grey, PhD
Frank Eichorn, MD • Deborah F. Harding, MD • Steven B. Harris, MD
Peter H. Langsjoen, MD, FACC • Dipnarine Maharaj, MD • Ralph W. Moss, PhD
Michael D. Ozner, MD, FACC • Jonathan V. Wright, MD, Xiaoxi Wei, PhD

CONTRIBUTORS

Ben Best • Maureen Fiona • Gary Goldfaden, MD • Robert Goldfaden
Steven Locke • Caroline Meyers • Timothy Rice • Kira Schmid, ND

ADVERTISING

Vice President of Marketing • Rey Searles • rsearles@lifeextension.com
National Advertising Manager • Leslie Stockton • 404-347-1755

VICE PRESIDENT OF SALES AND BUSINESS DEVELOPMENT

Ron Antriasian • rantriasian@lifeextension.com • 781-271-0089

CIRCULATION & DISTRIBUTION

Life Extension • 3600 West Commercial Blvd., Fort Lauderdale, FL 33309
Editorial offices: 954-766-8433 • fax: 954-491-5306

Customer Service: 800-678-8989

Email: customerservice@LifeExtension.com

Wellness specialists: 800-226-2370 • Wellness email: wellness@LifeExtension.com

At Life Extension Magazine® we value your opinion and welcome feedback.

Please mail your comments to *Life Extension Magazine®*,
Attn: Letters to the Editor, PO Box 407198, Fort Lauderdale, FL 33340
or email us: LEMagazine@LifeExtension.com

LIFE EXTENSION (ISSN 1524-198X) Vol. 22, No.8 ©2016 is published monthly except bi-monthly in April by LE Publications, Inc. at 3600 West Commercial Blvd., Fort Lauderdale, FL 33309-3338. **LE Publications, Inc. All rights reserved.** Published 13 times a year. Subscription rate: \$40 per year in the United States. US \$47 in Canada. US \$60 in other countries. Mail subscriptions or address changes to: LE Publications, Inc., P.O. Box 407198, Fort Lauderdale, FL 33340-7198, USA. Or phone us toll-free at: 1-800-841-5433. Canada Subscriptions: Publications mail agreement number 40028967. Return undeliverable Canadian addresses to PO Box 503, RPO West Beaver Creek, Richmond Hill, ON L4B4R6. You will be sent your first issue within six weeks after LE Publications, Inc. receives your subscription fee. Periodicals Postage paid at Fort Lauderdale, FL and at additional mailing offices. **POSTMASTER:** Send address changes to Life Extension, P.O. Box 407198, Ft. Lauderdale, Florida 33340-7198, USA. Printed in USA. The articles in this magazine are intended for informational purposes only. They are not intended to replace the attention or advice of a physician or other health-care professional. Anyone who wishes to embark on any dietary, drug, exercise, or other lifestyle change intended to prevent or treat a specific disease or condition should first consult with and seek clearance from a qualified health-care professional. **LEGAL NOTICE:** Health claims contained in articles and advertisements in this publication have not been approved by the FDA with the exception of FDA approved qualified health claims for calcium, antioxidant vitamins, folic acid and EPA and DHA omega-3 fatty acids, and selenium as noted where applicable. Life Extension® does not endorse any of the businesses or the products and/or services that may appear in advertisements for non-Life Extension branded products or services contained in Life Extension magazine® except to state that they are advertisers who may have paid Life Extension for placement of an advertisement in this publication. Life Extension disclaims any and all responsibilities or warranties as to the accuracy of information contained in advertisements for non-Life Extension branded products or services. For Canadian customers send change of address information and blocks of undeliverable copies to P.O. Box 1051, Fort Erie, ON L2A 6C7.

Dual-Action Support For Aging Joints!

As you grow older, age-related stiffness and discomfort in the joints becomes a fact of life. Activities once routine become a challenge as limited mobility hampers your every move.

You now have a more potent option to provide broad-spectrum support for aging joints.

ArthroMax® Advanced with UC-II® and AprèsFlex® is a multi-nutrient formula based on scientific data on natural support for joint health. The ArthroMax® formula provides more joint support than ever before, enhanced with two innovative, clinically validated ingredients: AprèsFlex® and UC-II®.

Better Absorption For Optimum Benefit

AprèsFlex® represents a quantum leap forward in the delivery to aging joints of **boswellia**, long prized for its ability to help with inflammatory issues. It is a superior inhibitor of the enzyme **5-lipoxygenase** or **5-LOX**.

Excess activity of 5-LOX results in the accumulation of **leukotriene B4**, a pro-inflammatory compound that affects aging joints. **Boswellia** has been shown to bind directly to the 5-LOX enzyme in our bodies, blocking it from facilitating production of pro-inflammatory leukotrienes.^{1,2}

AprèsFlex® boswellia *absorbs* into the blood **52%** **better** than previously available forms of boswellia, for superior effectiveness.

ArthroMax® Advanced With UC-II® And AprèsFlex®

In addition to AprèsFlex®, ArthroMax® contains a novel form of standardized cartilage: **UC-II®**. Data show it helps with immune issues that can impact joint discomfort and ease of motion in aging individuals.

ArthroMax® Advanced with UC-II® & AprèsFlex®

Item #01618 • 60 capsules

	Retail Price	Your Price
1 bottle	\$36	\$27
4 bottles		\$24 each

Non-GMO

AprèsFlex® is a registered trademark of Laila Nutraceuticals exclusively licensed to PL Thomas—Laila Nutra LLC. U.S. Patent No. 8,551,496 and other patents pending. FruiteX B® and OsteoBoron® are registered trademarks of VDF FutureCeuticals, Inc. U.S. Patent No. 5,962,049. UC-II® is a registered trademark of InterHealth N.I. U.S. patents 7,846,487; 7,083,820 and EPO patent EP 1435906B1; Canadian patent CA 2459981C; and Japanese patent JP 4800574B2.

References

1. *Wien Med Wochenschr.* 2002;152(15-16):373-8.
2. *J Ethnopharmacol.* 2006 Sep 19;107(2):249-53.



To order ArthroMax® Advanced with UC-II® and AprèsFlex® call 1-800-544-4440 or visit www.LifeExtension.com

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

LifeExtension[®]

Magazine

Gustavo Tovar Baez, MD, operates the Life Extension Clinic in Caracas, Venezuela. He is the first physician in Caracas to specialize in anti-aging medicine.

Ricardo Bernales, MD, is a board-certified pediatrician and general practitioner in Chicago, IL, focusing on allergies, bronchial asthma, and immunodeficiency.

Mark S. Bezzek, MD, FACP, FAARM, FAAEM, is board-certified in internal medicine, emergency medicine, and anti-aging/regenerative medicine. He is the director of Med-Link Consulting, which specializes in bioidentical hormone replacement therapy, natural alternatives, anti-aging, and degenerative diseases. He holds US patents for a multivitamin/mineral supplement, an Alzheimer's/dementia compilation, and a diabetic regimen.

Anna M. Cabeca, DO, FACOG, ABAARM, is a board-certified gynecologist and obstetrician, as well as board-certified in anti-aging and regenerative medicine, an expert in functional medicine, and an expert in women's health. She specializes in bioidentical hormone replacement therapy and natural alternatives, successful menopause and age-management medicine.

Thomas F. Crais, MD, FACS, a board-certified plastic surgeon, was medical director of the microsurgical research and training lab at Southern Baptist Hospital in New Orleans, LA, and currently practices in Sun Valley, ID.

William Davis, MD, is a preventive cardiologist and author of *Wheat Belly: Lose the Wheat, Lose the Weight* and *Find Your Path Back to Health*. He is also medical director of the online heart disease prevention and reversal program, *Track Your Plaque* (www.trackyourplaque.com).

Martin Dayton, MD, DO, practices at the Sunny Isles Medical Center in North Miami Beach, FL. His focus is on nutrition, aging, chelation therapy, holistic medicine, and oxidative medicine.

John DeLuca, MD, DC, is a 2005 graduate of St. George's University School of Medicine. He completed his internal medicine residency at Monmouth Medical Center in Long Branch, NJ, in 2008 and is board-certified by the American Board of Internal Medicine. Dr. DeLuca is a Diplomate of the American Academy of Anti-Aging Medicine and has obtained certifications in hyperbaric medicine, pain management, nutrition, strength and conditioning, and manipulation under anesthesia.

Sergey A. Dzugan, MD, PhD, was formerly chief of cardiovascular surgery at the Donetsk Regional Medical Center in Donetsk, Ukraine. Dr. Dzugan's current primary interests are anti-aging and biological therapy for cancer, cholesterol, and hormonal disorders.

Patrick M. Fratellone, MD, RH, is the founder and executive medical director of Fratellone Associates. He completed his internal medicine and cardiology fellowship at Lenox Hill Hospital in 1994, before becoming the medical director for the Atkins Center for Complementary Medicine.

Carmen Fusco, MS, RN, CNS, is a research scientist and clinical nutritionist in New York City who has lectured and written numerous articles on the biochemical approach to the prevention of aging and degenerative diseases.

Norman R. Gay, MD, is proprietor of the Bahamas Anti-Aging Medical Institute in Nassau, Bahamas. A former member of the Bahamian Parliament, he served as Minister of Health and Minister of Youth and Sports.

Mitchell J. Ghen, DO, PhD, holds a doctorate in holistic health and anti-aging and serves on the faculty of medicine at the Benemerita Universidad Autonoma De Puebla, Mexico, as a professor of cellular hemopoietic studies.

Gary Goldfaden, MD, is a clinical dermatologist and a lifetime member of the American Academy of Dermatology. He is the founder of Academy Dermatology of Hollywood, FL, and COSMESIS Skin Care.

Miguelangelo Gonzalez, MD, is a certified plastic and reconstructive surgeon at the Miguelangelo Plastic Surgery Clinic, Cabo San Lucas.

Garry F. Gordon, MD, DO, is a Payson, AZ-based researcher of alternative approaches to medical problems that are unresponsive to traditional therapies. He is president of the International College of Advanced Longevity Medicine.

Richard Heifetz, MD, is a board-certified anesthesiologist in Santa Rosa, CA, specializing in the delivery of anesthesia for office-based plastic/cosmetic surgery, chelation therapy, and pain management.

Roberto Marasi, MD, is a psychiatrist in Brescia and in Piacenza, Italy. He is involved in anti-aging strategies and weight management.

Maurice D. Marholin, DC, DO, is a licensed chiropractic physician and board-certified osteopathic family physician. While training at the University of Alabama, he completed Fellowships in Clinical Nutrition and Behavioral Medicine. He is currently in private practice in Clermont, FL.

Prof. Francesco Marotta, MD, PhD, of Montepapaleone Medical Center, Milan, Italy, is a gastroenterologist and nutrigenomics expert with extensive international university experience. He is also a consulting professor at the WHO-affiliated Center for Biotech & Traditional Medicine, University of Milano, Italy and hon. res. professor, Human Nutrition Dept, TWU, USA. He is the author of over 130 papers and 400 congress lectures.

Philip Lee Miller, MD, is founder and medical director of the Los Gatos Longevity Institute in Los Gatos, CA.

Michele G. Morrow, DO, FAAFP, is a board-certified family physician who merges mainstream and alternative medicine using functional medicine concepts, nutrition, and natural approaches.

Filippo Ongaro, MD, is board-certified in anti-aging medicine and has worked for many years as flight surgeon at the European Space Agency. He is considered a pioneer in functional and anti-aging medicine in Italy where he also works as a journalist and a writer.

Herbert Pardell, DO, FAAIM, practices internal medicine at the Emerald Hills Medical Center in Hollywood, FL. He is a medical director of the **Life Extension Foundation[®]**.

Lambert Titus K. Parker, MD, an internist and a board-certified anti-aging physician, practices integrative medicine from a human ecology perspective with emphasis on personalized brain health, biomarkers, genomics and total health optimization. He serves as the Medical Director of Integrative Longevity Institute of Virginia, a 501(c)3 Non-Profit Medical Research Institute. He also collaborates on education and research for Hampton Roads Hyperbaric Therapy.

Ross Pelton, RPh, PhD, CCN, is director of nutrition and anti-aging research for Intramedicine, Inc.

Patrick Quillin, PhD, RD, CNS, is a clinical nutritionist in Carlsbad, CA, and formerly served as vice president of nutrition for Cancer Treatment Centers of America, where he was a consultant to the National Institutes of Health.

Allan Rashford, MD, graduated from the University of Iowa Medical School. Upon completing medical training, he became chief of medicine at St. Francis Hospital in South Carolina, and he was later named president of the Charleston Medical Society.

Marc R. Rose, MD, practices ophthalmology in Los Angeles, CA, and is president of the Rose Eye Medical Group. He is on the staff of Pacific Alliance Medical Center, Los Angeles, and other area hospitals.

Michael R. Rose, MD, a board-certified ophthalmologist with the Rose Eye Medical Group in Los Angeles, CA, is on the staff of the University of Southern California and UCLA.

Ron Rothenberg, MD, is a full clinical professor at the University of California San Diego School of Medicine and founder of California HealthSpan Institute in San Diego, CA.

Roman Rozencwaig, MD, is a pioneer in research on melatonin and aging. He practices in Montreal, Canada, as research associate at Montreal General Hospital, Department of Medicine, McGill University.

Michael D. Seidman, MD, FACS, is the director of otolaryngology-head and neck surgery for the Bloomfield satellite of Henry Ford Health System (HFHS), Detroit, MI, co-director of the Tinnitus Center, and co-chair of the Complementary/ Alternative Medicine Initiative for HFHS.

Ronald L. Shuler, BS, DDS, CCN, LN, is involved in immunoncology for the prevention and treatment of cancer, human growth hormone secretagogues, and osteoporosis. He is board-certified in anti-aging medicine.

Paul Wand, MD, Fort Lauderdale, FL, is a clinical neurologist with special expertise in treating and reversing diabetic peripheral neuropathy and brain injuries from various causes.



Örn Adalsteinsson, PhD, is chairman of the **Life Extension®** Scientific Advisory board. He holds a master's and doctorate from the Massachusetts Institute of Technology (MIT). He has specialized in human therapeutics including vaccines, monoclonal antibodies, product development, nutraceuticals, formulations, artificial intelligence, hormones, and nutritional supplementation. He has also authored articles and contributed to peer-reviewed publications and served as an editor for the *Journal of Medicinal Food*.



John Boik, PhD, is the author of two books on cancer therapy, *Cancer and Natural Medicine* (1996) and *Natural Compounds in Cancer Therapy* (2001). He obtained his doctorate at the University of Texas Graduate School of Biomedical Sciences with research at the MD Anderson Cancer Center, focusing on screening models to identify promising new anti-cancer drugs. He conducted his postdoctoral training at Stanford University Department of Statistics. He is currently president of New Earth BioMed, a nonprofit cancer research corporation that studies mixtures of natural products.



Aubrey de Grey, PhD, is a biomedical gerontologist and Editor-in-Chief of *Rejuvenation Research*, the world's highest-impact peer-reviewed journal focused on intervention in aging. He received his BA and PhD from the University of Cambridge in 1985 and 2000 respectively. Dr. de Grey is a Fellow of both the Gerontological Society of America and the American Aging Association and sits on the editorial and scientific advisory boards of numerous journals and organizations.



Frank Eichorn, MD, is a urologist specializing in prostate cancer for 10 years. He has a private practice in Bad Reichenhall, Germany, and is prostate cancer consultant at the Urologische Klinik Castringius, Planegg, Munich. In his integrative approach to prostate cancer he works together with an international network of experts to improve treatment outcomes for prostate cancer patients with a special focus on natural and translational medicine.



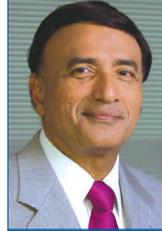
Deborah F. Harding, MD, is founder of the Harding Anti-Aging Center. She is double board-certified in internal medicine and sleep disorder medicine. She also earned the Cenegenics certification in age management medicine. She is a faculty member of the new University of Central Florida Medical School.



Steven B. Harris, MD, is president and director of research at Critical Care Research, a company that grew out of 21st Century Medicine in Rancho Cucamonga, CA. Dr. Harris participates in groundbreaking hypothermia, cryothermia, and ischemia research. His research interests include antioxidant and dietary-restriction effects in animals and humans.



Peter H. Langsjoen, MD, FACC, is a cardiologist specializing in congestive heart failure, primary and statin-induced diastolic dysfunction, and other heart diseases. A leading authority on coenzyme Q10, Dr. Langsjoen has been involved with its clinical application since 1983. He is a founding member of the executive committee of the International Coenzyme Q10 Association, a fellow of the American College of Cardiology, and a member of numerous other medical associations.



Dipnarine Maharaj MD, MB, ChB, FRCP (Glasgow), FRCP (Edinburgh), FRCPPath., FACP

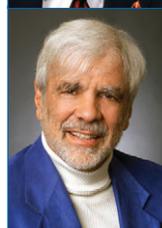
Dr. Dipnarine Maharaj is the Medical Director of the South Florida Bone Marrow Stem Cell Transplant Institute and is regarded as one of the world's foremost experts on adult stem cells. He received his medical degree in 1978 from the University of Glasgow Medical School, Scotland. He completed his internship and residency in Internal Medicine and Hematology at the University's Royal Infirmary.



Ralph W. Moss, PhD, is the author of books such as *Antioxidants Against Cancer*, *Cancer Therapy*, *Questioning Chemotherapy*, and *The Cancer Industry*, as well as the award-winning PBS documentary *The Cancer War*. Dr. Moss has independently evaluated the claims of various cancer treatments and currently directs The Moss Reports, an updated library of detailed reports on more than 200 varieties of cancer diagnoses.



Michael D. Ozner, MD, FACC, FAHA, is a board-certified cardiologist who specializes in cardiovascular disease prevention. He serves as medical director for the Cardiovascular Prevention Institute of South Florida and is a noted national speaker on heart disease prevention. Dr. Ozner is also author of *The Great American Heart Hoax*, *The Complete Mediterranean Diet* and *Heart Attack Proof*. For more information visit www.drozner.com.



Jonathan V. Wright, MD, is medical director of the Tahoma Clinic in Tukwila, WA. He received his MD from the University of Michigan and has taught natural biochemical medical treatments since 1983. Dr. Wright pioneered the use of bioidentical estrogens and DHEA in daily medical practice. He has authored or co-authored 14 books, selling over 1.5 million copies.



Xiaoxi Wei, PhD, is a chemist expert in supramolecular assembly and development of synthetic transmembrane nanopores with distinguished selectivity via biomimetic nanoscience. She has expertise in ion channel function and characterization. She founded X-Therma Inc., a company developing a radical new highway towards non-toxic, hyper-effective antifreeze agents to fight unwanted ice formation in regenerative medicine and reduce mechanical icing.

RESTORE YOUTHFUL COGNITIVE HEALTH



Dopamine is a neurotransmitter that regulates **mood** and **cognition**.

As we age, **dopamine** levels in our brain decline due to excess levels of the **MAO-B enzyme**.

Wild green oat extract inhibits the MAO-B enzyme to promote healthy dopamine levels.¹⁻⁴

The **wild green oat extract** in **Dopa-Mind™** is for aging individuals who wish to:

- **Maintain youthful mental performance**
- **Revive cognitive health**
- **Support longevity**

Dopa-Mind™

Item #02006 • 60 vegetarian tablets

	Retail Price	Your Price
1 bottle	\$48	\$36
4 bottles		\$32 each

To order **Dopa-Mind™**, call
1-800-544-4440
or visit **www.LifeExtension.com**

References

1. *J Neurosci*. 2013 Jan 2;33(1):286-91.
2. *Psychol Med*. 2011 Feb;41(2):419-29.
3. *Am J Psychiatry*. 1998 Mar;155(3):344-9.
4. *J Altern Complement Med*. 2011 Jul;17(7):635-7.

Non-GMO

Neuravena® is a registered trademark of Frutarom.

Government Inflicted Colon Cancer



BY WILLIAM FALOON

Life Extension Magazine[®] publishes articles that sometimes contain material your government prefers you not read. The reason they don't want this knowledge circulated is that it increases their medical costs in the short term.

The dilemma is that governments cannot **afford** their healthcare obligations.¹⁻⁸

This happened because free or subsidized medical care was promised, but costs are soaring and people are living longer.⁹⁻¹¹

Medical technology is improving, but not fast enough to bring about needed price reductions.¹²⁻¹⁴ The solution will be **curative** therapies that slash today's high price of chronic illness, but we are not there yet.

Governments are fighting this cost crisis by reducing healthcare outlays. The human group most targeted is over age **74**.

That age group represents many people reading this article, which obligates me to refute what is an illogical **rationing** of a **lifesaving** diagnostic procedure.

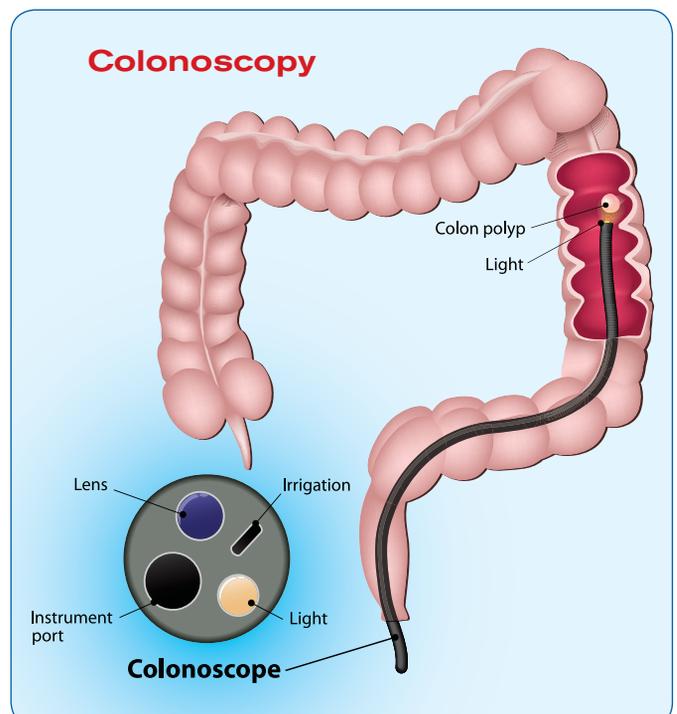
Earlier this year, a **Canadian** government task force recommended against screening **colonoscopies** for citizens aged 50 years and older. They suggested that a **fecal occult blood test** or **sigmoidoscopy** were sufficient. For people over **74**, the task force recommended **no** screening for polyps or early-stage colon cancer.¹⁵

The **US Preventive Service Task Force** made a similar recommendation in **2008** for people over **75**.¹⁶ This may be changing to let those aged **76-85** make

the choice. As of May 2016, however, draft guidance from the US Preventive Service Task Force recommends **against** routine screening for colorectal cancer in individuals **86** or over.¹⁷

You're going to learn about a substantial **increase** in **metastatic colon cancers** that will be inflicted by these senseless governmental edicts.

The encouraging news we report this month are documented ways to reduce your risk of developing **colon cancer** and curtail **metastasis** in case you are ever diagnosed with this malignancy.



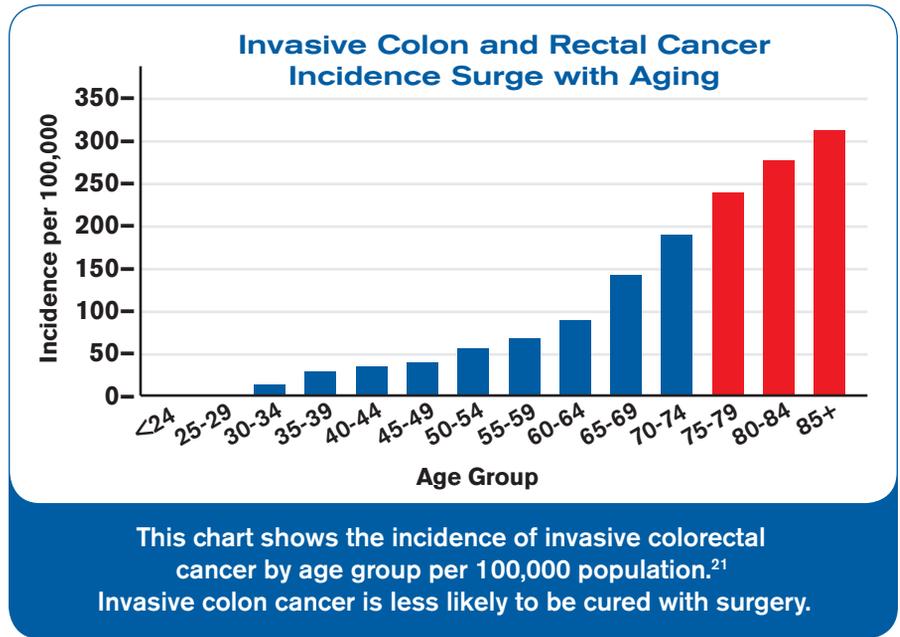
Humans are living longer, yet government policies seek to preclude older groups from accessing diagnostic testing.¹⁸⁻²⁰

New government recommendations make it appear that once people age past **74** their risk for diseases like **colon cancer** mystically disappears. As the chart on this page clearly shows, incidence of invasive **colon** and **rectal cancer** spirals upwards as people grow older.²¹

There is no reason to deprive a person over **74** of colon cancer **screening**. It is proven to be effective in detecting early-stage lesions before they progress and spread to the liver, lungs, and other organs.

If a polyp or other suspicious lesion is detected during a colonoscopy, it can usually be removed and biopsied without further procedures. Early removal of polyps reduces future colon cancers.²²

A **sigmoidoscopy** typically evaluates only **one-third** of the colon, whereas a **colonoscopy** examines the entire colon.²³⁻²⁶ Since the bowel preparation requirements are similar,²⁷⁻²⁹ we at **Life Extension®** have



long advocated **colonoscopy** over **sigmoidoscopy**.

We disagree with government recommendations regarding how often one should have a colonoscopy. We suggest individuals over age **40** consider a **colonoscopy** more frequently than current guidelines of once every **10 years**.¹⁶

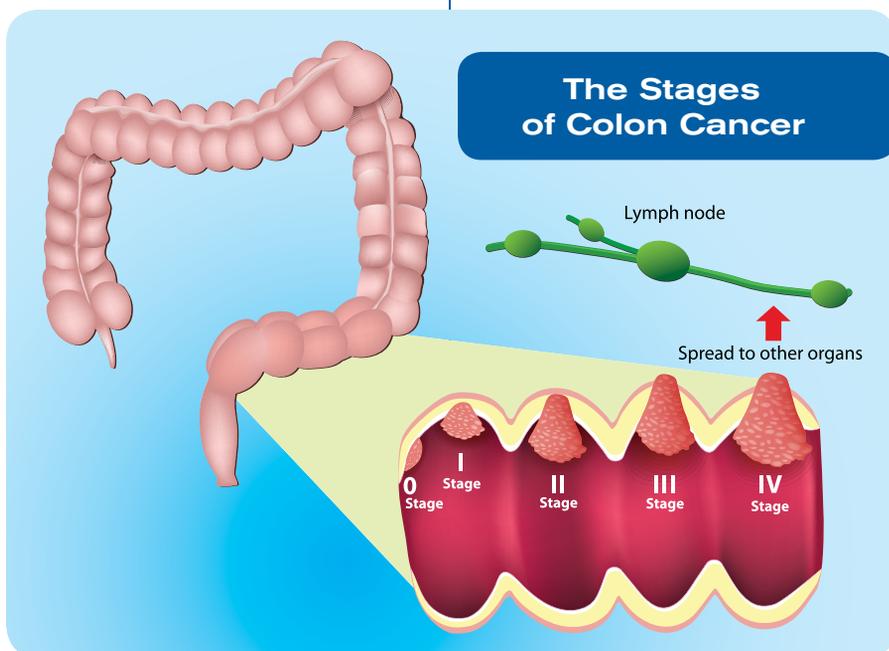
Troubling Statistics

Colorectal cancer usually (but not always) develops slowly,^{30,31} over a period of 10 to 20 years.^{32,33} Most cancers begin as a noncancerous growth called a **polyp** that develops on the inner lining of the colon or rectum.³⁴

When caught at these early stages, polyps and precancerous lesions can be easily removed during the **colonoscopy** procedure.³⁵⁻³⁷ So screening for colorectal cancer is not only diagnostic, but **therapeutic** when suspicious lesions/polyps are removed during the colonoscopy.^{38,39}

While colonoscopies have spared countless lives since widespread screening began, about **50,000** Americans still perish each year from metastatic **colorectal cancer**.^{40,41}

Most of these deaths would not have occurred had the victim utilized a proven diagnostic procedure. For example, in 2010 only **59%** of people age 50 or older reported having received colorectal cancer screening consistent with guide-



lines.⁴² Yet about **1 in 18** Americans will be diagnosed with colorectal cancer sometime in their life.⁴³

Colorectal cancer is the second leading cause of cancer death in the United States. The **American Cancer Society** estimates that about **135,000** people will be diagnosed with colorectal cancer this year.⁴⁴

In 2014, about **50,310** people died from the disease.⁴² This number represents a decline in colorectal deaths from previous years and probably reflects removal of polyps and other precancerous lesions during **colonoscopy**, along with finding early stage malignancies during colonoscopy that are curable.^{45,46}

With government groups now recommending *against* colonoscopies and the inconvenience of having it done, an increasing number of unwary humans will not avail themselves to this proven lifesaving procedure.

How Effective Are Colonoscopies?

Initial studies on high-risk patients undergoing colonoscopy indicated a reduced risk of colon cancer incidence by as much as **90%**.⁴⁷

An editorial published in *Annals of Internal Medicine* challenged this assertion. While acknowledging the importance of screening, it suggested that physicians inform patients that for lesions located in the descending or left side of the colon, high quality colonoscopies reduce the mortality risk by **60%** to **70%** and that limited data exists for reduction of right colon cancer mortality risk.⁴⁸

What we know is colon cancer diagnosis and death from metastatic disease remains rampant. Even curative treatment often inflicts permanent side effects.^{49,50} This means

you should not wait for symptoms to occur that necessitate surgical removal of a section of your bowel and follow up chemotherapy as a precaution against metastasis.

In seeking to identify the hard facts, we are disappointed by the lack of long-term high-quality studies to validate how many lives are really spared by colonoscopy, and what time intervals are ideal for average-risk individuals to have colonoscopies performed.⁵¹

I've therefore taken a **commonsense** approach in writing this editorial as to what health-conscious individuals should consider in order to reduce their risk of this common malignancy.

Comparing Screening Techniques

A number of investigative studies have been done to compare the benefits of **colonoscopy**, **sigmoidoscopy** and/or **fecal occult blood testing**.⁵²⁻⁵⁶

A **fecal occult blood test** is done at home by swiping a tiny amount of stool onto a card for

three consecutive days and delivering it to your doctor's office or sending to a laboratory. It is recommended that this test be done **annually**.⁵⁷⁻⁵⁹

If traces of blood are present in the stool then additional diagnostics are prescribed, typically colonoscopy.⁵⁹⁻⁶¹

When compared to **colonoscopies** done every **ten years**, high-sensitivity **fecal occult blood tests** performed **every year** showed similar benefits in detecting colorectal cancers.⁶²

Since **fecal occult blood tests** cost relatively little, health authorities view them as a way of reducing the higher cost of **colonoscopy** screening.

We take issue with this comparison data because we believe colonoscopies should ideally be done more frequently than every **10 years**.

When governments base recommendations to use **fecal occult blood tests** in lieu of **colonoscopy**, and compare it to **10-year** colonoscopy intervals, they are missing a potential benefit of more frequent colonoscopy screening.



Fecal occult blood test kit

We think **colonoscopies** performed about every **5 years** will yield superior results compared to annual **fecal occult blood testing**.

My reasons for suggesting more frequent colonoscopies include poor quality of prior colonoscopies and the benefit of more frequent removal of polyps and pre-malignant lesions (adenomas).^{63,64} There is also a small risk of what is termed “interval cancer,” which means a fast growing malignancy that occurs between colonoscopy screenings.^{65,66}

We’ve reprinted a chart at the end of this article that shows what is currently being recommended as far as **time intervals** between colonoscopy procedures. As I have written in the past, these kinds of recommendations are based on typical population groups. As a reader of this magazine, you expect better than “average” as far as your longevity is concerned.

Common-Sense Suggestions

The *Canadian Task Force on Preventive Health Care* is recommending **fecal occult blood screening** or sigmoidoscopy in lieu of **colonoscopy** for non-high-risk people aged **50-74**.⁶⁸ This recommendation is based on data showing only slightly better detection with **colonoscopy** compared to the cheaper **fecal occult blood** tests.

What’s being overlooked, however, is that when **blood** is detected in the stool, the patient may have been growing **polyps** or developing colorectal cancer years prior to this. We think more frequent colonoscopies will lead to more polyps and other suspicious lesions being identified and removed before early stage colon cancer develops.

While polyps can also bleed, and the blood can show up in the

stool, their diagnosis and removal still requires a scoping procedure of the colon.

Health “authorities” are now proposing that people forgo **colonoscopy** screening and instead deliver each year to their doctor’s office three consecutive days of stool samples to be **fecal occult blood tested**.

If traces of blood are detected, the patient is then usually told to undergo a **colonoscopy** to ascertain what is causing blood to appear in their stool.⁶⁹

Once a person reaches age **75**, the Canadian task force says no screening is needed,⁷⁰ which is analogous to what the **US Preventive Services Task Force** published in **2008**.¹⁶ We at **Life Extension** disagree with these recommendations for most healthy individuals.

We suggest that low-risk individuals over age 40 consider having a **colonoscopy** about every **5 years** to reduce risk of developing advanced or metastatic colorectal cancer.

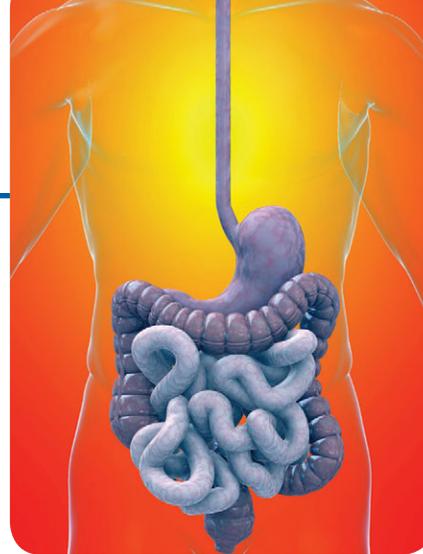
What’s missing from all these analyses is direct evidence, i.e. randomized, controlled trials over long time periods on large human populations showing how frequently colonoscopies should be performed.⁷¹

Rather than wait for results from conclusive trials, the common-sense approach we advocate is colonoscopies done more frequently than every **10 years** for low-risk individuals.

Putting This in Perspective

By the time the typical person (who neglects their health) ages past 74, they often suffer from so many chronic disorders that their lifespans are limited.

It may thus be understandable for a government task force to



recommend against colonoscopy for the **general population** over age 74 as they may not derive much real-world benefit.

When it comes to the general population, the government sees cost savings in sigmoidoscopy, which does not always require sedation as do most colonoscopies.^{72,73} Sedation adds costs to any medical procedure. Although serious complication rates are very low for both colonoscopies and sigmoidoscopies, colonoscopies have higher rates of serious complications compared to sigmoidoscopies.⁶² To save money in the short term, governments are looking to reduce the number of colonoscopies performed.

Readers of this publication are different. We take extraordinary measures to slow aging, prevent disease, and extend our healthy longevity. We don’t fit in with the “average” American or Canadian, who lets nature and deadly lifestyle choices accelerate degenerative processes.

We’re also privileged to know about **age reversal** research that may enable us to purge our bodies of multiple pathologies while our cells simultaneously become biologically younger. (Refer to article on page 54 of this month’s issue.)

Life Extension participants have huge incentives to protect against common disorders of today such as **colorectal cancer**, so they can remain healthy and alive to benefit from pending breakthrough biomedical advances.

What Should You Do?

Colonoscopies do not guarantee one will never die of colon cancer. The encouraging news is that the healthy lifestyle programs most of you practice have been shown to reduce (but not eliminate) colorectal cancer risk.

Colon cancer prevalence is so high that we suggest low-risk individuals should make an effort to undergo colonoscopy about every **5 years**. Those with certain risk factors like inflammatory bowel diseases or history of adenomas may want to have colonoscopies more frequently.

Governmental edicts that people over **age 74** should forgo colonoscopy may not apply to individuals who plan on living far longer, healthier lifespans.

Conventional medicine is recognizing the role of aspirin,⁷⁴⁻⁷⁶

calcium,^{77,78} vitamin D,^{79,80} and healthy dietary choices in reducing colorectal cancer risk. An article in this month's issue of *Life Extension Magazine* delves into this topic in detail.

The point I need to emphasize is that while many of the supplements you are taking have shown a profound effect in reducing colorectal cancer risk, they do not protect against the disease completely.

In addition, certain upper digestive tract malignancies are increasing in prevalence, such as **esophageal cancer**.⁸¹ If you are going to schedule a colonoscopy and be sedated anyway, it makes sense for people at risk to ask their gastroenterologist to perform an **endoscopy** and **colonoscopy** at the same visit.⁸²

By having these two procedures performed, lesions can be detected from the throat to the anus when

they are likely to be treatable before a **metastatic** disease manifests.

I realize these screening suggestions are in opposition to governmental guidelines that seek to cut back on healthcare outlays. There are also side-effect risks associated with any invasive diagnostic procedure such as colonoscopy and endoscopy.

My mission, however, is to keep you alive and healthy. I yearn for the day when the scourge of **cancer** will become a relic of the past, just as **smallpox** is today. Until that time, we should remain vigilant, even if we happen to be over **74 years** of age.

For longer life,



William Faloon

Conventional Recommendations for Surveillance and Screening Intervals in Individuals with Baseline Average Risk⁶⁷

Baseline colonoscopy: most advanced finding(s)	Recommended surveillance interval (years)	Quality of evidence supporting the recommendation
No polyps	10	Moderate
Small (<10 mm) hyperplastic polyps in rectum or sigmoid	10	Moderate
1–2 small (<10 mm) tubular adenomas	5–10	Moderate
3–10 tubular adenomas	3	Moderate
>10 adenomas	<3	Moderate
One or more tubular adenomas ≥10 mm	3	High
One or more villous adenomas	3	Moderate
Adenoma with HGD [high-grade dysplasia]	3	Moderate
Serrated lesions		
Sessile serrated polyp(s) <10 mm with no dysplasia	5	Low
Sessile serrated polyp(s) ≥10 mm, or sessile serrated polyp with dysplasia, or traditional serrated adenoma	3	Low

References

- Eklund W. Japan and its healthcare challenges and potential contribution of neonatal nurse practitioners. *J Perinat Neonatal Nurs.* 2010;24(2):155-66.
- Poses RM. A cautionary tale: the dysfunction of American health care. *Eur J Intern Med.* 2003;14(2):123-30.
- Fletcher T. The impact of physician entrepreneurship on escalating health care costs. *J Am Coll Radiol. JACR.* 2005;2(5):411-4.
- Suter E, Oelke ND, Adair CE, et al. Ten key principles for successful health systems integration. *Healthc Q. (Toronto, Ont.).* 2009;13 Spec No:16-23.
- Thomas D, Sarangi BL, Garg A, et al. Closing the health and nutrition gap in Odisha, India: A case study of how transforming the health system is achieving greater equity. *Soc Sci Med. (1982).* 2015;145:154-62.
- Okello DO, Lubanga R, Guwatudde D, Sebina-Zziwa A. The challenge to restoring basic health care in Uganda. *Soc Sci Med. (1982).* 1998;46(1):13-21.
- Wang C, Rao K, Wu S, et al. Health care in China: improvement, challenges, and reform. *Chest.* 2013;143(2):524-31.
- Grosios K, Gahan PB, Burbidge J. Overview of healthcare in the UK. *EPMA J.* 2010;1(4):529-34.
- Muka T, Imo D, Jaspers L, et al. The global impact of non-communicable diseases on healthcare spending and national income: a systematic review. *Eur J Epidemiol.* 2015;30(4):251-77.
- Kreatsoulas C, Anand SS. The impact of social determinants on cardiovascular disease. *Can J Cardiol.* 2010;26 Suppl C:8c-13c.
- Salomon JA, Wang H, Freeman MK, et al. Healthy life expectancy for 187 countries, 1990-2010: a systematic analysis for the Global Burden Disease Study 2010. *Lancet.* 2012;380(9859):2144-62.
- Kumar RK. Technology and healthcare costs. *Ann Pediatr Cardiol.* 2011;4(1):84-6.
- Arora S, Thornton K, Komaromy M, et al. Demonopolizing medical knowledge. *Acad Med.* 2014;89(1):30-2.
- Ventola CL. Challenges in evaluating and standardizing medical devices in health care facilities. *PT.* 2008;33(6):348-59.
- Care CTFoPH. Recommendations on screening for colorectal cancer in primary care. *CMAJ.* 2016;188(5):340-8.
- Available at: <http://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/colorectal-cancer-screening>. Accessed May 10, 2016.
- Available at: <http://www.uspreventiveservicestaskforce.org/Page/Document/draft-recommendation-statement38/colorectal-cancer-screening2>. Accessed May 10, 2016.
- Barberis M. America's elderly: policy implications. *Popul Bull.* 1981;35(4 Supplement):1-13.
- Christensen K, Doblhammer G, Rau R, et al. Ageing populations: the challenges ahead. *Lancet.* 2009;374(9696):1196-208.
- Jin K, Simpkins JW, Ji X, Leis M, et al. The critical need to promote research of aging and aging-related diseases to improve health and longevity of the elderly population. *Aging Dis.* 2015;6(1):1-5.
- Available at: http://seer.cancer.gov/archive/csr/1975_2012/browse_csr.php?sectionSEL=6&pageSEL=sect_06_table.11.html. Accessed May 11, 2016.
- Citarda F, Tomaselli G, Capocaccia R, et al. Efficacy in standard clinical practice of colonoscopic polypectomy in reducing colorectal cancer incidence. *Gut.* 2001;48(6):812-5.
- Macafee DAL, Scholefield JH. Population based endoscopic screening for colorectal cancer. *Gut.* 2003;52(3):323-6.
- Boland CR, Demarco DC. Invited commentary: Preventing colon cancer: looking over the horizon. *Proceedings (Baylor University. Medical Center).* 2003;16(3):344-5.
- Simon JB. Screening colonoscopy: is it time? *CMAJ.* 2000;163(10):1277-8.
- Church JM. Colon cancer screening update and management of the malignant polyp. *Clin Colon Rectal Surg.* 2005;18(3):141-9.
- Available at: <http://www.cancer.org/healthy/findcancerearly/examandtestdescriptions/faq-colonoscopy-and-sigmoidoscopy>. Accessed May 10, 2016.
- Available at: <http://www.mayoclinic.org/tests-procedures/flexible-sigmoidoscopy/basics/how-you-prepare/prc-20014697>. Accessed May 10, 2010.
- Available at: <http://www.mayoclinic.org/tests-procedures/colonoscopy/basics/how-you-prepare/prc-20013624>. Accessed May 10, 2016.
- Brenner H, Kloor M, Pox CP. Colorectal cancer. *Lancet.* 2014;383(9927):1490-502.
- Stathopoulos GP. Survival of untreated advanced colorectal cancer patients. *Oncol Lett.* 2011;2(4):731-3.



32. Available at: <http://www.cancer.org/acs/groups/cid/documents/webcontent/003170-pdf.pdf>. Accessed May 11, 2016.
33. Winawer SJ, Zauber AG. The advanced adenoma as the primary target of screening. *Gastro Endosc Clin N Am*. 2002;12(1):1-9, v.
34. Loeve F, Boer R, Zauber AG, et al. National polyp study data: evidence for regression of adenomas. *Int J Cancer*. 2004 Sept. 10;111(4):633-9.
35. Aarons CB, Shanmugan S, Bleier JJ. Management of malignant colon polyps: current status and controversies. *World J Gastroenterol*. 2014;20(43):16178-83.
36. Rutter MD. Evolving protocols in colorectal cancer surveillance. *Gastroenterol Hepatol (NY)*. 2008;4(2):114-6.
37. Delavari A, Mardan F, Salimzadeh H, et al. Characteristics of colorectal polyps and cancer; a retrospective review of colonoscopy data in iran. *Middle East J Dig Dis*. 2014;6(3):144-50.
38. Bari Z, Fakhri H, Sardarian H. Large bowel obstruction after colonoscopy; a case report. *Middle East J Dig Dis*. 2015;7(4):253-6.
39. Geiger TM, Ricciardi R. Screening options and recommendations for colorectal cancer. *Clin Col Rectal Surg*. 2009;22(4):209-17.
40. Litvak DA, Malad S, Wascher RA, et al. Laparoscopic splenectomy in colorectal cancer patients with chemotherapy-associated thrombocytopenia due to hypersplenism. *Case Rep Oncol*. 2012;5(3):601-7.
41. Fong Y, Cohen AM, Fortner JG, et al. Liver resection for colorectal metastases. *J Clin Oncol*. 1997;15(3):938-46.
42. Available at: <http://www.cancer.org/research/cancerfactsstatistics/colorectal-cancer-facts-figures>. Accessed May 11, 2016.
43. Amersi F, Agustin M, Ko CY. Colorectal cancer: epidemiology, risk factors, and health services. *Clin Colon Rectal Surg*. 2005;18(3):133-40.
44. Available at: <http://www.cancer.org/cancer/colonandrectumcancer/detailedguide/colorectal-cancer-key-statistics>. Accessed May 11, 2016.
45. Available at: <http://seer.cancer.gov/stat-facts/html/colorect.html>. Accessed May 11, 2016.
46. Available at: <http://www.cancer.org/acs/groups/content/documents/document/ac-spc-042280.pdf>. Accessed May 11, 2016.
47. Winawer SJ, Zauber AG, Ho MN, et al. Prevention of colorectal cancer by colonoscopic polypectomy. The National Polyp Study Workgroup. *The N Engl J Med*. 1993;329(27):1977-81.
48. Ransohoff DF. How much does colonoscopy reduce colon cancer mortality? *Ann Intern Med*. 2009 Jan 6;150(1):50-2.
49. Hellinger MD, Santiago CA. Reoperation for recurrent colorectal cancer. *Clin Colon Rectal Surg*. 2006 Nov;19(4):228-36.
50. Denlinger CS, Barsevick AM. The challenges of colorectal cancer survivorship. *J Natl Compr Canc Netw: JNCCN*. 2009;7(8):883-93; quiz 894.
51. Available at: <http://canadiantaskforce.ca/files/crc-screeningfinal031216.pdf>. Accessed May 12, 2016.
52. Holme O, Bretthauer M, Frøtheim A, et al. Flexible sigmoidoscopy versus faecal occult blood testing for colorectal cancer screening in asymptomatic individuals. *Cochrane Database Syst Rev*. 2013 Oct 1;9:Cd009259.
53. O'Leary BA, Olynyk JK, Neville AM, et al. Cost-effectiveness of colorectal cancer screening: comparison of community-based flexible sigmoidoscopy with fecal occult blood testing and colonoscopy. *J Gastroenterol Hepatol*. 2004;19(1):38-47.
54. Cheng TI, Wong JM, Hong CF, et al. Colorectal cancer screening in asymptomatic adults: comparison of colonoscopy, sigmoidoscopy and fecal occult blood tests. *J Formos Med Assoc*. 2002;101(10):685-90.
55. Graser A, Stieber P, Nagel D, et al. Comparison of CT colonography, colonoscopy, sigmoidoscopy and faecal occult blood tests for the detection of advanced adenoma in an average risk population. *Gut*. 2009;58(2):241-8.
56. Sung JJ, Chan FK, Leung WK, et al. Screening for colorectal cancer in Chinese: comparison of fecal occult blood test, flexible sigmoidoscopy, and colonoscopy. *Gastroenterology*. 2003;124(3):608-14.
57. Available at: <https://www.nlm.nih.gov/medlineplus/ency/article/003393.htm>. Accessed May 12, 2016.
58. Barry MJ. Fecal occult blood testing for colorectal cancer: a perspective. *Ann Oncol*. 2002;13(1):61-4.
59. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK445/>. Accessed May 12, 2016.
60. Available at: <https://www.ucsfhealth.org/tests/007008.html>. Accessed May 12, 2016.
61. Hubbard RA, Johnson E, Hsia R, et al. The cumulative risk of false-positive fecal occult blood test after 10 years of colorectal cancer screening. *Cancer epidemiol Biomarkers Prev*. 2013 Sep;22(9):1612-9.
62. Available at: <http://www.uspreventiveservicestaskforce.org/Page/Document/RecommendationStatementFinal/colorectal-cancer-screening>. Accessed May 12, 2016.
63. Jang JY, Chun HJ. Bowel preparations as quality indicators for colonoscopy. *World J Gastroenterol*. 2014 March 21;20(11):2746-50.
64. Chokshi RV, Hovis CE, Hollander T, et al. Prevalence of missed adenomas in patients with inadequate bowel preparation on screening colonoscopy. *Gastrointest Endosc*. 2012 Jun;75(6):1197-203.
65. Holt PR, Kozuch P, Mewar S. Colon cancer and the elderly: from screening to treatment in management of GI disease in the elderly. *Best pract Res Clin Gastroenterol*. 2009;23(6):889-907.
66. Samadder NJ, Curtin K, Tuohy TM, et al. Characteristics of missed or interval colorectal cancer and patient survival: a population-based study. *Gastroenterology*. 2014;146(4):950-60.
67. Lieberman DA, Rex DK, Winawer SJ, et al. Guidelines for colonoscopy surveillance after screening and polypectomy: a consensus update by the US Multi-Society Task Force on Colorectal Cancer. *Gastroenterology*. 2012;143(3):844-57.
68. Available at: <http://canadiantaskforce.ca/news/2016-02-22/canadian-medical-association-journal-publishes-ctfphcs-guideline-on-colorectal-cancer-screening-in-adults/>. Accessed May 12, 2016.
69. McLoughlin MT, Telford JJ. Positive occult blood and negative colonoscopy—should we perform gastroscopy? *Can J Gastroenterol*. 2007 Oct;21(10):633-6.
70. Available at: <http://canadiantaskforce.ca/ctfphc-guidelines/2015-colorectal-cancer/>. Accessed May 12, 2016.
71. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK65825/>. Accessed May 12, 2016.
72. Available at: <https://www.nlm.nih.gov/medlineplus/colonoscopy.html>. Accessed May 12, 2016.
73. Aljebreen AM, Almadi MA, Leung FW. Sedated vs unsedated colonoscopy: A prospective study. *World J Gastroenterol*. 2014 May;20(17):5113-8.
74. Drew DA, Cao Y, Chan AT. Aspirin and colorectal cancer: the promise of precision chemoprevention. *Nat Rev Cancer*. 2016;16(3):173-86.
75. Wakeman C, Keenan J, Eteuati J, et al. Chemoprevention of colorectal neoplasia. *ANZ J Surg*. 2015.
76. Jung YR, Kim EJ, Choi HJ, et al. Aspirin targets SIRT1 and AMPK to induce senescence of colorectal carcinoma cells. *Mol Pharmacol*. 2015 Oct;88(4):708-19.
77. Shaikat A, Scouras N, Schunemann HJ. Role of supplemental calcium in the recurrence of colorectal adenomas: a meta-analysis of randomized controlled trials. *Am J Gastroenterol*. 2005 Feb;100(2):390-4.
78. Han C, Shin A, Lee J, et al. Dietary calcium intake and the risk of colorectal cancer: a case control study. *BMC cancer*. 2015 Dec;15:966.
79. Klampfer L. Vitamin D and colon cancer. *World J Gastrointest Oncol*. 2014;6(11):430-7.
80. Pereira F, Larriba MJ, Munoz A. Vitamin D and colon cancer. *Endocr Relat Cancer*. 2012;19(3):R51-71.
81. Chai J, Jamal MM. Esophageal malignancy: a growing concern. *World J Gastroenterol*. 2012;18(45):6521-6.
82. Triadafilopoulos G, Aslan A. Same-day upper and lower inpatient endoscopy: a trend for the future. *Am J Gastroenterol*. 1991;86(8):952-5.



Does your multivitamin measure up? Two-Per-Day beats Centrum® in 10 ways!

Are You Getting The Maximum Potency From Your Daily Vitamin?

Life Extension®'s **Two-Per-Day** formulas are the highest potency multivitamins on the market. Compared to **Centrum**® Silver® Adults 50+, **Two-Per-Day** provides:

**Centrum®
Can't
Compete**

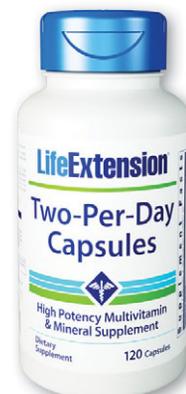
- 50 times more vitamin B1
- 12 times more vitamin B12
- 25 times more vitamin B6
- 10 times more biotin
- 10 times more selenium
- 8 times more vitamin C
- 2 times more vitamin D
- 2 times as much vitamin E
- 2.5 times as much vitamin B3
- 2.7 times as much zinc

Life Extension®'s **Two-Per-Day** contains superior forms of nutrients such as **5-MTHF** that is up to **7 times more bioavailable** than **folic acid**. These more **bioavailable** nutrients provide the body with greater biological **activity**, which is especially important as people age.

Two-Per-Day Capsules

Item #02014 • 120 capsules (2-month supply)

	Retail Price	Your Price
1 bottle	\$22	\$16.50
4 bottles		\$15 each



Two-Per-Day Tablets

Item #02015 • 120 tablets (2-month supply)

	Retail Price	Your Price
1 bottle	\$20	\$15
4 bottles		\$13.50 each



Non-GMO

Contains soybeans.

CAUTION: Individuals consuming more than 2,000 IU/day of vitamin D (from diet and supplements) should periodically obtain a serum 25-hydroxy vitamin D measurement. Vitamin D supplementation is not recommended for individuals with high blood calcium levels.

For the complete list of ingredients, trademarks, cautions, references, dosage and use, please visit www.LifeExtension.com. Two-Per-Day provides a small amount of gamma tocopherols as part of natural mixed tocopherols, which include natural vitamin E. NIAGEN® is a registered trademark of ChromaDex, Inc., Patents see: www.ChromaDexPatents.com.

To order **Life Extension Two-Per-Day Tablets** or **Two-Per-Day Capsules**, call **1-800-544-4440** or visit www.LifeExtension.com

Optimize Thyroid Function with **Triple Action Thyroid**

Feeling Fatigued and Forgetful?

A properly functioning thyroid helps support:

- Energy Levels
- Motivation
- Concentration
- Metabolism and
- Healthy Weight Management

Triple Action Thyroid includes three all-natural herbs, **Ashwagandha**, **Guggul**, and **Korean Ginseng**, to provide comprehensive support by optimizing the thyroid hormones T3 and T4.¹⁻³

The addition of **vitamin B12**, **iodine**, **tyrosine**, and **vitamin A** further enhances its benefits.

Non-GMO

References

1. *Ayurveda Integr Med.* 2014 Oct-Dec;5(4):241-5.
2. *Phytother Res.* 2005 Jan;19(1):78-80.
3. *Zhongguo Zhong Xi Yi Jie He Za Zhi.* 1999 Apr;19(4):209-11.

Due to the source of kelp, this product may contain fish and shellfish.

Sensoril® is protected under US Patent Nos. 6,153,198 and 6,713,092 and is a registered trademark of Natreon, Inc. GS15-4™ is a trademark of ILHWA N.A.

Caution: If you have a thyroid condition or are taking antithyroid medications, do not use without consulting your healthcare practitioner.



Triple Action Thyroid

Item #02003 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$36	\$27
4 bottles		\$24 each

To order **Triple Action Thyroid**,
call **1-800-544-4440**
or visit **www.LifeExtension.com**

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

THE OPTIMAL FORM OF VITAMIN E



Each softgel of Gamma E Tocopherol with Sesame Lignans provides:

Gamma tocopherol	230 mg
Delta tocopherol	90 mg
Alpha tocopherol	35 mg
Sesame lignan extract	20 mg



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**.^{1,2}

SESAME LIGNANS: The Vitamin E Booster!

Sesame and its lignans have been shown to protect against oxidation and help maintain already normal blood pressure.*

In a human study, a combination of **gamma tocopherol** and **sesame lignans** was **25% more effective** in suppressing tissue measurements for free radical and **inflammatory** damage.^{3,4}

Suggested dose is one softgel daily.

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

Gamma E Tocopherol with Sesame Lignans • Item #00759 • 60 softgels

	Retail Price	Your Price
1 bottle	\$32	\$24
4 bottles		\$21.75 each

Non-GMO

Contains soybeans.

References

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

Caution: If you are taking anti-coagulant or anti-platelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product.



Higher Vitamin D Levels Linked with Improved Prostate Cancer Survival

In an article in *Cancer Epidemiology, Biomarkers & Prevention*, researchers at the National Cancer Institute document an association between higher serum **vitamin D** levels and an increased chance of surviving prostate cancer.*

The current investigation included 1,000 participants in the Alpha-Tocopherol, Beta-Carotene Cancer Prevention Study who were diagnosed with prostate cancer following enrollment. Just over 360 subjects died from their disease over 23 years of follow-up from the time of diagnosis. Serum **25-hydroxyvitamin D** levels and other factors were measured upon enrollment and questionnaires concerning diet and medical history were completed by all participants.

Among men whose vitamin D levels were among the top **20%** of subjects, there was a **28% lower** average adjusted risk of dying from prostate cancer compared to those whose levels were among the bottom **20%**. The effect was stronger among those who survived more than 3.3 years.

Editor's Note: "Our findings that higher 25(OH)D reduces the most clinically relevant prostate cancer outcome, disease-specific mortality, if true, could have important public health implications, including whether vitamin D supplementation should be considered for men diagnosed with prostate cancer," the authors conclude.

**Cancer Epidemiol Biomark Prev.* 2016 Jan 25.



Philanthropist Donates \$250 Million to Fund Cancer Research

Sean Parker, 36, co-founder of the file-sharing computer service Napster and former president of Facebook, is donating \$250 million to cancer research via his philanthropic Parker Foundation.*

The newly created Parker Institute for Cancer Immunotherapy will concentrate on this developing field, which harnesses the body's immune system to fight cancer cells.

The institute comprises more than 300 researchers and more than 40 laboratories, including New York's Memorial Sloan Kettering; Stanford Medicine; the University of California, Los Angeles; the University of California, San Francisco; Houston's; University of Texas MD Anderson and the University of Pennsylvania in Philadelphia. The focus will be on three key areas of research: modifying a patient's own immune system T cells to target a tumor; studying ways to boost patient response to current immunotherapy drugs, and research to identify other novel targets to attack a tumor.

Parker hopes the grant will foster collaboration amongst various research centers. "Any breakthrough made at one center is immediately available to another center without any kind of intellectual property entanglements or bureaucracy," said Parker.

Editor's Note: Parker says the death of his friend Laura Ziskin, who produced the films *Pretty Woman* and *Spider-Man*, raised his awareness of the need to overhaul cancer research. Ziskin died of the disease in 2011.

*Available at: <http://www.latimes.com/science/sciencenow/la-sci-sn-cancer-immunotherapy-research-silicon-valley-20160412-story.html>. Accessed April 12, 2016.



© GETTY IMAGES



Decreased Bioavailable and Total Vitamin D Predict Greater Cardiovascular Event Risk

Results of a study presented at the American College of Cardiology Scientific Sessions reveal an association between vitamin D (total and bio-available) levels and lower risk of major adverse cardiovascular events including heart failure, stroke, and heart attack.*

“Many epidemiological studies have shown that [low] total circulating levels of **25-hydroxyvitamin D** [25(OH)D] are strongly associated with poor cardiovascular outcomes,” note Heidi T. May, PhD, MSPH, and colleagues. “However, **85%-90%** of circulating vitamin D is bound to vitamin D binding protein or albumin, and levels obtained may not be truly reflective of the vitamin D available to act on target cells.”

Acting on the results of observational studies, the researchers determined levels of total serum 25(OH)D and 25(OH)D bound to albumin and vitamin D binding protein in 4,200 men and women.

“Our study found that low levels of both total vitamin D and bioavailable vitamin D appear to be associated with poor cardiovascular outcomes,” Dr. May reported.

Editor’s Note: Bioavailable vitamin D includes that which is bound to albumin but not vitamin D binding protein.

*American College of Cardiology Scientific Sessions. 2016 Apr 2.

Meta-Analysis Affirms Association Between Omega-3 and Lowered Inflammation

A meta-analysis published in *PLOS One* found more evidence of a reduction in pro-inflammatory factors in association with supplementation with marine-derived **omega-3 fatty acids**.*

The researchers selected 18 randomized controlled trials involving 826 subjects for their review. Trials included those that evaluated the effects of fish oil (which contains EPA and DHA) or EPA alone on levels of prostaglandin E2, thromboxane B2, or leukotriene B4.

The meta-analysis uncovered a significant association between omega-3 supplementation in reduction in thromboxane B2 in the blood of participants at high risk of cardiovascular disease and in leukotriene B4 in the neutrophils of unhealthy subjects. Reductions in leukotriene B4 occurred only with 14 or more weeks of treatment with omega-3 fatty acids.

“High quality randomized controlled trials are needed to explore the effects of marine-derived omega-3s on different pro-inflammatory factors in subjects with different health status,” concluded the authors.

Editor’s Note: “Several arachidonic acid-derived eicosanoids exert their significant influence on the inflammatory response,” write the authors. Authors stated previous research that “Prostaglandin E2 is involved in the classic signs of inflammation and possesses both pro-inflammatory and anti-inflammatory actions; thromboxane A2, formed by platelets, macrophages and polymorphonuclear leukocytes, can induce vasoconstriction and promotes aggregation of platelets as well as adhesiveness of polymorphonuclear neutrophils; leukotriene B4 (LTB4) can not only increase vascular permeability and enhance local blood flow by stimulating neutrophil secretion, but also stimulate other inflammatory substances.”

**PLOS One*. 2016 Jan.

Whey Protein for Breakfast Could Help Manage Diabetes

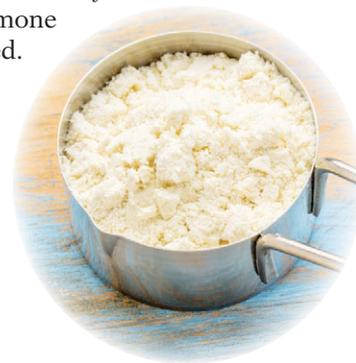
A study reported at the annual meeting of the Endocrine Society found a benefit for the inclusion of whey protein at breakfast by individuals with type II diabetes.*

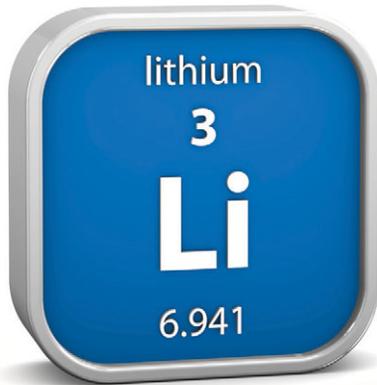
The study included 48 overweight and obese type II diabetics with an average age of 59. Participants were instructed to consume a large breakfast, medium-sized lunch and small dinner that provided the same amount of daily calories for 23 months. Subjects were assigned to breakfasts that consisted of whey protein shakes, other proteins such as eggs and tuna, or carbohydrates. Hemoglobin A1c and other factors were measured at the beginning and end of the study.

The group who received the whey protein shake for breakfast had less hunger, lower glucose spikes following meals and a greater reduction in hemoglobin A1c compared with the other groups. “The whey protein diet significantly suppresses the hunger hormone ghrelin,” lead author Janiela Jakubowicz, MD, noted.

Editor’s Note: “A high-calorie protein breakfast, medium-sized lunch, and small dinner is a proven successful strategy for weight loss, improved satiety, and reduced glucose spikes throughout the day in people with obesity and type II diabetes,” Dr. Jakubowicz stated. “However, the benefits of high protein content at breakfast also depend on the protein source and quality. Whey protein powder, which is a byproduct of milk during cheese production, induced greater satiety and reduction of glucose spikes after meals compared to other protein sources such as eggs, soy, or tuna.”

*Annual Meeting of the Endocrine Society. 2016 Apr 1.





Lithium Extends Life in Flies

A study published in *Cell Reports* found longer life for flies that were given **lithium**, a chemical element and drug used to stabilize mood in bipolar disorder.*

Researchers gave male and female flies high or low doses of lithium chloride or sodium chloride during adulthood or later in life. Flies that received low-dose lithium lived **16%** longer than average and had a maximum lifespan that was **18%** longer than controls, which the researchers attribute to blockage of **glycogen synthase kinase-3 (GSK-3)** and activation of a protein known as NRF2 that aids in the defense of cells against damage.

“We studied the responses of thousands of flies in different conditions to monitor the effects of lithium and how it extends life,” coauthor Dr. Ivana Bjedov stated. “We found low doses not only prolong life but also shield the body from stress and block fat production for flies on a high-sugar diet.”

Editor’s Note: “Identifying a drug target for aging is a crucial step in achieving this and by targeting GSK-3, we could discover new ways of controlling the aging process in mammals, including humans,” predicted first author Jorge Iván Castillo-Quan of University College London.

**Cell Rep.* 2016 Apr 7.

Testosterone Therapy Improves Sexual Function in Diabetic Men

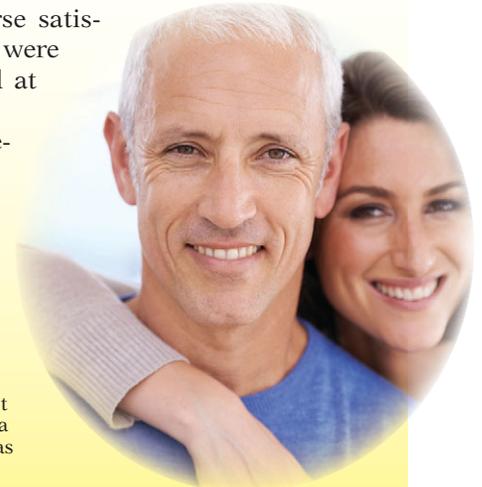
An article appearing in *BJU International* reveals better sexual function in men with type II diabetes and severely depressed testosterone levels who were treated with testosterone undecanoate.*

The study included 189 men, among whom 107 had mild hypogonadism and 87 had severe hypogonadism as demonstrated by low total and free serum testosterone levels. Participants received **1,000 mg** testosterone undecanoate or a placebo at the beginning of the study and at 6 and 18 weeks. Questionnaires that assessed erectile function, intercourse satisfaction, orgasmic function and desire were administered prior to treatment and at 6, 18 and 30 weeks.

After 30 weeks, significant improvement in erectile function was noted by the severely deficient men treated with testosterone. Intercourse satisfaction and desire also improved at 6, 18, and 30 weeks in the severely deficient testosterone-treated group.

Editor’s Note: Lead author Geoffrey Hackett of the Heart of England Foundation NHS Trust commented that “The study’s results also suggest that trials of testosterone therapy should be for a minimum of 6 months and not shorter periods as suggested by some guidelines.”

* *BJU Int.* 2016 Apr 28.



Meta-Analysis Shows Ginseng’s Benefits in Treating Diabetes

Researchers at China’s Zhejiang University provide evidence of a beneficial effect for ginseng in men and women with impaired glucose tolerance or type II diabetes.*

Yun-mei Yang and colleagues selected eight trials that included 195 participants treated with ginseng and an equal number of control subjects. The analysis revealed improvements in fasting glucose, post-meal insulin levels, and insulin resistance, as well as a reduction in triglycerides and total and low-density lipoprotein (LDL) cholesterol among ginseng-treated subjects.



Editor’s Note: As possible mechanisms for ginseng, Dr. Yang and colleagues suggest modulations of insulin production and secretion, glucose metabolism, glucose uptake and inflammation, and activation of the activated protein kinase (AMPK) pathway. Their research suggests that ginseng may be more beneficial for patients who have not started antidiabetic drug therapies than those already being treated.

**Medicine.* 2016 Feb.



Supplements May Prevent Hereditary Deafness

Findings from a study appearing in *Scientific Reports* suggest a role for a combination of free radical scavengers in the prevention of a common form of hereditary deafness caused by mutations in the connexin 26 gene.*

“Many babies born with a genetic mutation that causes deafness pass their newborn screening test but then lose their hearing later in life,” explained co-author Glenn Green, MD, CS. “These patterns suggest that for some children, there may be an opportunity to potentially save cells present at birth.”

In mice with a deletion of connexin 26, prenatal or postnatal supplementation with beta-carotene, vitamins C and E, and magnesium slowed hearing loss progression and improved hearing thresholds. “Our findings suggest that a particular high dose of mineral and vitamin supplements may be beneficial to one genetic mutation,” observed senior author Yehoash Raphael, PhD, of the University of Michigan.

Editor’s Note: Co-author Josef Miller, PhD, who developed the formula, noted that lowering overstimulation-related oxidative stress has been shown to protect the ear’s sensory hair cells. Additionally, free radical scavengers have been shown to preserve cell connections impacted by connexin 26 loss.

**Sci Rep.* 2016 Mar 11.

Decreased Vitamin C Levels Found in Patients Treated for Hematologic Malignancies

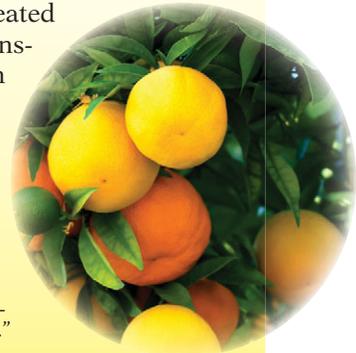
An article appearing in *Results in Immunology* documents the finding of decreased serum levels of **vitamin C** in patients with hematologic malignancies that include leukemia, multiple myeloma, and non-Hodgkin lymphoma.*

“As ascorbic acid has a major influence on (re)generation of immune cells *in vitro*, we executed an observational study in which ascorbic acid serum values of patients with hematological malignancies treated with and without hematopoietic stem cell transplantation were compared with those of healthy volunteers to see if low ascorbic acid levels should be considered of importance regarding immune recovery of these patients,” explained authors Mirelle Huijskens and colleagues.

The researchers measured vitamin C levels in 79 healthy volunteers and 42 hematology-oncology patients treated with chemotherapy or hematopoietic stem cell transplantation. While healthy subjects had median serum ascorbic acid levels of **1.14 mg/dL**, those with hematologic malignancies had levels of **0.36 mg/dL** per liter.

Editor’s Note: The authors add: “Since ascorbic acid might be crucial for immune function and for *in vitro* development and expansion of T and natural killer cells from stem cells, we are currently studying the function and recovery of immune cells while patients are on treatment for various malignancies, and determine the correlation of ascorbic acid serum and leukocyte levels and the possible effect of vitamin C supplementation.”

**Results Immunol.* 2016 Jan 12;6:8-10.



FDA Fast Tracks Potential Breakthrough Glioblastoma Treatment

A new treatment for a particularly aggressive form of brain cancer has been granted breakthrough status and is being fast-tracked by the Food and Drug Administration.*

As reported on CBS’s *60 Minutes*, the first test subject was 20-year-old Stephanie Lipscomb, who began experiencing headaches in 2011. She was diagnosed with a large glioblastoma tumor and told she had months to live. After having **98%** of the tumor removed, her cancer returned. Lipscomb then volunteered for an experimental treatment at Duke University. Today, she is free of a type of cancer that had previously been regarded as an almost-certain death sentence.

The treatment, devised by molecular biologist Matthias Gromeier, utilizes a genetically reengineered polio virus which attacks cancer cells but is harmless to the patient.

Gromeier explains that human cancers “develop a shield of protective measures that make them invisible to the immune system...by infecting the tumor, we are actually removing this protective shield, and enabling the immune system to come in and attack.”

Editor’s note: If you or someone you know has recurrent glioblastoma—that is, you have failed at least one conventional treatment, and you only have one area of tumor and are fully mentally coherent and able to walk and function normally, you may qualify to participate in the Duke University study. Go to www.lifeextension.com/glioblastoma for details.

*www.cbsnews.com/news/polio-cancer-treatment-duke-university-60-minutes-scott-pelley. Accessed May 16, 2016.



MAINTAIN YOUTHFUL MITOCHONDRIAL ACTIVITY

with
CoQ10

Super Ubiquinol CoQ10 combines the energy-activating power of *shilajit* into a formula that's more potent than a stand-alone CoQ10.

- Decline in mitochondrial activity is linked to accelerated brain aging.
- CoQ10 energizes aging cells and enhances mitochondrial function.
- Shilajit works with CoQ10 to increase cellular energy.

Non-GMO
Kaneka QH Ubiquinol® is a registered trademark of Kaneka Corporation.
PrimaVie® is a registered trademark of Natreon, Inc.

Super Ubiquinol CoQ10

Item #01426 • 60 softgels

	Retail Price	Your Price
1 bottle	\$62	\$46.50
4 bottles		\$39 each



To order **Super Ubiquinol CoQ10**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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



Vitamin D3 • 1,000 IU

Item #01751 • 250 softgels

Ideal for smaller individuals who also obtain **2,000-3,000 IUs** in a multi-formula

	Retail Price	Your Price
1 bottle	\$12.50	\$9.38
4 bottles		\$8.44 each



Vitamin D3 • 5,000 IU

Item #01713 • 60 softgels

Ideal for most people who take a multi-formula that contains **2,000-3,000 IUs** of vitamin D

	Retail Price	Your Price
1 bottle	\$10	\$7.50
4 bottles		\$6.50 each

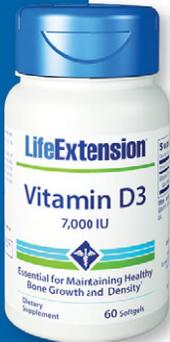


Vitamin D3 • 5,000 IU With Sea Iodine™*

Item #01758 • 60 capsules

With **1,000 mcg** iodine, this is ideal for those who may be iodine-deficient due to a salt-restricted diet

	Retail Price	Your Price
1 bottle	\$14	\$10.50
4 bottles		\$9.38 each



Vitamin D3 • 7,000 IU

Item #01718 • 60 softgels

For individuals who need higher levels, including those who weigh over 180 pounds

	Retail Price	Your Price
1 bottle	\$14	\$10.50
4 bottles		\$9.45 each



Vitamin D3 • 2,000 IU (Natural Mint Flavor)

Item #01732 • 1 ounce

Great for traveling or for those who have trouble swallowing a softgel or capsule
(Also available without mint. Item #00864)

	Retail Price	Your Price
1 bottle	\$28	\$21
4 bottles		\$18.75 each

VITAMIN D3

For Heart Health, Strong Bones, and a Vital Immune System



Find the Formula That's Right for You!

New research on **vitamin D** emerges daily. A simple, cost-effective blood test can help you identify your individual vitamin D needs. Life Extension®'s huge selection of vitamin D supplements allows you to customize your dosage.

To order **Vitamin D3** supplements, call **1-800-544-4440** or visit **www.LifeExtension.com**

Caution: Individuals consuming more than 2,000 IU/day of vitamin D (from diet and supplements) should periodically obtain a serum 25-hydroxy vitamin D measurement. Do not exceed 10,000 IU per day unless recommended by your doctor. Vitamin D supplementation is not recommended for individuals with high blood calcium levels.

*If you have a thyroid condition or are taking anti-thyroid medications, do not use without consulting your health care practitioner.

Not Eating Enough Veggies? No Problem!

Get All The Protective Benefits Of Cruciferous Vegetables

In One Easy-To-Take Supplement

Scientists continue to find healthy benefits—including DNA protection—in cruciferous plants.

But it's not always easy to get in five servings of cruciferous vegetables a day—and even if you do, cooking can destroy many of the protective compounds found in **broccoli, Brussels sprouts, cauliflower, and cabbage.**

Life Extension®'s **Triple Action Cruciferous Vegetable Extract** combines vital plant extracts into a comprehensive formula to provide optimal DNA protection. Each capsule also contains:¹⁻⁹

- **Indole-3-carbinol (I3C)** and **di-indolyl-methane (DIM)** to encourage liver detoxification and modulate estrogen metabolism
- Bioactive **cruciferous** compounds like **glucosinolates** that have favorable effects on healthy cell division
- **Apigenin**, a powerful plant flavonoid, that boosts cell protection¹⁻⁹

Those who want the additional benefits of *trans*-resveratrol can order **Triple Action Cruciferous Vegetable Extract with Resveratrol**. Each vegetarian capsule contains **20 mg** of *trans*-resveratrol in addition to the vegetable extract.

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.

Triple Action Cruciferous Vegetable Extract

Item #01468 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$24	\$18
4 bottles		\$16.50 each

Non-GMO

Triple Action Cruciferous Vegetable Extract with Resveratrol

Item #01469 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$32	\$24
4 bottles		\$22.20 each

Non-GMO



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



An Overlooked Cause of DEPRESSION

While great strides have been made in diagnosing and treating depression, millions of Americans still suffer with this difficult-to-treat disease. It is estimated that **15.7 million** American adults experienced depression in the past year.¹

Antidepressant use is on the rise with one in 10 Americans using them.^{2,3} Unfortunately, antidepressants only work as little as **50%** of the time and come with an array of side effects.^{4,5}

One reason that so many people continue to bear this burden is that most physicians are unaware of the well-documented link between **homocysteine** and **depression**.

A simple blood test that measures **homocysteine** levels, along with a metabolically active form of **folate**, better known as **5-MTHF**, may offer hope to millions of depression sufferers.

Homocysteine and Depression

Homocysteine is an amino acid that, when elevated, can damage the inner linings of the body's vessels leading to a range of cardiovascular conditions including stroke. There is also a correlation between high homocysteine levels and depression.^{6,7}

Researchers studied this association in a group of 924 middle-aged men. They found that individuals in the upper tertile for homocysteine levels had more than a **two-fold increased** risk for being **depressed** compared to those in the lowest tertile.⁷

Another study found that people with the highest levels of homocysteine (**>12 µmol/L**) have significantly lower amounts of **SAME**, a nutrient required for the synthesis of mood-enhancing neurotransmitters.⁸ Not surprisingly, they also had lower levels of mood-enhancing neurotransmitters.

The researchers also found that nearly a third of the depressed participants from the study had red blood cell folate levels **below** normal, and half of these participants had homocysteine levels **higher** than levels of the two control groups.⁸

The connection between low folate levels and imbalances in neurotransmitters shouldn't be surprising since folate is required for the production of neurotransmitters in the brain.

Improvement with 5-MTHF

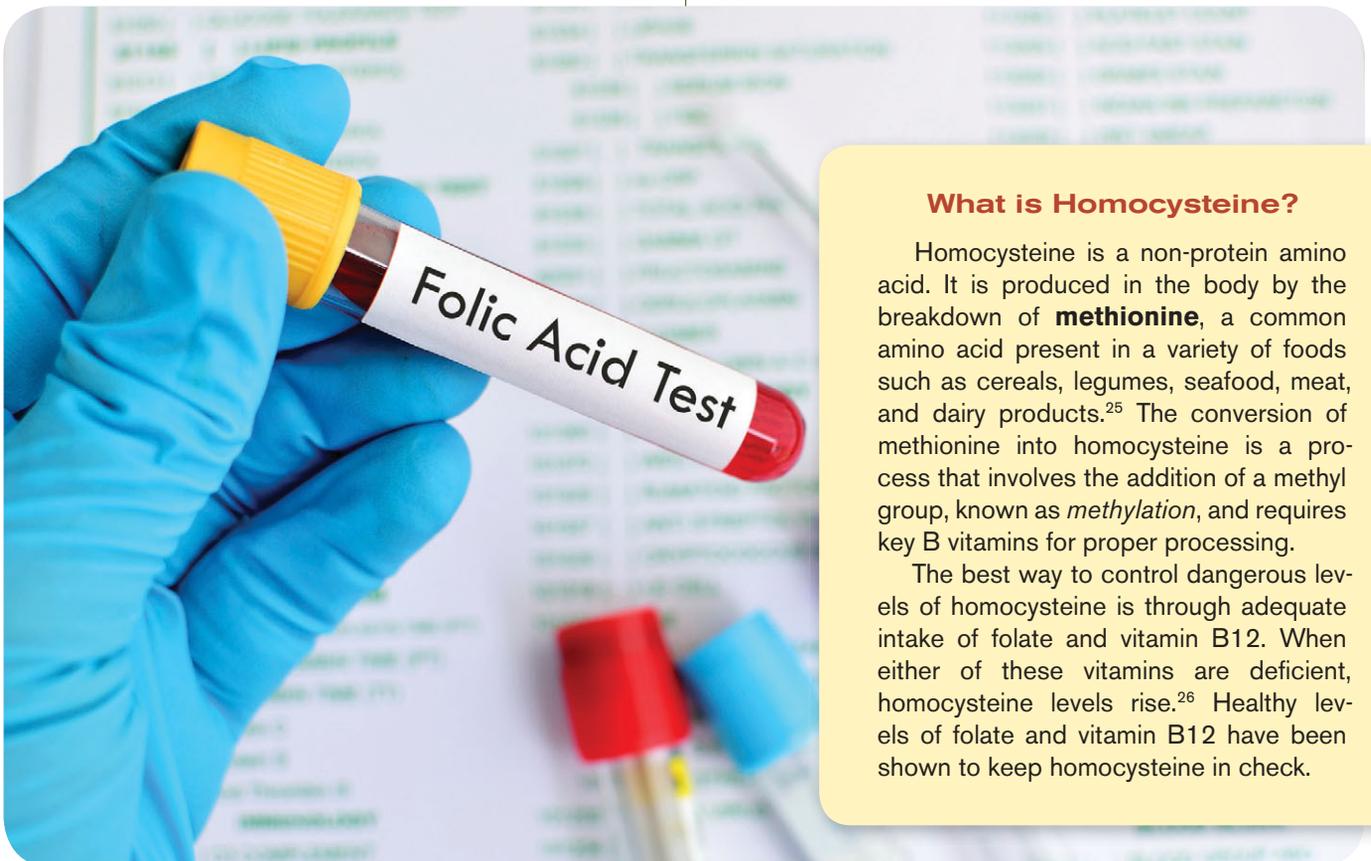
Once the connection between high homocysteine levels and the occurrence of depressive symptoms was discovered, researchers began to investigate whether the active form of folate, **5-MTHF**, could have a beneficial impact on major depression.^{9,10}

The results of these studies open a new door for the treatment of depression. What the scientists discovered was that taking 5-MTHF in addition to antidepressant drugs dramatically improved response rates and did so in a significantly shorter amount of time.

In one study, only **7%** of subjects taking an antidepressant drug experienced major improvement on a standard depression score. That number jumped to **19%** in those patients taking 5-MTHF in addition to an antidepressant drug.

The results were even more impressive in those with the most **severe** depression. Improvements of **40%** were experienced while taking **5-MTHF** in addition to their antidepressant drug, versus just **16%** of those only taking the drug by itself.¹⁰

In addition to improving symptom severity, 5-MTHF brought about these improvements significantly faster than antidepressant drugs alone. Specifically, the 5-MTHF group saw improvements in just **177** days, compared to **231** days in the



What is Homocysteine?

Homocysteine is a non-protein amino acid. It is produced in the body by the breakdown of **methionine**, a common amino acid present in a variety of foods such as cereals, legumes, seafood, meat, and dairy products.²⁵ The conversion of methionine into homocysteine is a process that involves the addition of a methyl group, known as *methylation*, and requires key B vitamins for proper processing.

The best way to control dangerous levels of homocysteine is through adequate intake of folate and vitamin B12. When either of these vitamins are deficient, homocysteine levels rise.²⁶ Healthy levels of folate and vitamin B12 have been shown to keep homocysteine in check.



control group. For those with the most severe depression, the results were even more dramatic. The median time to improvement took only **85** days, compared to **150** days in the control group. A key finding was that nearly twice as many people in the antidepressant-only group stopped therapy due to adverse events (**34%**) versus the 5-MTHF group (**17.9%**), providing evidence of the nutrient's superior safety profile.¹⁰

These results are very encouraging, especially considering the fact that major depression is notoriously difficult to treat, with only about **30%** of patients treated with a single antidepressant achieving resolution of their symptoms, a figure that rises to just **50%** to **55%** when a second drug is added.^{11,12}

This correlation makes supplementing with the metabolically active form of folate, 5-MTHF, a good idea for anyone suffering from depression.

Cognitive Decline, Alzheimer's Disease, and Aging

Depression by itself is bad enough. But now, scientists are finding that depression can lead to Alzheimer's disease and cognitive decline.^{13,14}

Research has confirmed the link between elevated homocysteine, cognitive decline, and Alzheimer's disease.¹⁵⁻¹⁷

Lowering Homocysteine Levels

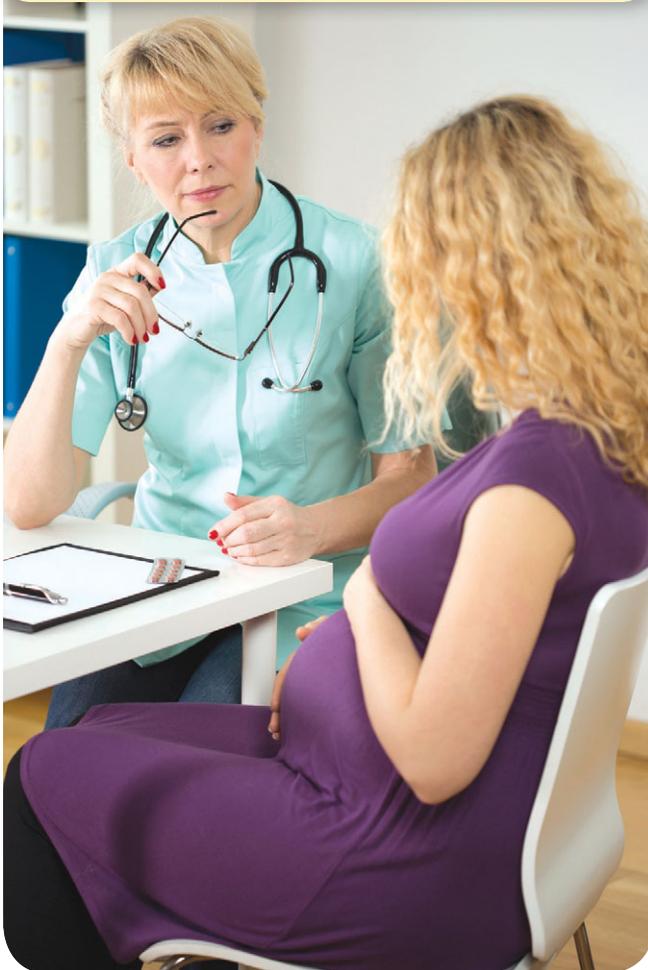
- Elevated homocysteine levels have been linked with a variety of health conditions including stroke, heart disease, macular degeneration, and dementia.
- New findings confirm the connection between elevated levels of homocysteine and depression as well as other psychiatric disorders.
- Factors such as commonly prescribed medications, alcohol, smoking, and advancing age contribute to elevated homocysteine levels.
- B-vitamins like folic acid, vitamin B6, and vitamin B12 can help the body maintain homocysteine levels within a healthy range.
- A significantly high proportion of the American population is at risk for carrying a genetic variant for the MTHFR gene mutation, leading to an impaired ability to metabolize folic acid to its active form.
- *5-methyltetrahydrofolate* or 5-MTHF, the active form of folic acid, is now available as a dietary supplement.
- Together with other complementary nutrients like vitamin B6 and vitamin B12, 5-MTHF provides advanced protection against homocysteine-related disorders.

Postpartum Depression

The well-documented relationship between elevated levels of homocysteine and depression has led scientists to propose using it as a risk biomarker in postpartum depression (PPD).²²

Postpartum depression has been shown to be present in **10%** to **15%** of new mothers.²³ Due to biochemical changes in their brains, mothers with postpartum depression experience fewer positive emotions relating to their pregnancy, and lack desire to actively care for and bond with their newborn children.

A recent study concluded that increased levels of homocysteine are not only related to postpartum depression, but are also associated with impaired production and metabolism of serotonin, an important mood enhancer. Likewise, newborns with mothers affected by postpartum depression scored lower on the APGAR scale (a common assessment of fetal well-being) compared to infants from mothers in a control group.²⁴



A study published in the *New England Journal of Medicine* following elderly participants with dementia found that more than **75%** of participants were eventually diagnosed with Alzheimer's disease over an eight-year follow-up period.¹⁸ The researchers reviewed participants' homocysteine levels and found that for those with homocysteine levels greater than **14 $\mu\text{mol/L}$** , the risk of Alzheimer's nearly doubled. They concluded that increased plasma levels of homocysteine are a strong, independent risk factor for the development of dementia and Alzheimer's disease.

Homocysteine alone is not the sole cause of age-related cognitive decline or memory impairment related to Alzheimer's disease, but there is a significant amount of evidence pointing to its role as a contributing factor.

Get Tested and Safely Lower Homocysteine

Those striving for longevity and seeking to safeguard their health can readily and safely prevent and modulate elevated homocysteine levels. Certain factors such as a diet poor in folate or other important B vitamins may be partially to blame. Sometimes other more complex issues need to be addressed as well to optimally maintain homocysteine levels. It is also important to avoid leading a sedentary lifestyle and consuming excessive amounts of foods rich in the amino acid **methionine**, such as red meats and dairy products. Decreasing or eliminating consumption of alcohol and smoking is also important.

Homocysteine levels should be part of a yearly battery of blood tests to ensure a healthy, long life. **Life Extension®** advises that the optimal range for homocysteine levels is **<7-8 $\mu\text{mol/L}$** , a much more aggressive cut-off than the currently accepted **<15 $\mu\text{mol/L}$** .

Individuals with elevated homocysteine levels should begin supplementation with 5-MTHF and retest levels after 3 months.

Summary

The buildup of homocysteine poses a major threat to one's health, raising the risk for cardiovascular and neurodegenerative diseases. New findings are confirming the connection between levels of homocysteine and psychiatric conditions like depression.

Those interested in guarding themselves from the devastating consequences of elevated homocysteine levels should begin proper supplementation with the bioactive form of folate, 5-MTHF, a convenient and low-cost nutrient that can provide the body with the best ammunition for controlling homocysteine.

Daily doses of **1,000 mcg** to **5,000 mcg (1-5 mg)** of bioactive folate are typically used in research studies to achieve clinically beneficial reductions in plasma homocysteine concentrations.

For optimal homocysteine reduction, adequate amounts of other B-vitamins such as B2, B6, and B12 are also required. These powerful and inexpensive nutritional strategies are readily available to help individuals prevent the wide array of chronic disorders and complications that can be traced to excessive levels of homocysteine. ●

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

5-MTHF Lowers Homocysteine Levels

The superiority of 5-MTHF for increasing blood levels of folate translates directly into lower levels of homocysteine.

In a study on healthy people, a low dose of folic acid (**100 mcg** per day) was compared to a low dose of 5-MTHF (**113 mcg** per day). After six months, the mean total homocysteine was reduced by **9.3%** in the folic acid group, compared to a **14.6%** mean reduction in the 5-MTHF group.¹⁹

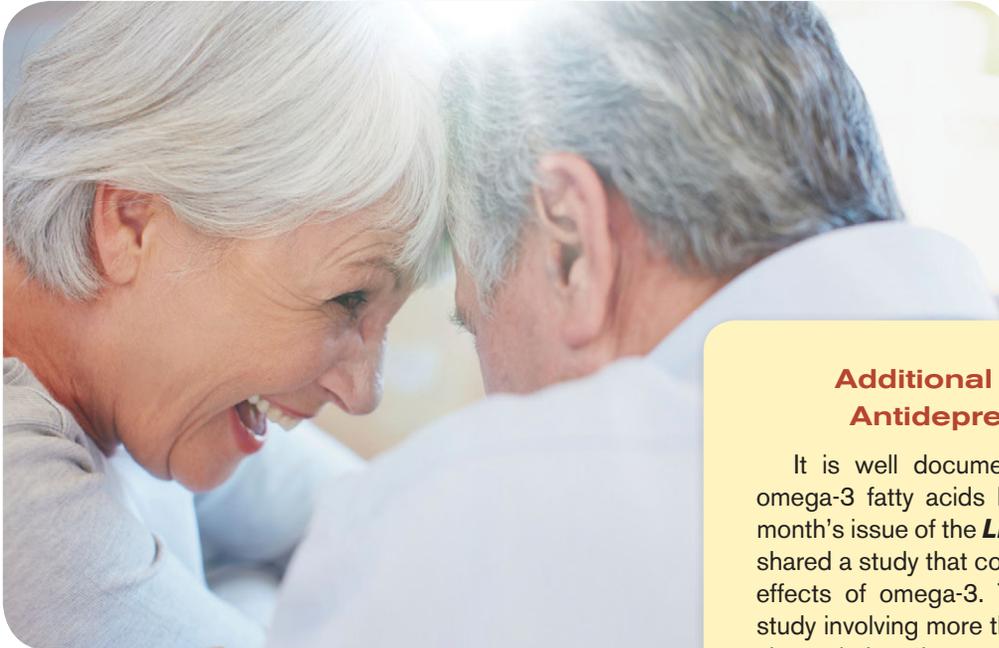
Numerous human studies have confirmed just how much more effective 5-MTHF is than folic acid. One particular crossover study examined cardiovascular patients with the gene variant for the weakened MTHFR enzyme. The researchers found that the subjects' peak serum folate levels were nearly **7 times** higher when taking the active 5-MTHF as they were when taking folic acid.²⁰

Another crossover study compared folic acid and the active 5-MTHF in women with both the typical gene variant for MTHFR and the weaker gene variant. Regardless of the women's genetic makeup, the total amount of folate in the blood, as well as the maximum concentration reached, were significantly higher when taking 5-MTHF.²¹

References:

1. Available at: <http://www.nlm.nih.gov/health/statistics/prevalence/major-depression-among-adults.shtml>. Accessed November 23, 2015.
2. Available at: http://well.blogs.nytimes.com/2013/08/12/a-glut-of-antidepressants/?_r=1. Accessed November 11, 2015.
3. Lindsley CW. The top prescription drugs of 2011 in the United States: antipsychotics and antidepressants once again lead CNS therapeutics. *ACS Chem Neurosci*. 2012;3(8):630-1.
4. Berney P. Dose-response relationship of recent antidepressants in the short-term treatment of depression. *Dialogues in Clinical Neuroscience*. 2005;7(3):249-62.
5. Available at: <http://www.webmd.com/depression/features/coping-with-side-effects-of-depression-treatment>. Accessed May 5, 2016.
6. Folstein M, Liu T, Peter I, et al. The homocysteine hypothesis of depression. *Am J Psychiatry*. 2007;164(6):861-7.
7. Tolmunen T, Hintikka J, Voutilainen S, et al. Association between depressive symptoms and serum concentrations of homocysteine in men: a population study. *Am J Clin Nutr*. 2004;80(6):1574-8.
8. Bottiglieri T, Laundry M, Crellin R, et al. Homocysteine, folate, methylation, and monoamine metabolism in depression. *J Neurosurg Psychiatry*. 2000;69(2):228-32.
9. Papakostas GI, Cassiello CF, Iovieno N. Folates and S-adenosylmethionine for major depressive disorder. *Can J Psychiatry*. 2012;57(7):406-13.
10. Ginsberg LD, Oubre AY, Daoud YA. L-methylfolate plus SSRI or SNRI from treatment initiation compared to SSRI or SNRI monotherapy in a major depressive episode. *Innov Clin Neurosci*. 2011;8(1):19-28.
11. Farah A. The role of L-methylfolate in depressive disorders. *CNS Spectr*. 2009;14(1 Suppl 2):2-7.
12. Wade RL, Kindermann SL, Hou Q, et al. Comparative assessment of adherence measures and resource use in SSRI/SNRI-treated patients with depression using second-generation antipsychotics or L-methylfolate as adjunctive therapy. *J Manag Care Pharm*. 2014;20(1):76-85.





Additional Nutrients with Antidepressant Effects

It is well documented in the literature that omega-3 fatty acids help fight depression. Last month's issue of the **Life Extension Magazine**[®] shared a study that confirmed the anti-depression effects of omega-3. This large "cross-sectional" study involving more than 2,000 men and women showed that those with higher blood levels of omega-3s had a **43%** lower risk of depression.²⁷ For the study, the researchers used a standard 20-question scale of depression symptoms that had been previously validated to show that scores of 16 and higher are present in people with relevant depressive symptoms.

And omega-3s supplementation in combination with commonly prescribed antidepressants is showing additional promising effects. In one study of patients with known major depressive disorder, the combination of omega-3 with antidepressant drug Celexa[®] proved to be superior to the drug alone in relieving depressive symptoms.²⁸

Of no surprise to **Life Extension** are the findings of a recent paper published by *The American Journal of Psychiatry* reporting on the benefits of improving the effectiveness of antidepressants in those with clinical depression through the use of nutrients like SAME, 5-MTHF (methylfolate), omega-3, and vitamin D.²⁹

13. Diniz BS, Butters MA, Albert SM, et al. Late-life depression and risk of vascular dementia and Alzheimer's disease: systematic review and meta-analysis of community-based cohort studies. *Psychiatry*. 2013;202(5):329-35.
14. Butters MA, Young JB, Lopez O, et al. Pathways linking late-life depression to persistent cognitive impairment and dementia. *Dialogues Clin Neurosci*. 2008;10(3):345-57.
15. 5-methyltetrahydrofolate. Monograph. *Altern Med Rev*. 2006;11(4):330-7.
16. Weir DG, Scott JM. Brain function in the elderly: role of vitamin B12 and folate. *Br Med Bull*. 1999;55(3):669-82.
17. Bottiglieri T, Parnetti L, Arning E, et al. Plasma total homocysteine levels and the C677T mutation in the methylenetetrahydrofolate reductase (MTHFR) gene: a study in an Italian population with dementia. *Mech Ageing Dev*. 2001;122(16):2013-23.
18. Seshadri S, Beiser A, Selhub J, et al. Plasma homocysteine as a risk factor for dementia and Alzheimer's disease. *N Engl J Med*. 2002;346(7):476-83.
19. Venn BJ, Green TJ, Moser R, et al. Comparison of the effect of low-dose supplementation with L-5-methyltetrahydrofolate or folic acid on plasma homocysteine: a randomized placebo-controlled study. *Am J Clin Nutr*. 2003;77(3):658-62.
20. Willems FF, Boers GH, Blom HJ, et al. Pharmacokinetic study on the utilisation of 5-methyltetrahydrofolate and folic acid in patients with coronary artery disease. *Br J Pharmacol*. 2004;141(5):825-30.
21. Prinz-Langenohl R, Bramswig S, Tobolski O, et al. [6S]-5-methyltetrahydrofolate increases plasma folate more effectively than folic acid in women with the homozygous or wild-type 677C->T polymorphism of methylenetetrahydrofolate reductase. *Br J Pharmacol*. 2009;158(8):2014-21.
22. Huang J, Zhang L, He M, et al. Comprehensive evaluation of postpartum depression and correlations between postpartum depression and serum levels of homocysteine in Chinese women. *Zhong Nan Da Xue Xue Bao Yi Xue Ban*. 2015;40(3):311-6.
23. Knudson-Martin C, Silverstein R. Suffering in silence: a qualitative meta-data-analysis of postpartum depression. *J Marital Fam Ther*. 2009;35(2):145-58.
24. Aishwarya S, Rajendiren S, Kattimani S, et al. Homocysteine and serotonin: association with postpartum depression. *Asian J Psychiatry*. 2013;6(6):473-7.
25. Available at: <http://nutritiondata.self.com/foods-000084000000000000000000.html>. Accessed May 13, 2016.
26. Klee GG. Cobalamin and folate evaluation: measurement of methylmalonic acid and homocysteine vs vitamin B(12) and folate. *Clin Chem*. 2000;46(8 Pt 2):1277-83.
27. Horikawa C, Otsuka R, Kato Y, et al. Cross-sectional association between serum concentrations of n-3 long-chain PUFA and depressive symptoms: results in Japanese community dwellers. *Br J Nutr*. 2016;115(4):672-80.
28. Gertsik L, Poland RE, Bresee C, et al. Omega-3 fatty acid augmentation of citalopram treatment for patients with major depressive disorder. *J Clin Psychopharmacol*. 2012;32(1):61-4.
29. Sarris J, Murphy J, Mischoulon D, et al. Adjunctive nutraceuticals for depression: a systematic review and meta-analyses. *Am J Psychiatry*. 0(0):appi.ajp.2016.15091228.

THE Neurologically Active Form of VITAMIN B12

Methylcobalamin is the form of **vitamin B12** active in the **central** and **peripheral nervous system**.

Methylcobalamin has been shown to protect against glutamate-induced "excitotoxic" neuronal damage.

Life Extension® offers **methylcobalamin**, the neurologically active form of vitamin B12, at remarkably low prices. Methylcobalamin lozenges come in a **good-tasting vanilla flavor**.



Methylcobalamin
1 mg dissolve-in-mouth
Item #01536 • 60 vegetarian lozenges

	Retail Price	Your Price
1 bottle	\$9.95	\$7.46
4 bottles		\$6 each

Non-GMO



Methylcobalamin
5 mg dissolve-in-mouth
Item #01537 • 60 vegetarian lozenges

	Retail Price	Your Price
1 bottle	\$32	\$24
4 bottles		\$18.75 each

Non-GMO

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

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.

“Turn On” Your Body’s Longevity Genes!

Optimized Resveratrol supports longevity mechanisms **four** different ways:

- **Resveratrol** facilitates youthful *gene expression* analogous to some longevity functions of **calorie restriction**.¹
- **NIAGEN® nicotinamide riboside** helps replenish cellular NAD+ to enhance energy efficiency of mitochondria to support physical performance.²
- **Quercetin*** helps purge the body of inflammation-inducing senescent cells and augment the beneficial gene expression effects of resveratrol.³
- **Pterostilbene, fisetin,** and other **fruit** compounds work synergistically with resveratrol to facilitate youthful gene expression and mimic calorie restriction.^{2,4}

* **Optimized Resveratrol with Nicotinamide Riboside** provides **2.5 times** the previous amount of **quercetin**.



Just one vegetarian capsule of **Optimized Resveratrol** provides:

Trans-Resveratrol (most biologically active form)	250 mg
Quercetin	150 mg
NIAGEN® Nicotinamide Riboside	100 mg
Red grape (fruit) and wild blueberry (fruit) blend	40 mg
Fisetin	10 mg
Trans-Pterostilbene (from pTeroPure®)	0.5 mg

Optimized Resveratrol with Nicotinamide Riboside

Item #02031 • 30 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$42	\$31.50
4 bottles		\$27 each

Non-GMO

To order **Optimized Resveratrol with Nicotinamide Riboside**, call **1-800-544-4440** or visit **www.LifeExtension.com**

References

1. *Cell Metab.* 2011 Nov 2;14(5):612-22.
2. *Aging Cell.* 2011 Oct;10(5):908-11.

3. *Aging Cell.* 2015 Aug;14(4):644-58.
4. *Neurobiol Aging.* 2012 Sep;33(9):2062-71.

CAUTION: If you are taking anti-coagulant or anti-platelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product.

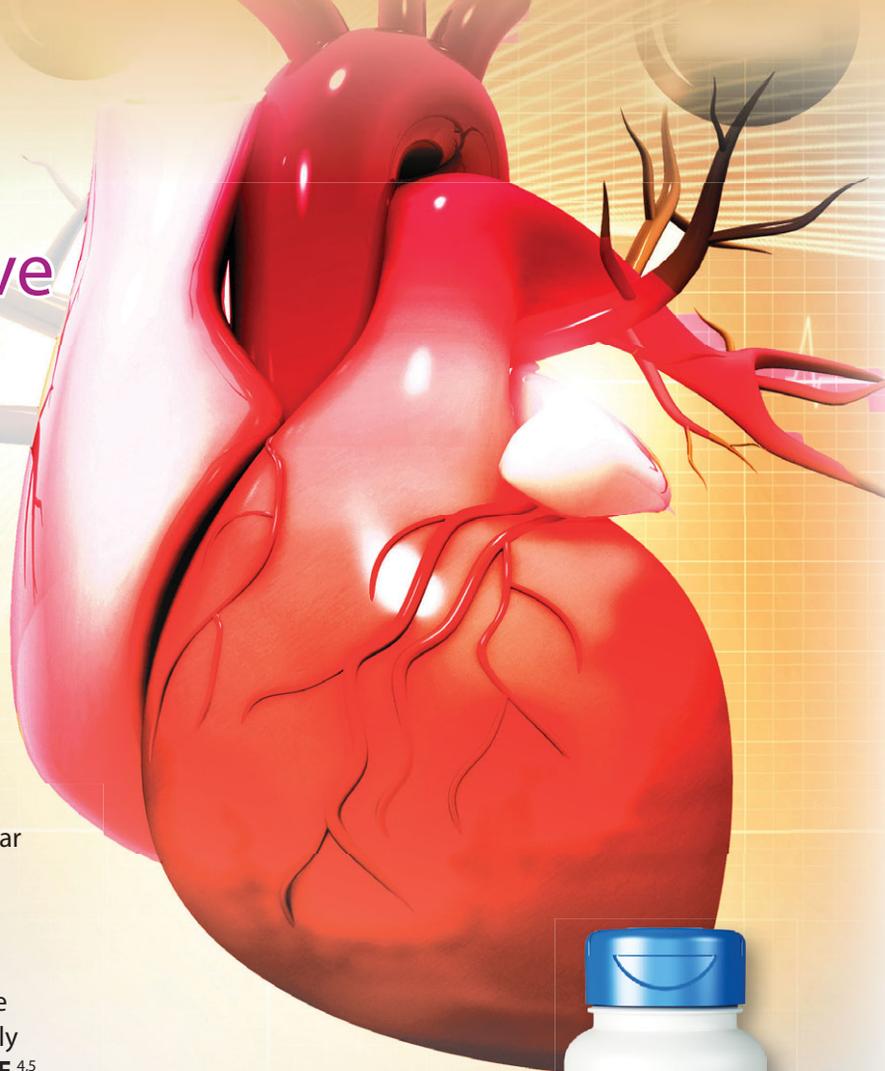
pTeroPure® and NIAGEN® are registered trademarks of ChromaDex, Inc. Patents see: www.ChromaDexPatents.com.

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

Metabolically Active

FOLATE

Supports Cardiovascular Health



Folate helps maintain homocysteine levels within the normal range,¹ thereby promoting cardiovascular health.² Folate also supports neurotransmitter synthesis—which in turn helps maintain cognitive abilities.³

However, not everyone has sufficient activity of the **enzyme** required to convert folate to its biologically active form, **5-methyltetrahydrofolate**, or **5-MTHF**.^{4,5}

For those whose homocysteine levels remain stubbornly high, the answer lies with the *bioactive* form of folate called **5-MTHF**, which is up to **7 times** more bioavailable than ordinary folic acid.⁶ This unique compound requires no enzymatic conversion to become metabolically active⁶—providing maximum support for both cardiovascular and cognitive health.

Optimized Folate provides metabolically active **5-MTHF** folate in **1,000 mcg** or **5,000 mcg** strengths.

The demand for **5-MTHF** has surged as more consumers have discovered its potent **homocysteine-lowering** effects.

Non-GMO

Quatrefolic® is a registered trademark of Gnosis, S.p.A.
Patent number 7,947,662.

References

1. *Am J Clin Nutr.* 2006 Apr;83(4):842-50.
2. *Am J Ther.* 2014; Epub Nov 17.
3. *Innov Clin Neurosci.* 2011 Jan;8(1):19-28.
4. *Am J Hum Genet.* 2002 Mar;70(3):758-62.
5. *Coll Antropol.* 2004 Dec;28(2):647-54.
6. *Br J Pharmacol.* 2004 Mar;141(5):825-30.

Optimized Folate (1,000 mcg)

Item #01939 • 100 vegetarian tablets

	Retail Price	Your Price
1 bottle	\$19	\$14.25
4 bottles		\$12.75 each



High Potency Optimized Folate (5,000 mcg)

Item #01913 • 30 vegetarian tablets

	Retail Price	Your Price
1 bottle	\$25	\$18.75
4 bottles		\$16.50 each



Caution: Do not use this product if treated with methotrexate, fluorouracil, phenytoin, phenobarbital, primidone, or levodopa prior to consultation with your personal physician.

To order either of these **Optimized Folate** formulas, call **1-800-544-4440** or visit **www.LifeExtension.com**





Healthy Choices *Protect Against* Colorectal Cancer

The **American Cancer Society** estimates that the majority of colorectal cancers, and subsequent deaths, could be prevented if we applied *existing knowledge* about this prevalent malignancy.¹

Simply making healthy diet and lifestyle choices can reduce the risk for developing **colorectal cancer**.

With a wealth of research available, too many people continue to needlessly die from this preventable disease.

This article describes strategies available for reducing one's risk of contracting colorectal cancer, which translates into living a longer, healthier life.

The Most Preventable Cancer

Colorectal cancer (cancer that occurs in the colon or rectum) represents the second leading cause of cancer death in men and women combined in the United States.^{2,3} According to the American Cancer Society, an approximate **135,000** new cases will be discovered and close to **50,000** deaths will occur in this year alone.⁴

Despite these statistics, colorectal cancer is one of the most preventable of all cancers. Lifestyle factors play a predominant role in the development of the disease, which means we have a tremendous amount of control over this second-leading cause of cancer deaths.

Cancers in general do not thrive when the body is operating at peak health. This includes active biological systems that work to stop cancer before it even starts by preventing and repairing DNA damage, identifying and destroying abnormal cells, and regulating normal cell growth. The best way to keep these systems in peak condition is through healthy diet and lifestyle choices, as well as incorporating a supplement regimen that includes powerful chemopreventive agents.

The authors of a study published in *Clinical Colorectal Cancer* said, “Improving the awareness of the population with regard to the benefits of a healthy lifestyle, including a balanced diet associated with exercise, could globally reduce CRC (colorectal cancer) risk.”⁵

A “healthy lifestyle” can include something as simple as regular brisk walking, which has been shown to reduce colorectal cancer risk by **18%-24%**.^{6,7}

The reason why exercise is always near the top of any health-optimization regimen is because increased physical activity significantly improves insulin resistance, and also improves levels of inflammatory mediators and inappropriate growth factors. All of these factors, if left untreated, tend to promote cancer development.^{5,8}



Studies show that these foods are associated with **increased** colorectal cancer risk.^{6,9,11}



Importance of Diet on Colorectal Cancer Incidence

Even more vital than exercise, *diet* is known to be one of the most important aspects of reducing colon cancer risk.⁹⁻¹¹ While dietary factors can impact **all** cancer types, they are especially critical in colorectal cancer because of the function and location of the large intestine (which includes the colon and rectum).

All food and drink that enters the body and has to be eliminated must pass through the large intestine. This exposes cells that line the large intestine to carcinogens that either directly damage DNA, or that increase inflammation, which causes free radicals that also damage DNA.

Eating foods cooked at very high temperatures, as *Life Extension Magazine*[®] has warned readers about for decades, can cause the formation of cancer-causing chemicals that damage cells of the large intestine.

Studies show that there are numerous foods (or components of foods) that are associated with **increased** colorectal cancer risk.^{6,9,11} These include red and processed meats, preserved foods, saturated fats, high-sugar foods, and refined carbohydrates (“white” starches).

Numerous dietary factors have been shown to have a *protective* effect against colorectal cancer. These include regular consumption of calcium, vitamin D, fruits, vegetables (especially cruciferous vegetables such as broccoli, cauliflower, and Brussels sprouts), fiber, and fish.^{6,9}

More specifically, regular fish consumption reduces the risk of colorectal cancer by **12%**, and eating fish while ingesting more than **20 grams** per day of fiber can reduce risk by **25%**.⁶

Cancer-Fighting Nutrients

A glance at the protective dietary factors above reveals that plants and fish products play an especially large role in colorectal cancer prevention. This is due in large part to the abundance of protective molecules that these organisms produce, including vitamins and a host of other biologically active compounds.

While healthy diet is critical, nutrient supplements also provide many compounds shown to help guard against cancer. The use of supplements to reduce the risk of cancer (known as **chemoprevention**) shows promise for preventing, slowing, and even reversing colon cancer development.

Here is a list of some of the best-studied nutrients in this area:

Higher **vitamin D** levels are associated with reduced colorectal cancer incidence and deaths. This is likely because vitamin D has immune-supportive functions that can boost the activity of cells that seek and destroy early cancer cells. Vitamin D also reduces the chronic inflammation that can promote colorectal cancer development.¹²

Studies show that **vitamin E**, especially in the form of *gamma-tocotrienol*, reduces colorectal cancer cell growth and even induces tumor cell death.^{13,14} In addition, vitamin E has strong free radical scavenging properties that may prevent DNA mutations from occurring to begin with.¹⁵

Folic acid is important in protecting DNA strands from damage, and has been shown to reduce colorectal cancer risk by **42%** in people with inflammatory bowel disease, a group at high risk for this type of cancer.¹⁶

Minerals, particularly calcium and selenium, also appear to factor heavily in colon cancer prevention.¹² In a large Korean study, people in the highest **25%** of **calcium** intake had an impressive **84%** reduction in colon cancer risk, compared with those in the lowest **25%** of intake.¹⁷ Lab studies reveal that **selenium** compounds can induce *autophagy*, a form of cancer cell death in which cells simply destroy themselves. This powerful mechanism is currently being targeted by drug developers.¹⁸

Fish oil, rich in omega-3, is a potent inhibitor of inflammation throughout the body, making it a natural fit for colon cancer prevention.^{19,20} Human studies demonstrate that supplementation with the omega-3 EPA (eicosapentaenoic acid, **2 grams** daily for 30 days prior to surgery for colorectal cancer) can reduce the formation of new blood vessels essential for tumor growth, and appeared to produce some increases in overall survival for the first 18 months post-operatively.²¹



What You Need to Know

Powerful Chemoprevention for Colorectal Cancer

- Colorectal cancers are the third most common cancer and the second deadliest cancer among US adults.
- Aggressive screening with colonoscopy has reduced rates of colorectal cancers and the resulting death rates, but even aggressive screening misses thousands of cases annually.
- Prevention is always better than treatment, and for colon cancer multiple nutrients are available that can slow or prevent colorectal cancer development.
- Several drugs, especially metformin, also show strong cancer chemopreventive effects.
- Given the clear and present risk of colorectal cancers in all of our lives, it makes sense to construct and follow a cancer-prevention strategy that includes lifestyle and diet changes as well as a thoughtful supplement regimen.

N-acetyl cysteine (NAC) is a strong natural free radical scavenger compound, making it appealing for use in cancer prevention of the colon.²² Studies have shown that people at risk for colorectal cancer who took NAC supplements, **600 mg** per day, had a **40%** reduction in the recurrence of colonic polyps.²² Biochemical and microscopic studies reveal that NAC augments the function of mitochondria (powerhouse of our cells), and produces microscopically detectable changes in early colon cancer in animals.²³

Cimetidine Prevents Colon Cancer Metastasis and Increases Survival

Cimetidine, commonly known as Tagamet®, is a well-known over-the-counter medication used historically to alleviate heartburn. A growing body of evidence has shown that it also has potent anti-cancer ability. It functions via several mechanisms to *inhibit* metastasis and *improve* survival in colon cancer patients.⁶⁸⁻⁷⁵

A study published in the *British Journal of Cancer* showed cimetidine's potent anti-cancer effects. For the study, 64 colon cancer patients received chemotherapy with or without cimetidine (**800 mg** per day) for one year. For the cimetidine group, the 10-year survival was close to **90%** compared to only **49.8%** for the control group. Remarkably, for those subjects with a more aggressive form of colon cancer, the 10-year survival was **85%** in those treated with cimetidine compared to **23%** in the control group.⁷⁴ These findings were confirmed in another study in which colorectal cancer patients were given cimetidine for just seven days at the time of their surgery, and 3-year survival increased from **59%** to **93%**.⁷⁵

Cimetidine does not reduce colon cancer incidence, so it should not be taken for prevention purposes. It has demonstrated powerful **treatment** benefits in those who contract colon cancer. Colon cancer patients should consider taking **800 mg** per day of cimetidine five days prior to surgical removal of their tumor and for one year after surgery to reduce metastatic risk. Some people take a 2-3 month course of cimetidine once a year to boost natural killer cell activity.

As beneficial as they are when taken separately, taking certain nutrients in combination has been shown to have additional value in colorectal cancer prevention. In a large clinical study, the combination of vitamins A, C, and E, along with the element selenium (**200 micrograms**), reduced the recurrence rate of precancerous polyps by **39%**.²⁴ Daily multivitamin use was associated with an **8%** reduction in colorectal cancer risk compared with nonusers.²⁵

The Protective Power of Plant-Based Nutrients

Phytonutrients are components of plants thought to promote human health. They help protect plants from dangers such as UV radiation and insect attacks. They also confer protection to those who consume the plants.

Not surprisingly, plant-based nutrients have multiple protective effects against colon cancer. We've included a summary of some of the best-studied phytonutrients.

Garlic, especially aged garlic, suppresses the excessive cell proliferation that occurs in the earliest stages of colon cancer. One animal study showed that supplementation with aged garlic extract significantly reduced the development of *aberrant crypt foci* (precursors of cancer).²⁶ It also reduced the total number of polyps and cancers that formed, a finding that was replicated in humans with a dose of **2.4 mL** (approximately half a teaspoon of liquid aged garlic extract) per day for one year.²⁷ Human studies show a **37%** reduction in colorectal cancer risk in people with the highest garlic consumption.²⁸

Ginger has effects similar to those of garlic, especially by blocking cell replication and boosting cancer cell self-destruction.²⁹⁻³² Doses of ginger at **2 grams** per day have produced significant favorable changes in the intestinal lining cells of people at increased risk for colorectal cancer.²⁹

Milk thistle extracts, particularly *silibinin* and *silymarin*, have numerous mechanisms that help prevent cancers from forming. These include triggering cancer cell suicide (apoptosis), reducing inflammatory changes, and blocking cell replication. Importantly, they also help prevent the spread of existing cancers by reducing the production of *protein-melting* enzymes that cancer cells use to invade and metastasize.³³⁻³⁵

Cruciferous vegetable extracts are those derived from broccoli, cabbage, Brussels sprouts, and related plants. These extracts are rich in molecules called *isothiocyanates*, which have been shown to promote cancer cell suicide and inhibit colorectal cancer development in animal and lab models.^{36,37} **Indole-3-carbinol (I3C)** is another component of these plants that has



been shown to reduce inflammation and attenuate colorectal tumor development in animal models. Human studies show that those with higher levels of indole-3-carbinol have significantly lower colorectal cancer risk.³⁸⁻⁴¹

Modified citrus pectin is a gel-like compound extracted from citrus peels that has direct tumor-killing effects on cancer cells. More importantly, animal studies show that it significantly prevents the spread of colorectal cancers to other parts of the body.^{42,43} This effect arises from pectin's ability to bind to and block adhesion molecules that metastatic tumor cells use to stick to, and invade, distant tissues.^{42,43} (Modified citrus pectin is usually taken by those already stricken with cancer. It is not considered a preventive.)

Coffee is rich in numerous bioactive molecules that appear to have anti-cancer properties, such as inducing cancer cell suicide.^{44,45} Studies show that people with the highest coffee consumption (more than 3 cups per day) have a **33%-79%** reduction in their risk for colorectal cancers compared to the lowest consumers.^{46,47}

Polyphenols Reduce Cancer Risk

Among the largest group of plant-based nutrients with demonstrated cancer-fighting potential is the **polyphenol** family.⁴⁸ Numerous epidemiological studies offer solid evidence that diets rich in polyphenols result in significantly lower incidence for many kinds of cancer—including colon cancer.^{49,50}

Understanding Colorectal Cancer

Cancers of the colon and rectum are collectively referred to as **colorectal cancer**. Both structures have similar cellular linings, which is where cancers arise. Although colorectal cancer has numerous causes, these all boil down to chronic chemical stress, inflammation, and loss of control over cell division common to all cancers.

Like all other types of cancer, colorectal cancer begins when DNA damage leads to mutations that deregulate DNA replication.^{76,77} This causes cells to begin to multiply out of control, which can lead to cancer. Colorectal cancer typically begins as a precancerous growth called a **polyp** (also known as an **adenoma**).⁷⁸⁻⁸⁰ Over time, inflammation and oxidative stress can cause benign polyps to progress to a specific kind of cancer called **adenocarcinomas**.^{76,81} This type of cancer can be especially dangerous because it is known for spreading to other parts of the body.

Colorectal cancer is typically a silent disease. By the time it produces noticeable symptoms (diarrhea, constipation, bloody stools), the disease is often already in its advanced stages. This highlights the importance of having a routine colonoscopy, which is associated with a **77%** mortality risk reduction over 10 years.⁸² But these screenings are not foolproof.

While it's true that colonoscopies have sharply reduced the death rates from colorectal cancers, we are still missing far too many dangerous colorectal cancers, often until it is too late.

That's why **prevention** is so important. Experts agree that lifelong colon health may be promoted safely, effectively, and naturally using **cutting-edge, low-cost** nutritional therapies.⁸³

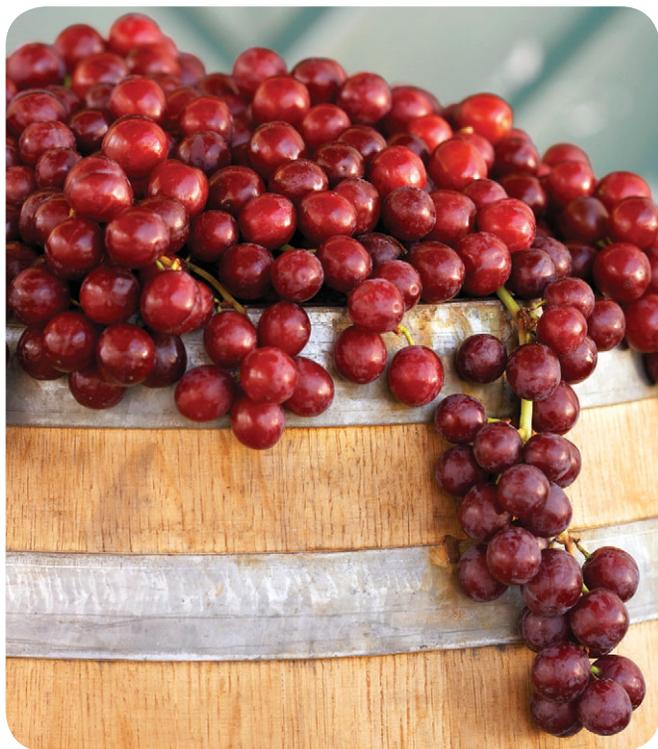
Three of the most potent polyphenols with colon cancer-preventive effects are resveratrol, quercetin, and curcumin.

Resveratrol is found in grape skins and other red-pigmented foods. It is one of the best-known polyphenols—and for good reason. It has actions against multiple stages of the cancer-development process that make it a potent chemopreventive agent. Resveratrol is a gene-modifying agent that enhances cells' resistance to oxidant- and inflammation-promoting stresses.⁵¹ It has multiple, complementary effects on colon cancer development, including switching "on" a tumor suppressor gene that gets switched "off" early in

carcinogenesis and inhibiting genes for invasion-promoting proteins.⁵² Preliminary human studies show that oral doses of resveratrol, ranging from **500 to 5,000 mg** per day, achieve levels in the intestine consistent with anti-carcinogenic effects.^{53,54} For people ingesting a wide variety of polyphenols from their healthy diet and supplements, lower doses (**100 to 250 mg** per day) of resveratrol may be all that is needed.

Quercetin is found in apples and onions. It has been shown to stop the cancer cell replicative cycle and is strongly associated with a reduction in inflammatory molecules that promote cancer development.^{55,56}

Curcumin is what gives the Indian spice turmeric its yellow color. This phenolic compound has an impressive suite of anti-cancer properties that prevent or fight cancer at multiple stages of the development process.⁵⁷ It blocks inflammation that promotes cancer progression, it stops the cancer cell replication cycle in its tracks, it increases the rate of cancer cell self-destruction, and it prevents tumor cells from developing invasive and metastatic potentials.⁵⁸⁻⁶⁰ In addition, lab studies reveal its ability to inhibit the recently-discovered **cancer stem cells**, which hide in tissues and are a major cause of cancer recurrence after treatment.⁶¹ And a human clinical trial showed that curcumin, **4 grams** per day for 30 days, significantly reduced the number of precancerous lesions found on colonoscopy, compared with controls.⁶² (Those obtaining nutrients from healthy diets and supplements usually need only **400 mg** per day of highly absorbable curcumin.)



Drugs that Prevent Colorectal Cancer

Several pharmaceutical agents are showing signs of effectiveness in preventing colorectal cancers. **Aspirin** has been associated with a reduction in colorectal cancer incidence and deaths, even at the low doses used for cardiovascular disease protection (**75 to 100 mg** per day).⁶³ One very large, long-term study has shown that regular aspirin use is associated with as much as a **19%** reduction in colorectal cancer rates.⁶⁴ Unfortunately, most doctors overlook aspirin's cancer-preventive properties.

Metformin is an antidiabetic drug with a longstanding safety record and a suite of actions more similar to natural supplements with multiple targets than to most single-targeted drugs (indeed, it is derived from the French Lilac tree). Because diabetes is a major risk factor for colorectal cancer, there has been great interest in metformin for cancer prevention, both in diabetes patients and in others.⁶⁵ A study demonstrated that metformin use is associated with a **10%** reduction in the incidence of colorectal cancers and a **32%** increase in survival, compared with nonusers.⁶⁶ In another study, metformin at a dose of **250 mg** daily showed a **40%** reduction in the prevalence of precancerous adenomas.⁶⁷

Summary

Colorectal cancer remains the second leading cause of cancer deaths.

Colorectal cancer risk may be greatly reduced through a combination of lifestyle and diet interventions.

A number of nutritional supplements have demonstrated colorectal cancer-preventive properties. These supplements are well-tolerated and widely available, and can form the backbone of a thoughtful, long-term, overall disease prevention strategy. ●

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

REFERENCES

- Available at: <http://www.cancer.org/acs/groups/content/documents/document/acspc-042280.pdf>. Accessed May 2, 2016.
- Available at: <http://www.ccalliance.org/get-information/what-is-colon-cancer/statistics/>. Accessed May 2, 2016.
- Available at: <http://www.cancer.gov/types/colorectal/patient/colorectal-prevention-pdq#section/all>. Accessed May 2, 2016.
- Available at: <http://www.cancer.org/cancer/colonandrectumcancer/detailedguide/colorectal-cancer-key-statistics>. Accessed May 11, 2016.
- Aran V, Victorino AP, Thuler LC, et al. Colorectal cancer: Epidemiology, disease mechanisms and interventions to reduce onset and mortality. *Clin Colorectal Cancer*. 2016.
- Baena R, Salinas P. Diet and colorectal cancer. *Maturitas*. 2015;80(3):258-64.
- Golshiri P, Rasooli S, Emami M, et al. Effects of physical activity on risk of colorectal cancer: a case-control study. *Int J Prev Med*. 2016;7:32.
- Lee DH, Kim JY, Lee MK, et al. Effects of a 12-week home-based exercise program on the level of physical activity, insulin, and cytokines in colorectal cancer survivors: a pilot study. *Support Care Cancer*. 2013;21(9):2537-45.
- Azeem S, Gillani SW, Siddiqui A, et al. Diet and colorectal cancer risk in Asia—a systematic review. *Asian Pac J Cancer Prev*. 2015;16(13):5389-96.
- Millen AE, Subar AF, Graubard BI, et al. Fruit and vegetable intake and prevalence of colorectal adenoma in a cancer screening trial. *Am J Clin Nutr*. 2007;86(6):1754-64.
- Tarraga Lopez PJ, Albero JS, Rodriguez-Montes JA. Primary and secondary prevention of colorectal cancer. *Clin Med Insights Gastroenterol*. 2014;7:33-46.
- Meeker S, Seamons A, Maggio-Price L, et al. Protective links between vitamin D, inflammatory bowel disease and colon cancer. *World J Gastroenterol*. 2016;22(3):933-48.
- Zhang JS, Li DM, Ma Y, et al. gamma-Tocotrienol induces apoptosis-like cell death in human colon carcinoma SW620 cells. *PLoS One*. 2013;8(2):e57779.
- Forster GM, Raina K, Kumar A, et al. Rice varietal differences in bioactive bran components for inhibition of colorectal cancer cell growth. *Food Chem*. 2013;141(2):1545-52.
- Campbell S, Stone W, Whaley S, et al. Development of gamma (gamma)-tocopherol as a colorectal cancer chemopreventive agent. *Crit Rev Oncol Hematol*. 2003;47(3):249-59.
- Burr NE, Hull MA, Subramanian V. Folic acid supplementation may reduce colorectal cancer risk in patients with inflammatory bowel disease : A systematic review and meta-analysis. *J Clin Gastroenterol*. 2016.
- Han C, Shin A, Lee J, et al. Dietary calcium intake and the risk of colorectal cancer: a case control study. *BMC Cancer*. 2015;15:966.
- Yang Y, Luo H, Hui K, et al. Selenite-induced autophagy antagonizes apoptosis in colorectal cancer cells in vitro and in vivo. *Oncol Rep*. 2016;35(3):1255-64.
- Sorensen LS, Thorlacius-Ussing O, Rasmussen HH, et al. Effects of perioperative supplementation with omega-3 fatty acids on leukotriene B(4) and leukotriene B(5) production by stimulated neutrophils in patients with colorectal cancer: a randomized, placebo-controlled intervention trial. *Nutrients*. 2014;6(10):4043-57.
- Ma CJ, Wu JM, Tsai HL, et al. Prospective double-blind randomized study on the efficacy and safety of an n-3 fatty acid enriched intravenous fat emulsion in postsurgical gastric and colorectal cancer patients. *Nutr J*. 2015;14:9.
- Cockbain AJ, Volpato M, Race AD, et al. Anticorectal cancer activity of the omega-3 polyunsaturated fatty acid eicosapentaenoic acid. *Gut*. 2014;63(11):1760-8.
- Ponz de Leon M, Roncucci L. Chemoprevention of colorectal tumors: role of lactulose and of other agents. *Scand J Gastroenterol Suppl*. 1997;222:72-5.
- Amrouche-Mekkioui I, Djerdjouri B. N-acetylcysteine improves redox status, mitochondrial dysfunction, mucin-depleted crypts and epithelial hyperplasia in dextran sulfate sodium-induced oxidative colitis in mice. *Eur J Pharmacol*. 2012;691(1-3):209-17.
- Bonelli L, Puntoni M, Gatteschi B, et al. Antioxidant supplement and long-term reduction of recurrent adenomas of the large bowel. A double-blind randomized trial. *J Gastroenterol*. 2013;48(6):698-705.
- Heine-Broring RC, Winkels RM, Renkema JM, et al. Dietary supplement use and colorectal cancer risk: a systematic review and meta-analyses of prospective cohort studies. *Int J Cancer*. 2015;136(10):2388-401.
- Jikihara H, Qi G, Nozoe K, et al. Aged garlic extract inhibits 1,2-dimethylhydrazine-induced colon tumor development by suppressing cell proliferation. *Oncol Rep*. 2015;33(3):1131-40.
- Tanaka S, Haruma K, Yoshihara M, et al. Aged garlic extract has potential suppressive effect on colorectal adenomas in humans. *J Nutr*. 2006;136(3 Suppl):821s-6s.
- Chiavarini M, Minelli L, Fabiani R. Garlic consumption and colorectal cancer risk in man: a systematic review and meta-analysis. *Public Health Nutr*. 2016;19(2):308-17.
- Citronberg J, Bostick R, Ahearn T, et al. Effects of ginger supplementation on cell-cycle biomarkers in the normal-appearing colonic mucosa of patients at increased risk for colorectal cancer: results from a pilot, randomized, and controlled trial. *Cancer Prev Res (Phila)*. 2013;6(4):271-81.
- Qi LW, Zhang Z, Zhang CF, et al. Anti-colon cancer effects of 6-shogaol through G2/M cell cycle arrest by p53/p21-cdc2/cdc25A crosstalk. *Am J Chin Med*. 2015;43(4):743-56.
- Tahir AA, Sani NF, Murad NA, et al. Combined ginger extract & Gelam honey modulate Ras/ERK and PI3K/AKT pathway genes in colon cancer HT29 cells. *Nutr J*. 2015;14:31.
- Wee LH, Morad NA, Aan GJ, et al. Mechanism of chemoprevention against colon cancer cells using combined Gelam honey and ginger extract via mTOR and wnt/beta-catenin pathways. *Asian Pac J Cancer Prev*. 2015;16(15):6549-56.
- Eo HJ, Park GH, Song HM, et al. Silymarin induces cyclin D1 proteasomal degradation via its phosphorylation of threonine-286 in human colorectal cancer cells. *Int Immunopharmacol*. 2015;24(1):1-6.
- Kauntz H, Bousserouel S, Gosse F, et al. Silibinin, a natural flavonoid, modulates the early expression of chemoprevention biomarkers in a preclinical model of colon carcinogenesis. *Int J Oncol*. 2012;41(3):849-54.
- Kauntz H, Bousserouel S, Gosse F, et al. Silibinin triggers apoptotic signaling pathways and autophagic survival response in human colon adenocarcinoma cells and their derived metastatic cells. *Apoptosis*. 2011;16(10):1042-53.
- Chen MJ, Tang WY, Hsu CW, et al. Apoptosis induction in primary human colorectal cancer cell lines and retarded tumor growth in SCID mice by sulforaphane. *Evid Based Complement Alternat Med*. 2012;2012:415231.
- Slaby O, Sachlova M, Brezkova V, et al. Identification of microRNAs regulated by isothiocyanates and association of polymorphisms inside their target sites with risk of sporadic colorectal cancer. *Nutr Cancer*. 2013;65(2):247-54.
- Kim YH, Kwon HS, Kim DH, et al. 3,3'-diindolylmethane attenuates colonic inflammation and tumorigenesis in mice. *Inflamm Bowel Dis*. 2009;15(8):1164-73.
- Lerner A, Grafi-Cohen M, Napsu T, et al. The indolic diet-derivative, 3,3'-diindolylmethane, induced apoptosis in human colon cancer cells through upregulation of NDRG1. *J Biomed Biotechnol*. 2012;2012:256178.
- Tse G, Eslick GD. Cruciferous vegetables and risk of colorectal neoplasms: a systematic review and meta-analysis. *Nutr Cancer*. 2014;66(1):128-39.

41. Yang G, Gao YT, Shu XO, et al. Isothiocyanate exposure, glutathione S-transferase polymorphisms, and colorectal cancer risk. *Am J Clin Nutr*. 2010;91(3):704-11.
42. Huang ZL, Liu HY. Expression of galectin-3 in liver metastasis of colon cancer and the inhibitory effect of modified citrus pectin. *Nan Fang Yi Ke Da Xue Xue Bao*. 2008;28(8):1358-61.
43. Liu HY, Huang ZL, Yang GH, et al. Inhibitory effect of modified citrus pectin on liver metastases in a mouse colon cancer model. *World J Gastroenterol*. 2008;14(48):7386-91.
44. Vitaglione P, Fogliano V, Pellegrini N. Coffee, colon function and colorectal cancer. *Food Funct*. 2012;3(9):916-22.
45. Choi DW, Lim MS, Lee JW, et al. The cytotoxicity of kahweol in HT-29 human colorectal cancer cells is mediated by apoptosis and suppression of heat shock protein 70 expression. *Biomol Ther (Seoul)*. 2015;23(2):128-33.
46. Budhathoki S, Iwasaki M, Yamaji T, et al. Coffee intake and the risk of colorectal adenoma: The colorectal adenoma study in Tokyo. *Int J Cancer*. 2015;137(2):463-70.
47. Nakamura T, Ishikawa H, Mutoh M, et al. Coffee prevents proximal colorectal adenomas in Japanese men: a prospective cohort study. *Eur J Cancer Prev*. 2015.
48. Lambert JD, Hong J, Yang GY, et al. Inhibition of carcinogenesis by polyphenols: evidence from laboratory investigations. *Am J Clin Nutr*. 2005;81(1 Suppl):284s-91s.
49. Mahmoud NN, Carothers AM, Grunberger D, et al. Plant phenolics decrease intestinal tumors in an animal model of familial adenomatous polyposis. *Carcinogenesis*. 2000;21(5):921-7.
50. Volate SR, Davenport DM, Muga SJ, et al. Modulation of aberrant crypt foci and apoptosis by dietary herbal supplements (quercetin, curcumin, silymarin, ginseng and rutin). *Carcinogenesis*. 2005;26(8):1450-6.
51. Buhrmann C, Shayan P, Popper B, et al. Sirt1 is required for resveratrol-mediated chemopreventive effects in colorectal cancer cells. *Nutrients*. 2016;8(3).
52. Yang S, Li W, Sun H, et al. Resveratrol elicits anti-colorectal cancer effect by activating miR-34c-KITLG in vitro and in vivo. *BMC Cancer*. 2015;15:969.
53. Patel KR, Brown VA, Jones DJ, et al. Clinical pharmacology of resveratrol and its metabolites in colorectal cancer patients. *Cancer Res*. 2010;70(19):7392-9.
54. Howells LM, Berry DP, Elliott PJ, et al. Phase I randomized, double-blind pilot study of micronized resveratrol (SRT501) in patients with hepatic metastases--safety, pharmacokinetics, and pharmacodynamics. *Cancer Prev Res (Phila)*. 2011;4(9):1419-25.
55. Bobe G, Albert PS, Sansbury LB, et al. Interleukin-6 as a potential indicator for prevention of high-risk adenoma recurrence by dietary flavonols in the polyp prevention trial. *Cancer Prev Res (Phila)*. 2010;3(6):764-75.
56. Atashpour S, Fouladdel S, Movahhed TK, et al. Quercetin induces cell cycle arrest and apoptosis in CD133(+) cancer stem cells of human colorectal HT29 cancer cell line and enhances anticancer effects of doxorubicin. *Iran J Basic Med Sci*. 2015;18(7):635-43.
57. Irving GR, Iwujii CO, Morgan B, et al. Combining curcumin (C3-complex, Sabinsa) with standard care FOLFOX chemotherapy in patients with inoperable colorectal cancer (CUFOX): study protocol for a randomised control trial. *Trials*. 2015;16:110.
58. Zhang Z, Chen H, Xu C, et al. Curcumin inhibits tumor epithelial-mesenchymal transition by downregulating the Wnt signaling pathway and upregulating NKD2 expression in colon cancer cells. *Oncol Rep*. 2016;35(5):2615-23.
59. Rajitha B, Belalcazar A, Nagaraju GP, et al. Inhibition of NF-kappaB translocation by curcumin analogs induces G0/G1 arrest and downregulates thymidylate synthase in colorectal cancer. *Cancer Lett*. 2016;373(2):227-33.
60. Huang YT, Lin YW, Chiu HM, et al. Curcumin induces apoptosis of colorectal cancer stem cells by coupling with CD44 marker. *J Agric Food Chem*. 2016;64(11):2247-53.
61. James MI, Iwujii C, Irving G, et al. Curcumin inhibits cancer stem cell phenotypes in ex vivo models of colorectal liver metastases, and is clinically safe and tolerable in combination with FOLFOX chemotherapy. *Cancer Lett*. 2015;364(2):135-41.
62. Carroll RE, Benya RV, Turgeon DK, et al. Phase IIa clinical trial of curcumin for the prevention of colorectal neoplasia. *Cancer Prev Res (Phila)*. 2011;4(3):354-64.
63. Di Francesco L, Lopez Contreras LA, Sacco A, et al. New insights into the mechanism of action of aspirin in the prevention of colorectal neoplasia. *Curr Pharm Des*. 2015;21(35):5116-26.
64. Cao Y, Nishihara R, Wu K, et al. Population-wide impact of long-term use of aspirin and the risk for cancer. *JAMA Oncol*. 2016.
65. Anwar MA, Kheir WA, Eid S, et al. Colorectal and prostate cancer risk in diabetes: Metformin, an actor behind the scene. *J Cancer*. 2014;5(9):736-44.
66. He XK, Su TT, Si JM, et al. Metformin is associated with slightly reduced risk of colorectal cancer and moderate survival benefits in diabetes mellitus: a meta-analysis. *Medicine (Baltimore)*. 2016;95(7):e2749.
67. Higurashi T, Hosono K, Takahashi H, et al. Metformin for chemoprevention of metachronous colorectal adenoma or polyps in post-polypectomy patients without diabetes: a multicentre double-blind, placebo-controlled, randomised phase 3 trial. *Lancet Oncol*. 2016.
68. Lefranc F, Yeaton P, Brotchi J, et al. Cimetidine, an unexpected anti-tumor agent, and its potential for the treatment of glioblastoma (review). *Int J Oncol*. 2006;28(5):1021-30.
69. Natori T, Sata M, Nagai R, et al. Cimetidine inhibits angiogenesis and suppresses tumor growth. *Biomed Pharmacother*. 2005;59(1-2):56-60.
70. Tomita K, Izumi K, Okabe S. Roxatidine- and cimetidine-induced angiogenesis inhibition suppresses growth of colon cancer implants in syngeneic mice. *J Pharmacol Sci*. 2003;93(3):321-30.
71. Adams WJ, Lawson JA, Morris DL. Cimetidine inhibits in vivo growth of human colon cancer and reverses histamine stimulated in vitro and in vivo growth. *Gut*. 1994;35(11):1632-6.
72. Adams WJ, Lawson JA, Nicholson SE, et al. The growth of carcinogen-induced colon cancer in rats is inhibited by cimetidine. *Eur J Surg Oncol*. 1993;19(4):332-5.
73. Adams WJ, Morris DL, Ross WB, et al. Cimetidine preserves non-specific immune function after colonic resection for cancer. *Aust N Z J Surg*. 1994;64(12):847-52.
74. Matsumoto S, Imaeda Y, Umamoto S, et al. Cimetidine increases survival of colorectal cancer patients with high levels of sialyl Lewis-X and sialyl Lewis-A epitope expression on tumour cells. *Br J Cancer*. 2002;86(2):161-7.
75. Adams WJ, Morris DL. Short-course cimetidine and survival with colorectal cancer. *Lancet*. 1994;344(8939-8940):1768-9.
76. Mariani F, Sena P, Roncucci L. Inflammatory pathways in the early steps of colorectal cancer development. *World J Gastroenterol*. 2014;20(29):9716-31.
77. Chong ES. A potential role of probiotics in colorectal cancer prevention: review of possible mechanisms of action. *World J Microbiol Biotechnol*. 2014;30(2):351-74.
78. Kowalczyk M, Siermontowski P, Mucha D, et al. Chromoendoscopy with a standard-resolution colonoscope for evaluation of rectal aberrant crypt foci. *PLoS One*. 2016;11(2):e0148286.
79. El Zoghbi M, Cummings LC. New era of colorectal cancer screening. *World J Gastrointest Endosc*. 2016;8(5):252-8.
80. Floer M, Meister T. Endoscopic improvement of the adenoma detection rate during colonoscopy - where do we stand in 2015? *Digestion*. 2016;93(3):202-13.
81. Hamilton SR. The adenoma-adenocarcinoma sequence in the large bowel: variations on a theme. *J Cell Biochem Suppl*. 1992;16g:41-6.
82. Brenner H, Chang-Claude J, Seiler CM, et al. Protection from colorectal cancer after colonoscopy: a population-based, case-control study. *Ann Intern Med*. 2011;154(1):22-30.
83. Johnson JJ, Mukhtar H. Curcumin for chemoprevention of colon cancer. *Cancer Lett*. 2007;255(2):170-81.

Enhanced Support for Healthy Inflammation Response

SUPER Bio-Curcumin®

Super Bio-Curcumin® features **BCM-95®**, a patented curcumin that absorbs up to **7 times** better than standard curcumin,¹ making this product ideal for those seeking to:

- Inhibit the pro-inflammatory effects of NF-kappaB activation,²
- Inhibit the pro-inflammatory activities of cyclooxygenase, lipoxygenase, and cytokines,³
- Support normal cell-cycle growth,⁴
- Maintain immune system health,⁵
- Promote healthy brain function,⁶
- Help maintain normal, healthy platelet function,⁷
- Support natural defenses against estrogen-mimicking chemicals,⁸ and
- Benefit joint health and bowel function.⁹

Life Extension® also provides **Advanced Bio-Curcumin® with Ginger & Turmerones** for those seeking the additional benefits of added ginger:^{10,11}

- Nearly *twice* the support for a healthy inflammatory response,
- Nearly *twice* the immune health support, and
- About *twice* the free radical fighting support.



Super Bio-Curcumin®

Item #00407 • 60 vegetarian capsules

Non-GMO.

	Retail Price	Your Price
1 bottle	\$38	\$28.50
4 bottles		\$26.25 each

Advanced Bio-Curcumin® with Ginger and Turmerones

Item #01924 • 30 softgels

Non-GMO.

	Retail Price	Your Price
1 bottle	\$30	\$22.50
4 bottles		\$20.25 each



References

1. *Indian J Pharm Sci.* 2008 Jul-Aug;70(4):445-9.

2. *AAPS J.* 2014 Jul;16(4):649-57.

3. *Ceska Slov Farm.* 2014 Feb;63(1):26-31.

4. *AAPS J.* 2009 Sep;11(3):495-510.

5. *Anesth Analg.* 2014 Jun;118(6):1336-44.

6. *Zhongguo Zhong Yao Za Zhi.* 2014 Oct;39(19):3846-9.

7. *Platelets.* 2011;22(4):270-6.

8. *Phytother Res.* 2014 Oct;28(10):1553-60.

9. *Pharmacol Res.* 2015 May-Jun;95-96C:71-81.

10. *Int J Pharmacol.* 2009;5(6):333-45.

11. *Food Nutr Res.* 2009;48(3):148-52.

CAUTION: Do not take if you have gallbladder problems or gallstones. If you are taking anti-coagulant or anti-platelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product.

Bio-Curcumin® and BCM-95® are registered trademarks of Dolcas-Biotech, LLC. U.S. Patent Nos. 7,883,728, 7,736,679 and 7,879,373.

To order **Super Bio-Curcumin®** or **Advanced Bio-Curcumin® with Ginger & Turmerones**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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.

IFOS Highest Independent 5 Star Rating International Fish Oil Society For Over Nine Years

HIGHLY PURIFIED ALASKAN FISH OIL Super Omega-3

FISH OIL + OLIVE EXTRACT + SESAME LIGNANS

Broad-spectrum, Mediterranean health benefits of fish oil, olive oil polyphenols, and sesame lignans for heart and brain health.

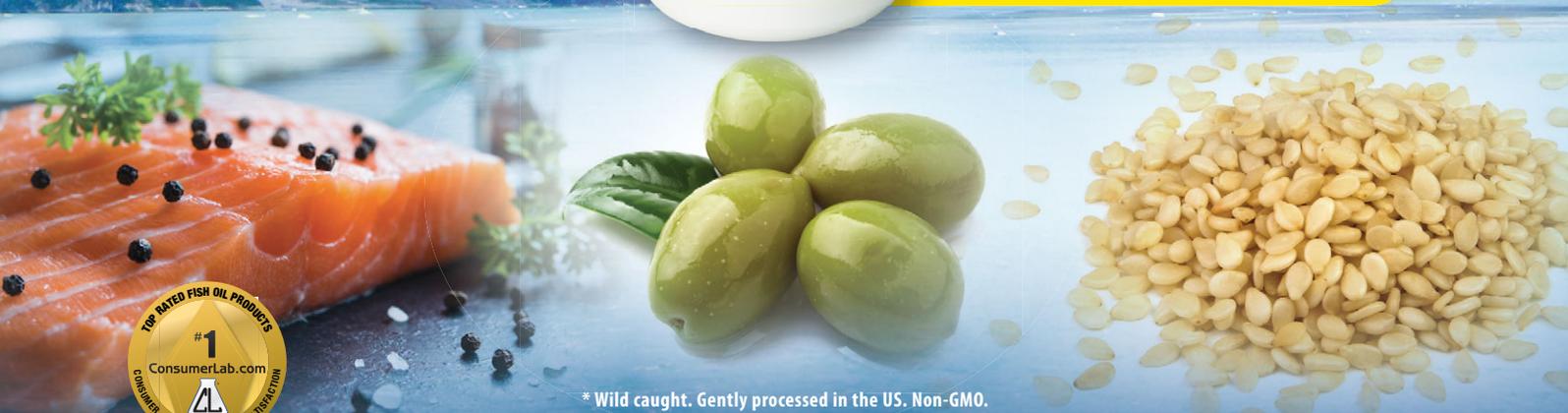
- Pure fish oil from sustainable sources in pristine waters in Alaska*, **highest 5-star rating** by leading independent third-party testing organization (**IFOS**).
- Provides the polyphenol equivalent of **8 to 12 tablespoons** of heart-healthy **extra virgin olive oil**.
- Specialized support against free radical oxidation with **sesame lignans**, a novel component of the heart-healthy **Mediterranean diet**.



Super Omega-3

Item #01982 • 120 softgels

	Retail Price	Your Price
1 bottle	\$32	\$24
4 bottles		\$21 each
10 bottles		\$17.05 each



* Wild caught. Gently processed in the US. Non-GMO.



To order **Super Omega-3**, call **1-800-544-4440** or visit **www.LifeExtension.com**

CAUTION: If you are taking anti-coagulant or anti-platelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product. Supportive but not conclusive research shows that consumption of EPA and DHA omega-3 fatty acids may reduce the risk of coronary heart disease.

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

INCREASE **SKELETAL STRENGTH** AND BOOST **BONE DENSITY**

It takes more than **calcium** to build strong bones...a host of skeletal-promoting nutrients are needed.

Bone Restore supplies a highly *absorbable* form of **calcium** in a combination of critical bone-building compounds:

- **Boron,**
- **Vitamin D3,**
- **Magnesium,**
- **Manganese,**
- **Zinc,**
- **Silicon, and**
- **Vitamin K2 (long-acting MK-7 form).**

Vitamin K2 is essential to maintain skeletal integrity.



Bone Restore with Vitamin K2

Item #01727 • 120 capsules

	Retail Price	Your Price
1 bottle	\$24	\$18
4 bottles		\$16.50 each

Bone Restore without Vitamin K

For those taking **Super K** or **Health Booster**, additional vitamin K2 is not needed. **Bone Restore** (without vitamin K) is available as **Item #01726** at a slightly lower price.

To order **Bone Restore with Vitamin K2**, call **1-800-544-4440** or visit **www.LifeExtension.com**

Non-GMO

CAUTION: If you are taking anti-coagulant or anti-platelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product.

Note: Those taking anticoagulant drug Coumadin® (warfarin) should use Bone Restore without K2.

Fruitex B® and OsteoBoron® are registered trademarks of VDF Futureceuticals, Inc. U.S. patent #5,962,049. DimaCal® and TRAACS® are registered trademarks of Albion Laboratories, Inc. Malate is covered by U.S. Patent 6,706,904 and patents pending.

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.



Novel Method Uncovers Younger-Looking Skin

Microdermabrasion is one of the top five minimally invasive topical procedures performed in the United States.¹

This mechanical form of exfoliation has increased in popularity in recent years due to its effectiveness in revitalizing aging skin on the face and across the entire body.^{2,3}

Body microdermabrasion can transform dull, thick, and peeling skin into soft, silky, and smooth youthful skin.

However, this in-office treatment can be expensive and also carries potential health risks that leave many individuals seeking safer, effective alternatives.

Fortunately, a unique body exfoliator has been formulated with ingredients that provide skin rejuvenating effects similar to microdermabrasion but without the significant drawbacks. The result is a safe, affordable, and gentler at-home alternative that delivers the appearance of younger, healthier, and more radiant skin.

Microdermabrasion: Multiple Benefits and Drawbacks

The outermost layer of the skin, called the *stratum corneum*, is composed of dead cells that form a protective outer barrier and prevent moisture loss.⁴ As new replacement cells migrate to the surface, these dead cells slough off. This continuous natural exfoliation maintains skin integrity and function.⁵

As the years pass, the migration of replacement cells to the surface of the skin slows down, leading to the accumulation of dead cells in the stratum corneum that produce changes associated with premature aging.⁶

So, in an attempt to overcome this problem, a growing number of individuals have turned to a non-surgical procedure originally developed in Italy in 1985 called **microdermabrasion** that primarily targets the facial area.⁷ This technique uses a machine to spray micro-crystals onto the face and then gently remove them simultaneously with aged surface skin.⁸ Since the body interprets this as a mild injury, it speeds up repair mechanisms to renew the skin surface—revealing fresh, younger cells.^{8,9}

Additionally, removing dead skin cells opens the door for more effective delivery of moisturizers, antioxidants, and collagen-boosting compounds deeper into the layers of the skin for a substantial anti-aging effect.¹⁰

Despite its effectiveness, microdermabrasion has several drawbacks. First, the machine is shared among patients, and, without proper sanitation measures, increases the risk of blood-borne pathogen transmission.¹¹ Second, most procedures are performed by non-physician personnel, which heightens the potential for human error. For instance, there is a greater likelihood of crystals entering the patient's eye and causing irritation. Third, treatments in a qualified professional setting such as a certified medical clinic can cost up to \$300 per session, with a total of 12 sessions typically required for results.¹² This quickly adds up to \$3,600 without taking into account other expenses like the initial consultation fee.

Development of a Safe and Affordable Alternative

But suppose you had access to an effective means of achieving similar results as machine-administered microdermabrasion without all the downsides. Better yet, what if it was designed to target the entire surface of the body, rather than solely the face, as in traditional microdermabrasion?

Life Extension[®] readers will be gratified to learn that such an intervention is now available in a body exfoliator—*featuring scientifically proven ingredients*—that can be safely used in the shower, especially on the neck and décolleté, hands, and back.





Body Exfoliator Rejuvenates Aging Skin

- A minimally invasive exfoliant procedure called microdermabrasion has increased in popularity in recent years due to its remarkable ability to rejuvenate aging facial skin.
- Its potential health risks, along with a hefty price tag, have many individuals seeking alternatives.
- A safe, affordable, and gentler at-home alternative in a body exfoliator has been developed that delivers the same results as microdermabrasion without all the notable drawbacks.
- This unique body exfoliator provides cleansing benefits, exfoliating properties, and stimulation of the skin.
- It is formulated with amber crystals and jojoba beads, as well as potent antioxidants and effective moisturizers that translate into fresher, more radiant, younger-looking skin.

This whole-body cleanser removes the buildup of dirt, grease, and other impurities—leaving your skin with a soft and clean feeling accompanied by a healthy, bright glow. Its exfoliating properties eliminate the rough, dry, and crepey skin characteristic of aging. Lastly, this body exfoliator effectively treats skin nuisances like calluses and back acne.

Let's now discuss the natural substances incorporated into this unique formula, and how they work in a complementary way to leave behind a more youthful appearance.

Amber Crystals and Jojoba Beads

Amber is a natural organic substance derived from fossilized resin trees in the present territory of central and northern Europe almost 45 million years ago. This stone has been historically used in the Baltic Sea region to treat a variety of ailments, from asthma to rheumatism to jaundice.¹³ Skin rejuvenation can now be added to this already impressive list of benefits thanks to the terpene compounds present in amber,¹³ which provide powerful free-radical scavenging and anti-aging properties.¹⁴

Amber crystals have been shown time after time to combat the tell-tale signs of aging. In dermatologist test cases, the daily use of an exfoliant containing amber crystals for one to three weeks was shown to produce the following improvements in patients with aging skin:¹⁵

- Improvement in **skin roughness** by **50%**
- Reduction in **dark hyperpigmentation** by **30%**
- Improvement in **sun-damaged skin** by more than **70%**

Another substance with exfoliating properties is **jojoba beads** (also known as jojoba esters) derived from jojoba oil. Since jojoba beads are biodegradable and possess a unique spherical shape,¹⁶ they are an ideal choice for gently exfoliating the skin without leaving behind any scratches or cuts. Additionally, jojoba esters fit the bill perfectly for improving skin hydration as its chemical composition closely resembles human sebum—the oily substance that naturally moisturizes and soothes the skin.¹⁷ In one clinical study, jojoba esters increased skin suppleness by approximately **25%** after eight hours.¹⁸ As an added bonus, jojoba esters have anti-inflammatory actions.^{19,20}

Antioxidant Tea Blend

Repeated exposure to ultraviolet radiation produces a chronic stream of free radicals that induce oxidative stress, which in turn slowly destroys key skin structures and triggers the formation of wrinkles, sagginess, and fine lines.^{21,22} Regular exfoliation allows extracts of red, green, and white tea to better penetrate the skin to combat free radicals and protect against oxidative stress.^{23,24}

Most of green tea's skin benefits have long been attributed to the potent polyphenol **epigallocatechin-3-gallate (EGCG)**, which quenches inflammation and prevents photoaging.²⁵ Red tea extract boasts a noteworthy one-two punch of aspalathin, a rare flavonoid that enhances concentration of free radical-fighting **superoxide dismutase (SOD)**,²⁶ and **alpha-hydroxy acids** that stimulate new cell growth and repair following exfoliation.²⁷

Lastly, white tea extract has been shown to inhibit enzymes that break down the structural proteins collagen and elastin, as well as possessing antioxidant power comparable to the aforementioned teas.²⁸



Seaweed Extract

Chondrus crispus, a form of red seaweed found in the Atlantic Ocean, acts as an emollient to help soothe, smooth, and hydrate aging skin.²⁹ With its wealth of anti-aging nutrients, from proteins, vitamins, and minerals to its high content of the moisturizing compound carrageenan, *Chondrus crispus* has the repertoire to nourish and restore the skin's healthy appearance.^{30,31}

Peppermint Oil

Peppermint oil contains a high concentration of **menthol**, which produces a cool, soothing sensation when applied to the skin.³² This refreshing effect, combined with its pain-relieving,³³ antimicrobial,³⁴ and antifungal properties,³⁵ makes peppermint oil an ideal compound to enhance the skin's healing capacity after exfoliation.

Summary

Microdermabrasion refers to a type of mechanical exfoliation that has been highly successful in revitalizing aging facial skin. While microdermabrasion has been proven to visibly improve the health and appearance of aging skin, its high cost and potential health risks have left many individuals seeking safer and effective alternatives.

Fortunately, a low-cost, at-home alternative exfoliator has been developed that offers deep cleansing benefits and overall stimulation of the skin all over the body. This unique body exfoliator contains ultra-fine amber crystals and jojoba beads, along with potent antioxidants and moisturizers, that work together to gently rub away skin imperfections to restore a youthful appearance. ●

Gary Goldfaden, MD, is a clinical dermatologist and lifetime member of the American Academy of Dermatology. He is the founder of Academy Dermatology in Hollywood, FL, and Cosmesis Skin Care. Dr. Goldfaden is a member of **Life Extension®**'s Medical Advisory Board.

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

References

- Available at: <http://www.plasticsurgery.org/Documents/news-resources/statistics/2015-statistics/top-five-cosmetic-plastic-surgery-procedures-2015.pdf>. Accessed April 11, 2016.
- Spencer JM. Microdermabrasion. *Am J Clin Dermatol*. 2005;6(2):89-92.
- Karimipour DJ, Karimipour G, Orringer JS. Microdermabrasion: an evidence-based review. *Plast Reconstr Surg*. 2010;125(1):372-7.
- Nemes Z, Steinert PM. Bricks and mortar of the epidermal barrier. *Exp Mol Med*. 1999;31(1):5-19.
- Madison KC. Barrier function of the skin: "la raison d'être" of the epidermis. *J Invest Dermatol*. 2003;121(2):231-41.
- Farage MA, Miller KW, Elsner P, et al. Characteristics of the aging skin. *Adv Wound Care (New Rochelle)*. 2013;2(1):5-10.
- Loesch MM, Somani AK, Kingsley MM, et al. Skin resurfacing procedures: new and emerging options. *Clin Cosmet Investig Dermatol*. 2014;7:231-41.
- Kirkland EB, Hantash BM. Microdermabrasion: molecular mechanisms unraveled, part 2. *J Drugs Dermatol*. 2012;11(9):e10-7.
- Kirkland EB, Hantash BM. Microdermabrasion: molecular mechanisms unraveled, part 1. *J Drugs Dermatol*. 2012;11(9):e2-9.
- Freedman BM. Topical antioxidant application enhances the effects of facial microdermabrasion. *J Dermatolog Treat*. 2009;20(2):82-7.
- Shelton RM. Prevention of cross-contamination when using microdermabrasion equipment. *Cutis*. 2003;72(4):266-8.
- Available at: <http://www.skinabrasion.net/cost.html>. Accessed April 11, 2016.
- Available at: <http://www.centerchem.com/Products/DownloadFile.aspx?FileID=6805>. Accessed April 11, 2016.
- Gonzalez-Burgos E, Gomez-Serranillos MP. Terpene compounds in nature: a review of their potential antioxidant activity. *Curr Med Chem*. 2012;19(31):5319-41.
- Clinical case histories provided by Gary Goldfaden, MD. August 5, 2015.
- Available at: <http://thejobabaoil.com/jojoba-beads/>. Accessed April 14, 2016.
- Available at: <http://www.centerchem.com/Products/DownloadFile.aspx?FileID=6962>. Accessed April 14, 2016.
- Christensen MS, Packman EW. Skin surface softening effects of jojoba and its derivatives. *Proceedings from the seventh international conference on jojoba and its uses*. Champaign, IL: American Oil Chemists' Society; 1988.
- Pazyar N, Yaghoobi R, Ghassemi MR, et al. Jojoba in dermatology: a succinct review. *G Ital Dermatol Venereol*. 2013;148(6):687-91.
- Habashy RR, Abdel-Naim AB, Khalifa AE, et al. Anti-inflammatory effects of jojoba liquid wax in experimental models. *Pharmacol Res*. 2005;51(2):95-105.
- Fisher GJ, Kang S, Varani J, et al. Mechanisms of photoaging and chronological skin aging. *Arch Dermatol*. 2002;138(11):1462-70.
- Imokawa G. Recent advances in characterizing biological mechanisms underlying UV-induced wrinkles: a pivotal role of fibroblast-derived elastase. *Arch Dermatol Res*. 2008;300 Suppl 1:S7-20.
- Katiyar SK, Afaq F, Perez A, et al. Green tea polyphenol (-)-epigallocatechin-3-gallate treatment of human skin inhibits ultraviolet radiation-induced oxidative stress. *Carcinogenesis*. 2001;22(2):287-94.
- Camouse MM, Domingo DS, Swain FR, et al. Topical application of green and white tea extracts provides protection from solar-simulated ultraviolet light in human skin. *Exp Dermatol*. 2009;18(6):522-6.
- Katiyar SK. Skin photoprotection by green tea: antioxidant and immunomodulatory effects. *Curr Drug Targets Immune Endocr Metabol Disord*. 2003;3(3):234-42.
- Baba H, Ohtsuka Y, Haruna H, et al. Studies of anti-inflammatory effects of Rooibos tea in rats. *Pediatr Int*. 2009;51(5):700-4.
- Yamamoto Y, Uede K, Yonei N, et al. Effects of alpha-hydroxy acids on the human skin of Japanese subjects: the rationale for chemical peeling. *J Dermatol*. 2006;33(1):16-22.
- Thring TS, Hili P, Naughton DP. Anti-collagenase, anti-elastase and anti-oxidant activities of extracts from 21 plants. *BMC Complement Altern Med*. 2009;9:27.
- Available at: <http://jalgalbiomass.com/paper14vol3no4.pdf>. Accessed April 14, 2016.
- Available at: http://www.herbs2000.com/herbs/herbs_irish_moss.htm. Accessed April 14, 2016.
- Available at: <http://drs.nio.org/drs/handle/2264/489>. Accessed April 14, 2016.
- Eccles R. Menthol and related cooling compounds. *Journal of Pharmacy and Pharmacology*. 1994;46(8):618-30.
- Gaudio C, Hao J, Martin-Eauclaire MF, et al. Menthol pain relief through cumulative inactivation of voltage-gated sodium channels. *Pain*. 2012;153(2):473-84.
- Scholz Z, Molnar J, Hohmann J. Antimicrobial and antiplasmodial activities of essential oils. *Fitoterapia*. 2006;77(4):279-85.
- Pattnaik S, Subramanyam VR, Bapaji M, et al. Antibacterial and antifungal activity of aromatic constituents of essential oils. *Microbios*. 1997;89(358):39-46.



Restore Youthful-Looking Skin from the Inside Out

As skin ages, it loses suppleness and smoothness. A prime reason is loss of **ceramides** that are required for skin to retain its **moisture**.

Skin Restoring Phytoceramides contains wheat-derived ceramide oils in an **oral** capsule that **hydrates** the deepest dermal layers to nourish the entire body's skin.



Skin Restoring Phytoceramides with Lipowheat® • Item #01596
30 liquid vegetarian capsules • Non-GMO

	Retail Price	Your Price
1 bottle	\$25	\$18.75
4 bottles		\$17.25 each

To order **Skin Restoring Phytoceramides with Lipowheat®**,
call **1-800-544-4440**
or visit **www.LifeExtension.com**

Contains wheat. Gluten free. Lipowheat® is a registered trademark of Arco, Robertet Group, France.

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

Microdermabrasion is a form of exfoliation that **revitalizes** aging skin.^{1,2}

By removing the tough layer of dead cells on the skin's surface, **microdermabrasion** can help restore skin back to the soft, silky, smooth look and feel of youth.

The new **Ultimate MicroDermabrasion** provides:

- **Amber** crystals to smooth skin and help even out discoloration,³
- **Jojoba** crystals that exfoliate and boost skin suppleness,⁴
- **Green, red, and white tea** extracts to support skin-cell growth and collagen, elastin, and antioxidant levels,⁵⁻⁷
- **Seaweed** to smooth and hydrate the skin,⁸ and
- **Peppermint oil**, which provides a cooling, soothing sensation to exfoliated skin.⁹

Designed to be used **at home** for the entire body, **Ultimate MicroDermabrasion** can be used daily until youthful appearance and softness is restored. It can then be used as needed to maintain benefits.

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

For external use only. Avoid contact with eyes. If product contacts eyes, rinse thoroughly with water. If rash or irritation occurs, please discontinue use.

References

1. *Am J Clin Dermatol.* 2005;6(2):89-92
2. *Plast Reconstr Surg.* 2010;125(1):372-7
3. Case histories. Gary Goldfaden, MD.
4. Proceedings from the seventh international conference on jojoba and its uses. Champaign, IL: American Oil Chemists' Society; 1988.
5. *Curr Drug Targets Immune Endocr Metabol Disord.* 2003;3(3):234-42.
6. *Pediatr Int.* 2009;51(5):700-4.
7. *BMC Complement Altern Med.* 2009;9:27.
8. Available at: <http://jalgalbiomass.com/paper14vol3no4.pdf>. Accessed May 16, 2016.
9. *Journal of Pharmacy and Pharmacology.* 1994;46(8):618-30.

Skin-Rejuvenating Effects of MicroDermabrasion

Ultimate MicroDermabrasion
Item #80162 • 8 fl. oz. (240 ml.)

	Retail Price	Your Price
1 bottle	\$39	\$29.25
4 bottles		\$26.25 each



A line of skin care products developed by renowned dermatologist Gary Goldfaden, MD

Developed Exclusively for Life Extension®





A NEW ERA!

Curing Aging in Our Lifetime

by William Faloon,
Co-Founder of Life Extension Foundation®

Many of you know I've spent my career pursuing scientific methods to eradicate **biological aging** in humans.

By networking with like-minded individuals, I've helped introduce discoveries that have had profound impacts on human longevity.

In our early years, **Life Extension Foundation®** introduced the anti-aging properties of **coenzyme Q10** and the heart attack-prevention effects of **aspirin**.

These innovations spared countless lives. What was lacking up to now were therapies to reverse systemic **aging** in the elderly.

A pivotal turning point emerged in **2015**. The game-changers were laboratory findings demonstrating that **systemic rejuvenation** can be induced in old animals.

This prompted scientists we work with to design **clinical studies** whereby techniques that have **reversed aging** in the animal model can be tested in elderly people.

As we delved into the prospect of **human age reversal**, we began finding others who were planning similar research endeavors. We were not alone in conjecturing that a cure for human aging might be attainable relatively soon.

This article will enlighten you to an exciting conference in San Diego where a host of **age-reversal** initiatives will be presented.

Since our inception in 1980, the **Life Extension Foundation** has funded **biomedical research** aimed at extending lifespans beyond conventional expectations.

What's happening today with rejuvenation research is nothing short of a **medical renaissance**. It will likely have a far greater impact on human longevity than the discovery of antibiotics.

On **August 4-7, 2016**, in **San Diego, California**, you will have the opportunity to interact with the **physicians** and **scientists** who will oversee human rejuvenation studies.

The name of the event is **Revolution against Aging and Death** conference. It is sponsored by several non-profit groups. The focus is **human age-reversal**.

Before describing this unprecedented scientific gathering, let me discuss a few recent timelines so you can appreciate the rapid pace with which **age-reversal** technology has progressed.

We Were Desperate in 2005

In **2005**, I hosted a Sunday afternoon get-together with some of our dedicated supporters. I'll never forget an educated woman approaching me with a complaint. She was annoyed that she had an outlandish amount of money at her disposal but nowhere to contribute it to that had any chance of reversing biological aging.

She asked me if there were any direct donations she could make to **Life Extension Foundation** that would accelerate human age-reversal research. My response to her **blank check offer** was that I suffered the same dilemma.

Life Extension® was doing modestly well, but there were no projects in which meaningful human age-reversal theories could be clinically tested.

This was not the first time a wealthy person offered us donations for impossible-to-conduct clinical research. There were in fact instances of hostility



expressed to me by individuals saying, *"I'm offering to pay you to do human age reversal research... why aren't you taking my money?"*

The reality in those early years was that there were no viable human age-reversal concepts that could be clinically tested...no matter how much funding was offered.

The "Emergency" Summit of 2013

In **January 2013**, a group of anti-aging doctors and motivated individuals invited me to what they described as an **"emergency summit"** in South Florida. The credibility of the people involved mandated my attendance.

An inspirational speech given by the lead physician enlightened the group to the nature of the "emergency." Put simply, this doctor made it clear we were all **"aging to death,"** yet no one was doing anything to reverse this lethal trend. The group united behind the premise that scientific methods to reverse pathological aging had to be fast-tracked. They then turned and asked me, *"What should we do?"*

I reminded the group that they were already slowing premature aging via their healthy lifestyle choices, along with their hormone and supplement regimens. They understood this, but the question remained: What could be done to reverse the aging process that was slowly killing us all?

The group wanted to hear something positive, so I reminded them of data showing that the drug **metformin** had anti-aging properties that mimicked the longevity benefits of **calorie restriction**. One doctor stood up and offered to lead a clinical trial to document metformin's benefits. I replied that this would be a waste of money because there was already sufficient data to support the use of **metformin**.

After a few subsequent meetings, this South Florida group disbanded because there was nothing practical at the time that could be funded that would lead to rapid reversal of human aging.

Age Reversal goes Live in 2015!

Move forward to **2015** and a paradigm shift emerged. For the first time, research initiatives to induce meaningful, systemic human age reversal were evolving into scientific reality.

To put this into perspective, I'd been attending anti-aging conferences since **1978**, when the focus was on disease prevention. The concept of **age reversal** was viewed as something that would occur in the future.

In **2015**, that future arrived in the form of a **biomedical revolution!**



Unlike typical scientific conferences where you often get lost in the crowd, this unique event will bring like-minded individuals close together, with a range of age-reversal strategies presented by avant-garde scientists and lots of group follow-up discussions.

A listing of the speakers scheduled for **Revolution against Aging and Death** appears at the end of this article. We reserve the right to bring in last minute scientists who may be on the verge of major discoveries. What may surprise you is how far along these physician/scientists are in potentially making older people grow functionally younger.

Exponential Knowledge Eruption

Scientists at this moment are pushing through **human** studies with the objective of biologically transforming older people into measurably younger individuals. Experts believe that many aspects of pathologic aging can be eradicated using these newly developed therapies. The integrity of these findings is what prompted the **Life Extension Foundation** to cosponsor **Revolution against Aging and Death**, our first national **conference on aging** since 1986.

If it were not for the many scientists now engaged in human age reversal initiatives, **Life Extension** would NOT be involved in this conference. In other words, if all you were going to learn at this event was more information on preventing disease and slowing aging, this conference would not be that different from others you may have attended.

What distinguishes this conference from all others is a focus on **human age reversal** and how it can be accelerated through group collaboration.

A Different Kind of Gathering

If you consider what's included in this multi-day program, you'll realize what a bargain it is. In addition to the impressive speakers, the registration fee covers **healthy lunches** and **dinners** for two straight days in a fabulous hotel resort ballroom. This will allow for productive interactions to occur throughout the multi-day program and eliminates awkward breaks where you wonder what to do for an hour or two. The group is staying together throughout most of this historic meeting.

Multiple Reasons for You to Attend

There are multiple benefits beyond the illuminating experience you will gain by attending the **Revolution against Aging and Death** conference.

One opportunity is making personal connections with physicians who may already be planning to incorporate **age reversal** into their practice on an individual basis. This means you might gain access to regenerative therapies before formal clinical trials are initiated. Another reason for attending is to demonstrate solidarity amongst a huge group of like-minded people. This is important because there may be hundreds of representatives from Wall Street, Silicon Valley and Big Pharma at this conference looking for investment opportunities, along with dozens of media.

We expect over **1,000 people** will attend the conference. Can you imagine how much more energy will be generated if we pack the room with **4,000** individuals whose priority is to gain quick access to age-reversal therapies? This single event could open the floodgates (i.e. billions of dollars) of private investment aimed at rapidly developing validated **age-reversal** treatments.

In case you haven't figured this out yet, we are seeking to bring the science of age reversal to the public forefront. We want to transform **regenerative medicine** into the most important avenue of research that humanity pursues.

Discounted Registration Fee

Most conferences are put on by commercial entities looking to turn a profit, so they charge substantial fees over and above their expenses.

In contrast, this unique conference is sponsored by several **nonprofit** organizations. The goal is to unite activists, enthusiasts, scientists and entrepreneurs who want to accelerate widespread availability of **age-reversal** technologies. By eliminating the profit incentive, we've been able to make participation in this event affordable.

Through July 14, the registration fee is **\$577** each, but we've arranged for **Life Extension** supporters to save \$100 and pay just **\$477**. From July 15 to August 3, the cost is **\$627**, **\$527** for **Life Extension** supporters. Couples who sign up get an additional **\$50** off these promotions. Again, the fee includes two days of healthy lunches and dinners.

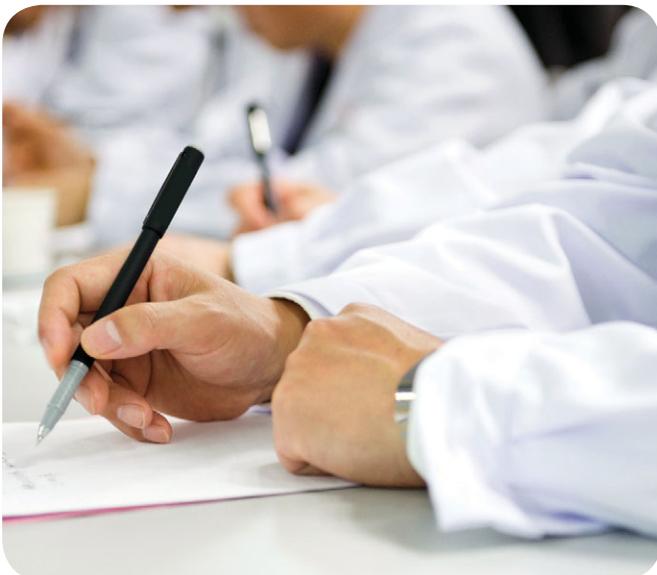
To get the best price, enroll by **July 14** and pay only **\$477**.

Dates and Location

The location of this unprecedented event is the **Town and Country Resort in San Diego, California**. This historic gathering begins **Thursday evening, August 4**, and ends **Sunday, August 7** at 1 p.m. Based on the number of pre-registrations, we expect over **1,000** attendees.

To enroll at the discounted rate, log on to **www.RAADfest.com**. Enter "**LEF**" in the promotional code box, and \$100 will be subtracted from your registration fee.

If you don't like dealing with websites, just call **1-866-595-6577** (24 hours). A live operator will register you at the **discounted** price.



Uniting Longevity Enthusiasts

Life Extension Foundation is the world's **largest** consumer-based longevity group. We fund **millions of dollars** each year in **research** aimed at radically extending the healthy human life span.

Major leaps in knowledge about **aging** have occurred over the past two years. These advances mandate that groups involved in **longevity research** unite to make **age reversal** a reality for those alive today.

In this era of electronic communication, it's vital to have **in-person** interactions with like-minded people to share information, inspiration and insight. This helps create a sense of connection and belonging that strengthens our resolve to gain control of aging in our lifetime. For the first time in medical history, the prospect of human age reversal has become a realistic probability. This represents a renaissance not just in the scientific arena, but also in the way society views human longevity potential.

I hope to personally meet many of you during this unprecedented educational venture **August 4-7** in **San Diego**.

For longer life,

William Faloon

P.S. There are additional perks included with the registration fee that you can review by visiting the official **Revolution against Aging and Death** website at **www.RAADfest.com**.

In addition to **Life Extension's** William Faloon, presenters at the event include top anti-aging researchers and experts such as:

Ray Kurzweil, Dr. Bill Andrews, Ilana Arzt, Joe Bardin, Greta Blackburn, Bernadeane Brown, Dr. Kristen Comella, Jose Luis Cordeiro, Dr. Brant Cortright, Dr. Aubrey de Grey, Maria Entraigues-Abramson, Dr. Greg Fahy, Dr. Ben Goertzel, Michael Greve, Laurie Handlers, Dr. Steve Hruby, Zoltan Istvan, Dr. Ron Klatz, Dr. Moss Jackson, David Kekich, Dr. Dobri Kiproff, Dr. Mark Gordon, Dipnarine Maharaj, MD, Nuno R.B. Martins, Dr. Max More, Peter Nygard, Liz Parrish, Dr. Kirk Parsley, Steve Perry, Molly Sheridan, James Strole, Suzanne Somers, Neal VanDeree, Dr. Natasha Vita-More, Dr. Kristen Willeumier, and Dr. Alex Zhavoronkov.

Age Reversal Research Initiatives at RAADfest

There will be over 30 speakers at RAADfest 2016 making presentations on a wide range of health topics. Listed on this page are speakers that will describe age reversal research initiatives they are directly spearheading.



Dipnarine Maharaj, MD

Reversal of pathologic aging processes along with restoration of functional immune status using stem cell mobilized plasma components from young donors.

The bone marrow of healthy 21-year-olds contains an abundance of youthful stem cells, immune cells, and paracrine factors including plasma exosomes, proteins, and cytokines.

Using an FDA-approved factor that mobilizes bone marrow and a proprietary protocol for its administration, rapid release of pro-youth factors is enhanced. Through a process known as apheresis, these pro-youth factors are harvested in the plasma of the young donors and administered to older people to induce restoration of healthy immune function.

This study is to be conducted at the South Florida Bone Marrow/Stem Cell Institute. It plans to initially enroll about 30 study subjects who meet the eligibility criteria to receive the mobilized plasma from blood-matched healthy, young donors.

Dr. Maharaj is a hematologist/oncologist stem cell transplant physician and director of the South Florida Bone Marrow Stem Cell Institute in Boynton Beach, Florida.



Greg Fahy, PhD

Reversal of human immunological aging, along with suppression of autoimmune factors

Immune senescence is initiated by the atrophy of the thymus gland near the time of puberty. Thymic atrophy can be reversed in animals and in young AIDS patients, but this has not been shown in normal aging humans.

Dr. Fahy reports on an ongoing human aging reversal trial at Stanford University he is overseeing using FDA-approved agents that in combination should safely regenerate not only the thymus, but also other organs as well. Side effects have been minimal, and subjective responses have been positive. Spinoff applications include elimination of diabetes, other autoimmune diseases, and transplant rejection.

Greg Fahy is a world renowned authority on aging and cryobiology and is chief scientific officer at 21st Century Medicine, a wholly owned subsidiary of the Life Extension Foundation.



Dobri Kiprov, MD

Inhibition of senescent cell behavior in elderly subjects via removal of aged plasma and infusion of young plasma components...Prophylactic plasma exchange in an attempt to suppress inflammation and induce age reversal

Dr. Kiprov's team was the first in the world to demonstrate that replacing the systemic milieu restores the function of impaired immunocompetent cells in patients with autoimmune diseases.

Based on this observation, he developed a model whereby removing senescent plasma and replacing it with factors derived from young plasma may reset the responsible biochemical pathways to their younger states.

Dr. Kiprov's team has received an Internal Review Board (IRB) approval to conduct a trial study on Intermittent Heterochronic Plasma Exchange as a Modality for Preventing Cellular Senescence that will seek to prove age reversal in a group of older volunteer (human) study subjects.

Dobri Kiprov is an internationally recognized pioneer and expert in the field of therapeutic apheresis. He is chief of the Division of Immunotherapy at California Pacific Medical Center in San Francisco and medical director of Apheresis Care Group.



Aubrey de Grey, PhD

Why aging's days are numbered, and the diseases of aging even more so

Over 15 years ago, I proposed that age-related disease and disability might be easier to postpone and prevent by periodically repairing the accumulating damage that causes it, rather than by slowing the body's creation of that damage as almost all researchers had historically assumed. This concept, which is now widely accepted as promising by the expert community, is being pursued by laboratories worldwide and especially by SENS Research Foundation's in-house and extramural laboratory research programs. Progress is accelerating, as illustrated both by increasingly high-profile academic publications and by the translation of these projects to spin-out startup companies.

Aubrey de Grey is Chief Scientific Officer at SENS Research Foundation and internationally recognized leader in the field of regenerative medicine research.



Steve Perry
Growth Differentiation Factor 11 (GDF11)

GDF11 is a naturally occurring peptide that has been cited in several papers as having a capacity to restore aging muscles, hearts and brains in mice, while researchers at Harvard proclaim it may have significant age reversal effects.

GDF11 had never been tested in humans until Steve Perry became patient zero on June 7, 2014.

Parabiosis research strongly suggests that blood borne peptides, especially GDF11, are a key component of the “aging program.” Steve and about 20 volunteers are working on a GDF11 dosing strategy as well as measuring key biomarkers in the cardiac, pulmonary, cognitive, skin elasticity and immune areas to determine GDF11’s effectiveness. Results of this study will be presented.

Steve Perry founded GDF11 Rejuvenation, Inc. in June, 2014, to explore the efficacy of GDF11 supplementation.



Liz Parrish and Bill Andrews, PhD
Extending telomeres to reverse the clock to aging and declining health

Lengthening of telomeres using the enzyme telomerase is a protocol that has demonstrated age reversal in experimental models.

Longer telomeres not only increased the energy in old mice, but also restored their ability to breed, restored their brain function, revitalized their internal organs, and made them look and behave young again.

Elizabeth Parrish and Bill Andrews will be discussing their joint venture, called BioViva Fiji, to use gene therapy to make this technology of telomere lengthening possible for humans in the very near future.

Elizabeth Parrish is the Founder and CEO of BioViva USA. She is a humanitarian, entrepreneur and innovator, and a leading voice for genetic cures for aging. Recently she became patient zero in the highly publicized first gene therapy effort to reverse aging.

Bill Andrews is the Founder and CEO of Sierra Sciences. His scientific background for 35 years in the biotech industry has led to many medical breakthroughs including the discovery of human telomerase. His career for the last 17 years has focused specifically on finding ways to extend human lifespan and health span through the intervention of telomere shortening.



Alex Zhavoronkov, PhD
Deep learning and artificial intelligence advances that can accurately evaluate efficacy of age-reversal therapies

Dr. Zhavoronkov leads an international group of scientists using advances in deep learning and artificial intelligence to develop comprehensive and accurate

biomarkers of human aging to measure the effectiveness of many interventions. His team is also developing a set of molecules aimed at reversing age-related changes while targeting cancer and common degenerative disorders. Over the past 3 years his group published over 60 peer-reviewed publications in leading scientific journals and two books including *The Ageless Generation* by Palgrave Macmillan.

Alex Zhavoronkov is chief science officer of the Biogerontology Research Foundation, a UK-based registered charity supporting aging research worldwide and a CEO of Insilico Medicine at the Emerging Technology Centers located at the Johns Hopkins University.

His talk at RAADFest will focus on developing comprehensive and actionable biomarkers of human aging, where ensembles of deep neural networks trained on multimodal data coming from tens of thousands of patients predict both chronological and biological age of the patient at any point in time. He will expand on how artificially intelligent systems in coup with human domain expertise in drug discovery can be used to discover, personalize and test geroprotectors intended to slow down or even repair age-related damage.



Ben Goertzel, PhD
Using advanced artificial intelligence to identify pathways to super-longevity and super-abundant health

Dr. Goertzel has been working for decades on advancing artificial intelligence technology and using artificial intelligence to analyze longevity-related datasets, with exciting results.

He will indicate the kinds of greater insights that would be expected to emerge from larger and more comprehensive initiatives using artificial intelligence-based analysis/simulation of the biological underpinnings of aging and health.

Ben Goertzel is the chief scientist of Hanson Robotics, a Hong Kong robotics company, and Aidya Limited, an investment management firm. He also leads the open-source OpenCog Artificial General Intelligence (AGI) project and the annual AGI conference series. In the area of longevity research, he has been the foremost advocate and practitioner of the application of artificial intelligence technology to the analysis of genomics data regarding longevity.



Dr. Kristin Comella,
Harnessing your body's own stem cells to reverse the effects of disease and aging

The field of regenerative medicine is rapidly growing with an emphasis on stem cell therapies and the promise of cures for everything from acute injuries to chronic degenerative diseases. The list of indications is expanding as more studies are published to demonstrate safety and efficacy.

Regenerative medicine is the process of replacing or regenerating human cells, tissues or organs to restore or establish normal function. The concept is that damaged tissue can be restored by using the body's own healing mechanism to promote repair. This new branch of medicine may change the course of chronic diseases and standard clinical therapies. The team at US Stem Cell is harvesting stem cells from fat tissue for use in orthopedics, autoimmune diseases, neurological disorders and more.

Kristin Comella is a world-renowned expert on regenerative medicine with a focus on adipose-derived stem cells. She was named Number 1 on the Academy of Regenerative Practices list of Top 10 Stem Cell Innovators.



Peter Nygård
Somatic cell nuclear transfer to reverse cellular aging in vivo

Peter Nygård is a successful entrepreneur who has invested millions of dollars in the pursuit of research aimed at transitioning aging human cells in the living body back into an autologous embryonic state via a process known as Somatic Cell Nuclear Transfer (SCNT).

In 2009, as he got started on the path to reverse aging, he suddenly realized that the trip from 70 to 80 was not a nice trip—that something bad can and will happen on that trip. He wisely decided “not to go on that trip.”

His transition from an old 69-year-old to a young 75-year-old is a motivating example of the potential of this technology.

Peter Nygård practices and supports all paths to regenerative medicine and is best known for his SCNT work. He is one of the few humans who have used SCNT technology on themselves. He has also written laws for countries to make the use legal. He is heavily financing SCNT translational research throughout the world.



William Faloon
Connecting the dots may be all that's required to cure aging

A cure for common bacterial infections was published in 1929, yet widespread availability of antibiotics did not occur until 1946. A similar situation exists today whereby aging has been reversed in experimental animal models, but transforming this into human rejuvenation therapies has been delayed.

Bill Faloon will lay out a plan whereby merely connecting the various areas of expertise and commitment will result in laboratory research being rapidly translated into clinical trials and, when successful, made widely available to spare humans today from the lethal impact of age-related disorders.

Bill Faloon is co-founder of the Life Extension Foundation, the world's largest consumer-based longevity organization.



Suzanne Somers
Enlightening the world to the prospect of age reversal

Public acceptance of the medical concept of age reversal will accelerate greater allocation of resources to the field of longevity research.

Suzanne Somers, author, lecturer, entertainer, entrepreneur, is the second most recognized female celebrity (after Oprah) in the United States and has exerted a powerful influence on public opinion as it relates to taking charge of one's personal health and well-being.

She has written 26 books, 14 *New York Times* best sellers courageously taking on mainstream medicine by introducing America and then the world to the remarkable benefits to health and aging with bioidentical hormone replacement.

Suzanne's passion for alternatives and non-drug approaches to health and aging include new thinking about cancer and battling environmental pollution and chemicals. To access longevity and quality of life requires a shift in thinking and Suzanne clearly lives her message. She is a breathtaking example of health, sexuality and as she calls it... A NEW WAY TO AGE.

To Attend This Historic Conference

To enroll at the **Revolution against Aging and Death** conference in San Diego (**August 4-7, 2016**) at the discounted rate, log on to www.RAADfest.com. Enter “LEF” in the promotional code box, and **\$100** will be subtracted from your registration fee. If you don't like dealing with websites, just call **1-866-595-6577** (24 hours). A live operator will register you at the **discounted** price.

AMPK

PROMOTES LONGEVITY FACTORS

Importance of AMPK

Studies show **increased** AMPK activity supports:

- Reduced fat storage,¹
- New mitochondria production,²
- The promotion of healthy blood glucose and lipids already within normal range.³

AMPK Activator provides nutrients shown to significantly **boost** AMPK activity.

Gynostemma Pentaphyllum

In one study, researchers documented a 1-inch reduction in **abdominal circumference** in overweight individuals who took **450 mg** daily of *G. pentaphyllum* extract for 12 weeks.⁴

Trans-Tiliroside

Trans-tiliroside promotes healthy blood glucose levels and body weight already within normal range.⁵

References

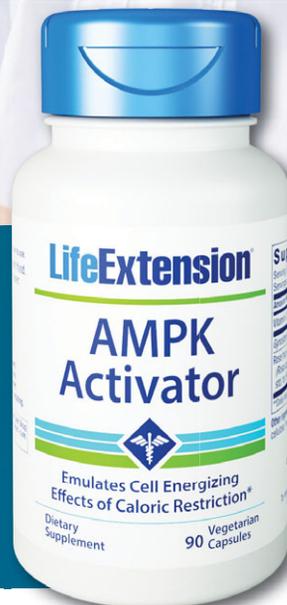
1. *Age (Dordr)*. 2014 Apr;36(2):641-63.
2. *Clin Sci (Lond)*. 2013 Apr;124(8):491-507.
3. *J Mol Med (Berl)*. 2011 Jul;89(7):667-76.
4. *Biotechnol Lett*. 2012 Sep;34(9):1607-16.
5. *Obesity (Silver Spring)*. 2014 Jan;22(1):63-71.

Non-GMO
ActivAMP® is a registered trademark of Gencor.

AMPK Activator

Item #01907 • 90 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$48	\$36
4 bottles		\$33 each



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

This supplement should be taken in conjunction with a healthy diet and regular exercise program. Individual results are not guaranteed and results may vary.

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



PREMIER

Giving more value to those we value most.

You already get 2% LE Dollars back on every purchase as a part of Your Healthy Rewards. But Premier gives you so much more:



12 months of FREE unlimited shipping. Anywhere in the United States ... including Alaska and Hawaii.*



DOUBLE rewards. That's 4% LE Dollars back on every purchase — twice as much as regular customers.†



Instant \$50 Dollar Bonus. This bonus offsets the cost of Premier, giving more value to those we value most.

Premier. Simply. More.

Just \$49.95 | \$59.95 for international customers

Call **1-888-224-8239** to enroll

Visit www.LifeExtension.com/Premier for details

Mention code **YRX605A**

LifeExtension



*FREE unlimited standard delivery (3 to 5 business days) to any mailing address within the 50 U.S. states, excluding U.S. territories. Also includes discounts on non-standard shipping and shipping outside of the U.S. Excludes blood test products and gift cards.

†Earn LE Dollars on all Life Extension purchases (except shipping fees, Life Extension Magazine® subscriptions, CHOICE and Premier program fees, and purchases made with LE Dollars or gift cards). Redeem LE Dollars to purchase products, blood tests, sale items, and shipping fees at the rate of 1 LE Dollar equal to \$1 U.S. Dollar at checkout. LE Dollars cannot be redeemed for CHOICE and Premier program fees or to purchase gift cards or Life Extension Magazine® subscriptions. LE Dollars have no cash value and are not redeemable for cash, transferable or assignable for any reason.

Prices subject to change without notice. Cannot be combined with any other offer. Offer not available to international customers serviced by distributors of Life Extension products.



Fight Stress and *Support* Adrenals with Adaptogens

It would be impossible to overstate the devastating impact chronic stress has on Americans' health. As many as **75%** to **90%** of primary care physician visits may be related to acute or chronic stress.¹

Chronic stress can be harmful to our emotional and physical well-being. It increases the risk for cardiovascular disease, diminished immune function, impaired cognitive function, and depression—and evidence shows that it contributes directly to ***accelerated aging*** and ***premature death***.¹⁻⁴

We now recognize that chronic stress leads to a body-wide imbalance of critical hormones and other essential signaling molecules. While eliminating stress is not practical in today's society, it *is* possible to restore the body's natural hormonal balance and reduce the impact of stress on cells, tissues, and organs throughout the body.¹

Adaptogens are nutrients that have been used for thousands of years to help counteract the negative effects of stress by restoring the body's natural balance.

Modern research has now confirmed that specific **adaptogens** can help the body fight the harmful biochemical and emotional effects of stress. The result is improved mood, greater energy, reduced incidence of chronic disease, and increased longevity.^{5,6}

In this article, we'll discuss the four adaptogens best-suited to help naturally neutralize the effects of **chronic stress**.

How Chronic Stress Affects the Body

The human body is well-equipped to respond to short-term, acute stress. It has a built-in alarm system that, when triggered, releases a surge of hormones that are designed to help ensure survival.

But chronic, unrelenting stress can wreak havoc on the body. Chronic stress leads to higher risk for cardiovascular disease and diminished immune function. It also causes harmful shifts in the human gut microbiome, the community of trillions of bacterial cells in our intestinal tracts that are intimately involved not only with intestinal health but also with normal balance and metabolism throughout the body.^{1,7,8}

Studies show that psychological stress is the **strongest** risk factor that predicts future cardiac events—including heart attacks—in people with existing coronary artery disease.⁹ And it leads to serious impairment of immune function, leaving us at major risk for cancers or severe viral infections.^{1,10,11}

There is also evidence that chronic stress directly contributes to accelerated aging and premature death.²⁻⁴

One recent alarming study showed that stress is associated with shortened **telomeres**. As most of you know, **telomeres** are stretches of DNA at the end of chromosomes that shorten with aging and can contribute to cellular senescence, or cell aging.⁴

One of the main reasons why chronic stress is so harmful is because it causes fundamental shifts in the way our cells maintain their balanced function, called **homeostasis**.¹² This imbalance makes us feel edgy, tired, or depressed. It has even been associated with obesity, diabetes, osteoporosis, hypertension, cardiovascular disease, infectious disease, cancer, gastrointestinal complaints, neurological disorders, sexual dysfunction, psychological problems, suppressed immunity, and more.^{4,5,12-15}



How to Protect Against Chronic Stress

It is nearly impossible to eliminate stress from our lives, which makes it important to supply our bodies with the tools necessary to protect against the long list of consequences stress can produce.

Substances called **adaptogens** have been used for hundreds—and some, even thousands—of years to help restore balance to the body, counteracting the harmful effects of chronic stress.

While adaptogens are not primary stress relievers, what they do is help to naturally fight the harmful effects stress has on the body. Studies show this leads to improved mental and physical performance, reduced incidence of chronic disease, and increased longevity.^{5,6}

We've included in this article the latest information on nutrients with adaptogenic properties best suited for rebalancing the body's stress responses.

Let's look at each of them individually.



Ashwagandha Improves Strength and Endurance

Ashwagandha (*Withania somnifera*) has been used for millennia as an adaptogenic herb in the Indian Ayurvedic system of medicine.¹⁶ In that system, it is considered a *Rasayana*, or tonic, that can improve endurance and stamina, promote longevity, improve immunity, and enhance fertility.^{16,17}

In animal studies, ashwagandha root extract has been found to improve measures of immune function and increase endurance. It has also been shown to prevent stress-induced gastric ulcers, reduce the inflammatory response to foreign proteins, and improve body weight lost as a result of stress.¹⁶

Recent studies in humans have confirmed ashwagandha's adaptogenic ability to improve the body's response to chronic stress—particularly in the area of physical activity.

Moderate exercise has multiple health benefits, but it also induces injury from muscle cell damage and inflammation. There is an important role, therefore, for adaptogens such as ashwagandha in mitigating exercise-induced stress.

Villagers living high in the Himalayas have used adaptogens for centuries to enhance their resistance to the effects of oxygen deprivation and to boost their endurance from strenuous tasks.¹⁸ Now, modern research offers solid evidence that ashwagandha helps improve the body's response to stressful exercise and resistance training.

In one study, healthy athletic adults taking ashwagandha (**300 mg** twice daily) experienced a **2-fold** increase in their oxygen consumption after 12 weeks, compared with placebo recipients.¹⁷ Increased oxygen consumption is an indicator of improved cardiorespiratory endurance. The supplemented subjects also experienced significant improvement in quality of life scores, compared with the placebo group.

A second study confirmed and extended these findings.¹⁹ Subjects taking ashwagandha root extract (**300 mg** twice daily) showed remarkable improvements in muscle strength compared to placebo recipients. After just eight weeks of training:

- Supplemented men bench-pressed **101.2 pounds**, compared to only **58 pounds** in the placebo group,
- On leg-extension exercises, supplemented men could lift **31.9 pounds**, compared to **21.6 pounds** in the placebo group, and
- Muscle size grew significantly more in the supplemented group than in the placebo group.

Most telling of ashwagandha's adaptogenic properties were the biochemical findings. Compared to the placebo recipients, men taking ashwagandha experienced greater reduction of exercise-induced muscle damage, greater decrease in body fat percentage, and a whopping **4.3-fold** increase in testosterone levels.

These studies are a powerful demonstration of ashwagandha's ability to relieve the physiological effects of physical stress.



What You Need to Know

Adaptogens Combat Effects of Chronic Stress

- Chronic stress can underlie many of the diseases we associate with aging.
- Chronic stress can lead to imbalances in metabolic processes throughout the body that can accelerate aging.
- Adaptogens can help the body rebalance its hormones and neurotransmitters, resulting in significant protection against stress.
- Ashwagandha, bacopa, cordyceps, and holy basil each work by different mechanisms to help the body adapt and protect itself against chronic stress, balancing cortisol levels and supporting adrenal gland function.
- People under chronic stress should consider supplementing with these nutrients as a means of restoring a healthy balance in their bodies and protecting themselves against the deleterious effects of stress.

Bacopa Improves Cognitive Performance

Bacopa monnieri (also called brahmi, or water hyssop) is an herb native to South Asia. It has a lengthy history in Ayurvedic medicine for use in the treatment of **cognitive impairment** and **nervousness**, both of which are closely related to stress.²⁰⁻²² Early studies have validated its adaptogenic effects in animal models, and recent human studies have demonstrated its practical impact in those suffering from chronic stress.

For example, one study showed that when rats were subjected to either acute or chronic stress they experienced an imbalance in their blood cortisol levels and in the levels of “fight-or-flight” neurotransmitters in their brains. Treatment with bacopa extracts helped to **normalize** these stress-induced changes, confirming its **adaptogenic** ability. Bacopa has also been shown to help protect against the depressive effects of chronic stress, help prevent the brain degeneration that arises from chronic stress,²³ and help fight the harmful effects of oxidative stress that can be caused by emotional and physical stresses.²²

Impressive animal studies like these inspired human research into bacopa as a valuable stress-protective adaptogen—and the results did not disappoint. One study showed that bacopa (**125 mg** twice daily) produced significant improvements in mental control, logical memory, and learning.²¹

Another study showed that bacopa can provide protective effects against **multitasking**, an inherently stressful activity that can cause deterioration in cognitive performance. In this human study, bacopa extracts improved both cognitive performance and mood in as little as one to two hours.²⁴

Together, these results demonstrate that bacopa has powerful adaptogenic effects, both in its ability to regulate cortisol levels during stress and in its ability to improve cognitive performance in those facing chronic stress.

Cordyceps Improves Immune Function

Cordyceps is a versatile medicinal mushroom that has been used as an adaptogen and health promoter for centuries. It is notable for its ability to battle oxidative stress and inflammation, and extracts of the **cordyceps fungi** have been shown to help enhance immune function in animals facing stress.²⁵⁻²⁹

This is critical, since **diminished immune function** is one of the primary consequences of chronic stress, and has been shown to leave us at major risk for both cancer and severe viral infection.^{1,10,11}

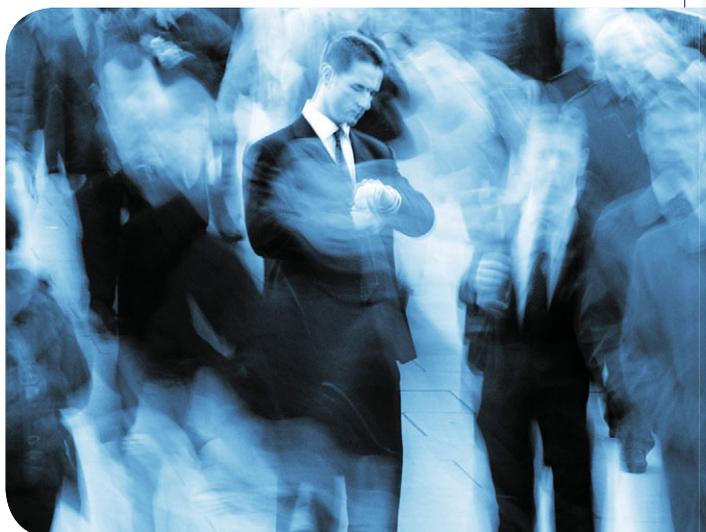
Cordyceps’s impact on other types of physiological stress is no less dramatic. For example, in mice subjected to a forced swimming test, cordyceps polysaccharide treatment significantly prolonged swimming time (a measure of endurance), increased natural protective systems in blood and tissue, and lowered markers of oxidized cells.²⁹

Another common effect of chronic stress is depression. **Cordycepin**, the major bioactive component of cordyceps, has been shown to have antidepressant-like effects in mice subjected to chronic mild stress.³⁰ Prior to treatment with cordycepin, the chronic stress induced depression-like symptoms, reduced levels of **brain-derived neurotrophic factor (BDNF)**, a growth factor for brain cells, and decreased serotonin and dopamine in the brain. But when animals received a cordycepin supplement, their behaviors **normalized**, as did all of the imbalanced neurotransmitters and elevated markers of stress. The low levels of BDNF also rose significantly, especially in the hippocampus, the brain’s memory/cognition center.

What About Adrenal Fatigue?

Although not a diagnosis recognized by the conventional medical establishment, some innovative doctors characterize “adrenal fatigue” as a condition that shares some symptoms with Addison’s disease, such as tiredness, depression, muscle pain, poor concentration, low blood sugar, craving for stimulants, and difficulty sleeping. However, in adrenal fatigue it is thought that the adrenal glands are unable to perform, normally due to exposure to chronic stress.³⁹

Life Extension® believes that symptoms often attributed to “adrenal fatigue” arise from multifactorial pathological processes involving, among other systems, the **hypothalamic-pituitary-adrenal (HPA)** axis, and that these conditions must be treated as such.



These results demonstrate that cordyceps extracts have adaptogenic effects with a focus on improved immune function, an essential feature for any supplement aimed at protecting the body from the long-term effects of stress.

Holy Basil Protects Against Stress-Induced Anxiety and Depression

Holy basil (*Ocimum sanctum*), also known as *tulsi*, is an aromatic plant that has been used for more than 3,000 years in Ayurvedic medicine. It is considered by many experts to be the preeminent adaptogen of Ayurvedic medicine.³¹ Now, modern science has confirmed holy basil's adaptogenic ability to protect the body against the effects of chronic stress.

The leaf extract of the plant contains several novel compounds that have been shown to normalize high blood sugar and cortisol levels while reducing markers of stress-induced tissue breakdown and excessive adrenal gland size.³² It also provides strong protection against oxidative and radiation stress on cells, which can affect many aspects of human health.^{33,34}

Importantly, animal studies show that holy basil helps the body maintain **homeostasis** of the stress hormone **cortisol**.³⁵ It rebalances mood-associated neurotransmitters in the brain—specifically ones that can produce feelings of hopelessness, which is a key component of stress-induced depression.³⁶

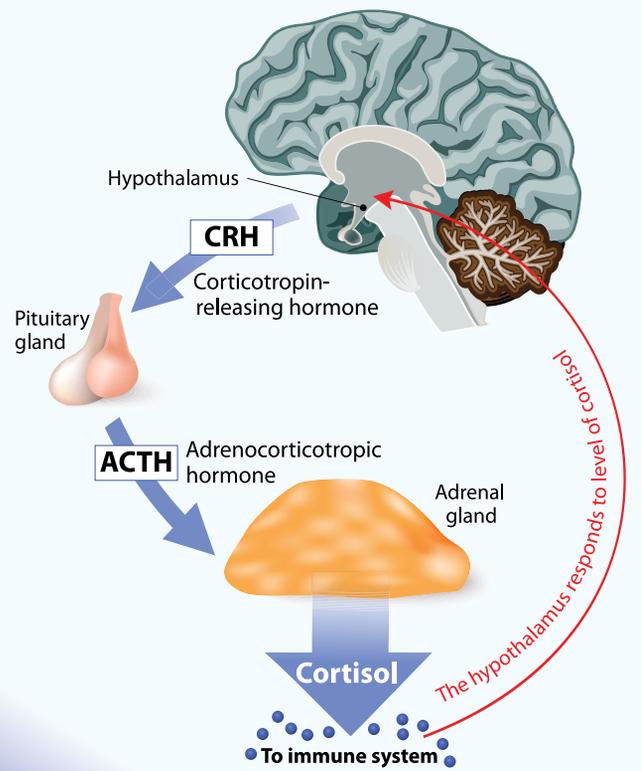
Human studies show that holy basil protects against the psychological effects of stress. In one study, holy basil extract (**500 mg** twice daily) significantly reduced the severity of generalized anxiety disorder, stress, and depression in middle-aged adults. It also improved the “willingness to adjust,” an important psychological characteristic that helps people adapt to circumstances beyond their control.³⁷

Taking this adaptogenic impact of holy basil one step further, in a human study, researchers demonstrated that **300 mg** a day of holy basil leaf extract produced a significant improvement in many different cognitive parameters, including reaction time and error rates on standardized tests, as well as on a test that measures anxiety.³⁸

These findings together highlight the importance of holy basil as an adaptogen capable of controlling the anxiety and depressive effects of chronic stress.

Summary

Chronic stress is an important factor in Americans' lives today. In addition to the unpleasant feelings and mood that it can cause, chronic stress also underlies many of the disorders associated with aging.



Human Stress Response System

Release of cortisol stimulated by hormones from the hypothalamus and pituitary gland in response to stress. Under normal conditions, this system keeps homeostasis in check, but under stressful conditions, it is constantly activated.

The impact of chronic stress on the adrenal gland can be enormous, causing the gland to pump out constant, high levels of stress-adaptive hormones. Over prolonged periods, these hormones, particularly cortisol, produce undesirable consequences—including weight gain, diabetes, and immunosuppression—which can become contributors to further health problems on their own.

We can now help protect our bodies against the impact of chronic stress by using a group of adaptogenic nutritional supplements. Ashwagandha, bacopa, cordyceps, and holy basil have each been in use for hundreds to thousands of years as powerful adaptogens that influence the brain and its stress responses. Each acts by a different mechanism, and each has now been shown to significantly attenuate the effects of chronic stress, resulting in improved mood, greater energy, and better adaptability to the stressful circumstances of life.

Anyone living with the effects of chronic stress should consider supplementing with these proven, stress-relieving adaptogens in order to protect their hormonal status and adrenal glands. ●

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

References

- Head KA, Kelly GS. Nutrients and botanicals for treatment of stress: adrenal fatigue, neurotransmitter imbalance, anxiety, and restless sleep. *Altern Med Rev*. 2009;14(2):114-40.
- Nielsen NR, Kristensen TS, Schnohr P, et al. Perceived stress and cause-specific mortality among men and women: results from a prospective cohort study. *Am J Epidemiol*. 2008;168(5):481-91; discussion 92-6.
- Carroll BJ. Ageing, stress and the brain. *Novartis Found Symp*. 2002;242:26-36; discussion -45.
- Wikgren M, Maripuu M, Karlsson T, et al. Short telomeres in depression and the general population are associated with a hypo-cortisolemic state. *Biol Psychiatry*. 2012;71(4):294-300.
- Epel ES. Psychological and metabolic stress: a recipe for accelerated cellular aging? *Hormones (Athens)*. 2009;8(1):7-22.
- Panosian A, Wikman G. Evidence-based efficacy of adaptogens in fatigue, and molecular mechanisms related to their stress-protective activity. *Curr Clin Pharmacol*. 2009;4(3):198-219.
- Huis in 't Veld JH. Gastrointestinal flora and health in man and animal. *Tijdschr Diergeneeskd*. 1991;116(5):232-9.
- Lizko NN, Silov VM, Syrych GD. Events in the development of dysbacteriosis of the intestines in man under extreme conditions. *Nahrung*. 1984;28(6-7):599-605.
- Allison TG, Williams DE, Miller TD, et al. Medical and economic costs of psychologic distress in patients with coronary artery disease. *Mayo Clin Proc*. 1995;70(8):734-42.
- Andersen BL, Farrar WB, Golden-Kreutz D, et al. Stress and immune responses after surgical treatment for regional breast cancer. *J Natl Cancer Inst*. 1998;90(1):30-6.
- Pike JL, Smith TL, Hauger RL, et al. Chronic life stress alters sympathetic, neuroendocrine, and immune responsiveness to an acute psychological stressor in humans. *Psychosom Med*. 1997;59(4):447-57.
- Kyrou I, Tsigos C. Stress mechanisms and metabolic complications. *Horm Metab Res*. 2007;39(6):430-8.
- Schneiderman N, Ironson G, Siegel SD. Stress and health: psychological, behavioral, and biological determinants. *Annu Rev Clin Psychol*. 2005;1:607-28.
- Kyrou I, Tsigos C. Chronic stress, visceral obesity and gonadal dysfunction. *Hormones (Athens)*. 2008;7(4):287-93.
- Azuma K, Adachi Y, Hayashi H, et al. Chronic psychological stress as a risk factor of osteoporosis. *J uoeh*. 2015;37(4):245-53.
- Singh N, Bhalla M, de Jager P, et al. An overview on ashwagandha: a rasayana (rejuvenator) of Ayurveda. *Afr J Tradit Complement Altern Med*. 2011;8(5 Suppl):208-13.
- Choudhary B, Shetty A, Langade DG. Efficacy of ashwagandha (*Withania somnifera* [L.] Dunal) in improving cardiorespiratory endurance in healthy athletic adults. *Ayu*. 2015;36(1):63-8.
- Zhang ZH, Feng SH, Hu GD, et al. Effect of *Rhodiola kirilowii* (Regel.) Maxim on preventing high altitude reactions. A comparison of cardiopulmonary function in villagers at various altitudes. *Zhongguo Zhong Yao Za Zhi*. 1989;14(11):687-90, 704.
- Wankhede S, Langade D, Joshi K, et al. Examining the effect of *Withania somnifera* supplementation on muscle strength and recovery: a randomized controlled trial. *J Int Soc Sports Nutr*. 2015;12:43.
- Kumar SS, Saraswathi P, Vijayaraghavan R. Effect of bacopa monniera on cold stress induced neurodegeneration in hippocampus of wistar rats: a histomorphometric study. *J Clin Diagn Res*. 2015;9(1):Af05-7.
- Raghav S, Singh H, Dalal PK, et al. Randomized controlled trial of standardized Bacopa monniera extract in age-associated memory impairment. *Indian J Psychiatry*. 2006;48(4):238-42.
- Dhanasekaran M, Tharakan B, Holcomb LA, et al. Neuroprotective mechanisms of ayurvedic antidementia botanical Bacopa monniera. *Phytother Res*. 2007;21(10):965-9.
- Chowdhuri DK, Parmar D, Kakkar P, et al. Antistress effects of bacosides of Bacopa monnieri: modulation of Hsp70 expression, superoxide dismutase and cytochrome P450 activity in rat brain. *Phytother Res*. 2002;16(7):639-45.
- Benson S, Downey LA, Stough C, et al. An acute, double-blind, placebo-controlled cross-over study of 320 mg and 640 mg doses of Bacopa monnieri (CDRI 08) on multitasking stress reactivity and mood. *Phytother Res*. 2014;28(4):551-9.
- Zhu JS, Halpern GM, Jones K. The scientific rediscovery of an ancient Chinese herbal medicine: cordyceps sinensis: part I. *J Altern Complement Med*. 1998;4(3):289-303.
- Li SP, Li P, Dong TT, et al. Anti-oxidation activity of different types of natural cordyceps sinensis and cultured cordyceps mycelia. *Phytomedicine*. 2001;8(3):207-12.
- Ng TB, Wang HX. Pharmacological actions of cordyceps, a prized folk medicine. *J Pharm Pharmacol*. 2005;57(12):1509-19.
- Liu JY, Feng CP, Li X, et al. Immunomodulatory and antioxidative activity of cordyceps militaris polysaccharides in mice. *Int J Biol Macromol*. 2016;86:594-8.
- Yan F, Wang B, Zhang Y. Polysaccharides from cordyceps sinensis mycelium ameliorate exhaustive swimming exercise-induced oxidative stress. *Pharm Biol*. 2014;52(2):157-61.
- Tianzhu Z, Shihai Y, Juan D. Antidepressant-like effects of cordycepin in a mice model of chronic unpredictable mild stress. *Evid Based Complement Alternat Med*. 2014;2014:438506.
- Cohen MM. Tulsi - *Ocimum sanctum*: A herb for all reasons. *J Ayurveda Integr Med*. 2014;5(4):251-9.
- Gupta P, Yadav DK, Siripurapu KB, et al. Constituents of *Ocimum sanctum* with antistress activity. *J Nat Prod*. 2007;70(9):1410-6.
- Rai V, Iyer U, Mani UV. Effect of Tulasi (*Ocimum sanctum*) leaf powder supplementation on blood sugar levels, serum lipids and tissue lipids in diabetic rats. *Plant Foods Hum Nutr*. 1997;50(1):9-16.
- Subramanian M, Chintalwar GJ, Chattopadhyay S. Antioxidant and radioprotective properties of an *Ocimum sanctum* polysaccharide. *Redox Rep*. 2005;10(5):257-64.
- Jothie Richard E, Illuri R, Bethapudi B, et al. Anti-stress activity of *Ocimum sanctum*: Possible effects on hypothalamic-pituitary-adrenal axis. *Phytother Res*. 2016;30(5):805-14.
- Ahmad A, Rasheed N, Chand K, et al. Restraint stress-induced central monoaminergic & oxidative changes in rats & their prevention by novel *Ocimum sanctum* compounds. *Indian J Med Res*. 2012;135(4):548-54.
- Bhattacharyya D, Sur TK, Jana U, et al. Controlled programmed trial of *Ocimum sanctum* leaf on generalized anxiety disorders. *Nepal Med Coll J*. 2008;10(3):176-9.
- Sampath S, Mahapatra SC, Padhi MM, et al. Holy basil (*Ocimum sanctum* Linn.) leaf extract enhances specific cognitive parameters in healthy adult volunteers: A placebo controlled study. *Indian J Physiol Pharmacol*. 2015;59(1):69-77.
- Ahn YW. Adrenal exhaustion and fatigue due to chronic stress. *J Korean Med Assoc*. 2011;54(1):81-7.





Blueberry Extract

Boosts DNA Function

Studies show that blueberries delay the aging process through a variety of mechanisms, including **maintaining healthy DNA structure** and favorably modulating **genes** associated with aging.^{1,2}

Chockfull of **anthocyanins**, the **blueberry** provides health-boosting benefits shown to:

- Enhance heart health³
- Maintain brain function^{4,5}
- Sustain healthy blood sugar levels already within normal range⁶
- Support smooth, firm skin⁷
- Improve movement and coordination⁸

Blueberry extract is more potent than the whole berry, providing greater metabolic support throughout the body and without the excess sugar of raw fruit.⁹

Blueberry Extract Capsules consist of concentrated **extracts** from **wild blueberries**, which possess up to **10 times** the antioxidant capacity of cultivated berries.

Blueberry Extract Capsules

Item #01214 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$22.50	\$16.88
4 bottles		\$15 each



Suggested dose is one capsule daily for most individuals.

References

1. *Curr Pharm Des.* 2013;19(34):6094-111.
2. *Int J Mol Sci.* 2013;14(11):21447-62.
3. *PLoS One.* 2009;4(6):e5954.
4. *Nutr Neurosci.* 2005 Apr;8(2):111-20.
5. Available at: <http://www.scientificamerican.com/article.cfm?id=your-brain-on-blueberries>. Accessed January 28, 2015.
6. *Georgian Med News.* 2006 Dec;(141):66-72.
7. *J Cosmet Dermatol.* 2009 Jun;8(2):147-51.
8. *Applied Physiol Nutr Met.* 2015;40(6):543-9.
9. *J Agric Food Chem.* 2010 Apr 14;58(7):3970-6.

Non-GMO

AuroraBlue® is a registered trademark of Denali BioTechnologies, Inc.

To order **Blueberry Extract Capsules**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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



IMPROVED FORMULA! LOW PRICE!



Turn Back the **CLOCK** and **FEEL** **ENERGIZED**

Aging robs us of the energy of our youth. Age-related changes in cellular energy production occurs in our mitochondria, the critical energy power plant of our cells.

Mitochondrial Energy Optimizer with BioPQQ® provides the essential nutrients to help neutralize these changes.

Mitochondrial Energy Optimizer with BioPQQ®

Item #01868 • 120 capsules

	Retail Price	Your Price
1 bottle	\$72	\$54
4 bottles		\$48 each



To order **Mitochondrial Energy Optimizer with BioPQQ®**, call **1-800-544-4440** or visit www.LifeExtension.com

Non-GMO. BioPQQ® is a registered trademark of MGC (Japan). Bio-Enhanced® is a registered trademark of Geronova Research, Inc.

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

Maintain Normal Balance During STRESS



Stress disrupts multiple biochemical pathways which can deplete the adrenal glands—making it difficult for the body to restore its natural equilibrium.

Life Extension® has created a state-of-the-art formulation of **four botanical adaptogens**, each shown to modulate different stress-related changes.

Adrenal Energy Formula provides:

- **Ashwagandha**, supports muscle strength¹ and increases oxygen consumption,² an indicator of cardiorespiratory endurance,
- **Bacopa**, supports mental control, logical memory, and learning,³ and enhances cognitive performance and mood,⁴
- **Cordyceps**, promotes endurance⁵ and enhanced immune function,⁶ and
- **Holy basil**, supports balanced levels of neurotransmitters⁷ and blood sugar⁸ for those already within the normal range, and enhances mood.⁹

References

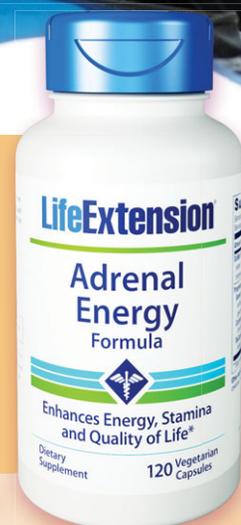
1. *J Int Soc Sports Nutr.* 2015;12:43.
2. *Ayu.* 2015 Jan-Mar;36(1):63-8.
3. *Indian J Psychiatry.* 2006 Oct;48(4):238-42.
4. *Phytother Res.* 2014 Apr;28(4):551-9.
5. *Pharm Biol.* 2014;52(2):157-61.
6. *Evid Based Complement Alternat Med.* 2014;2014:438506.
7. *Indian J Med Res.* 2012 Apr;135(4):548-54.
8. *J Nat Prod.* 2007 Sep;70(9):1410-6.
9. *Nepal Med Coll J.* 2008 Sep;10(3):176-9.

Adrenal Energy Formula

Item #01630 • 120 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$46	\$34.50
4 bottles		\$31.50 each

Non-GMO



BACOGNIZE® ULTRA is a registered trademark of Verdure Sciences, Inc. OciBest® is a registered trademark of Natural Remedies Private Limited. Sensoril® is protected under US Patent Nos. 6,153,198 and 6,713,092 and is a registered trademark of Natreon, Inc.

To order **Adrenal Energy Formula**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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



Sleep Medicine CONFERENCE

Why do people need sleep? Sleep allows the brain to restore essential nutrients (like glycogen) and remove waste products^{1,2} (like the beta-amyloid protein that causes Alzheimer's disease).^{3,4} Sleep replenishes proteins depleted during wakefulness.⁵ Sleep is important for memory consolidation.^{6,7}

Although the brain is approximately **2%** of the body's weight, it uses about **20%** of the body's energy.⁸ The major source of that energy is the molecule **adenosine triphosphate (ATP)**.

With increasing hours of wakefulness, there are decreasing amounts of ATP and increasing amounts of **adenosine** from which the high-energy phosphates have been removed. Adenosine causes sleepiness.⁹ Caffeine blocks adenosine action, reducing sleepiness and increasing alertness,¹⁰ while impeding learning and memory.¹¹

How much sleep do people need? The need for sleep diminishes with age. During the first six months of life, infants sleep more than half the time.¹² Medications account for many of the sleeping problems of the elderly,¹³ but healthy older adults (age 65-76) tolerate sleep deprivation better than young adults (age 18-29).¹⁴ More than half of sleep-related car accidents occur in drivers under age 30, compared to only **5%** in drivers over age 65.¹⁴ Adolescents tend to go to bed late and sleep late when they are allowed to, but as people get older they tend to go to bed earlier and awaken earlier.¹⁵

Independent of diabetes and high blood pressure, men who slept less than 6 hours per night were found to have **four times** the risk of dying compared to those who slept more (although this effect was not seen in women).¹⁶ Young healthy adults randomized to spend 4, 6, or 8 hours in bed for 14 nights showed reduced cognitive performance in proportion to lack of sleep for those with bed times of 6 hours or less.¹⁷

Healthy adults who habitually sleep more than the minimum necessary to avoid sleepiness and impaired function have shown greater development of the cerebral cortex.¹⁸ However, many studies have shown that excessive sleep is associated with increased risk of dying, although this is controversial.¹⁹ Post-menopausal women sleeping more than 9 hours per night exhibited increased inflammation and blood clotting, which are risk factors for cardiovascular disease.²⁰

One study showed that napping for 10 minutes improves alertness and cognitive performance over the subsequent 3 hours, whereas a 5-minute nap is of little benefit, and a 20-minute nap results in "sleep inertia," which is the period of confusion, sleepiness, and disorientation that is experienced upon awakening from sleep.²¹ A 16-week supplementation of healthy children with **600 mg** daily of the omega-3 fatty acid DHA improved their sleep (longer sleep with fewer awakenings).²²

On June 6-10, 2015, I attended the annual meeting of the American Academy of Sleep Medicine and the Sleep Research Society in Seattle, Washington.

Effects of Sleep Loss

David Dinges, PhD, (Chief of the Division of Sleep and Chronobiology, Perelman School of Medicine, Philadelphia, Pennsylvania) studies the effects of sleep



loss. Between 1985 and 2012, average nightly sleep time reported by American adults declined from 7.40 to 7.18 hours.²³ The reported prevalence of insomnia among US adults increased from **17.5%** in 2004 to **19.2%** in 2012. The largest relative increases occurred among diabetics and those in the 25-34 age group.²⁴ An estimated **15%** of Americans sleep less than 6 hours per night.²⁵ Dr. Dinges has demonstrated that healthy adults restricted to four hours sleep per night for five nights ate more food and gained weight.²⁶ Sleep restriction has been shown to reduce blood levels of the satiety hormone **leptin**,²⁷ and to increase the hunger-producing hormone **ghrelin**,²⁸ thereby increasing appetite. Dr. Dinges also showed that an extended night of sleep following the five nights of restricted sleep was not adequate to restore alertness or prevent sleepiness.²⁹

Rachel Leproult, PhD, (Researcher, Universite libre de Bruxelles, Brussels, Belgium) studies the relationship between diabetes and quantity or quality of sleep.³⁰



In one of her studies, she found that persons who slept less than 5 or 6 hours per night had a **28%** greater incidence of diabetes, those who had difficulty initiating sleep had a **57%** greater incidence, and those who had difficulty maintaining sleep had an **84%** greater incidence.³¹ Dr. Leproult conducted an experiment in which healthy adults were either simply restricted to less than 5 hours sleep per night for eight days, or restricted to less than 5 hours sleep combined with disruption of the 24-hour circadian rhythm cycle. Both groups showed increased inflammation and insulin resistance, but the effect was substantially greater when sleep restriction was combined with circadian disruption. Insulin sensitivity dropped **39%** for those who were only sleep restricted, but more than **53%** for those who also experienced circadian disruption.³²

Circadian disruption due to flying across more than three time zones ("jet lag") is most effectively treated by exposure to bright daylight in the morning, and administration of melatonin in the evening.³³ In another experiment, Dr. Leproult asked healthy adults who normally sleep less than 7 hours per night to increase their time in bed by one hour per day. After 40 days the subjects showed substantially improved insulin sensitivity.³⁴



Carol Everson, PhD, (Professor, Medical College of Wisconsin, Milwaukee, Wisconsin) has studied the effects of sleep deprivation in rats. She has shown



that sleep deprivation weakens the immune system³⁵ and increases pathogenic bacteria in rats.³⁶ People sleeping less than 7 hours per night have been shown to be nearly **3 times** more likely to develop a common cold than people who sleep 8 hours or more.³⁷ Totally sleep-deprived rats will die within about two or four weeks, with over **20%** weight loss.³⁸ Dr. Everson subjected rats to 10 days of sleep restriction six times, separated by two days of rest each time. The rats lost **15%** of their body weight despite eating more than the usual amount of food. Even after four months of recovery, the rats were consuming **20%** more food and **35%** more water.³⁸ In another experiment, rats totally deprived of sleep for 10 days showed a **39%** increase in oxidative DNA damage, particularly in the liver, where this damage was almost **2.5 times** greater than in controls.³⁹

Cognitive Effects of Sleep Deprivation

Hans van Dongen, PhD, (Research Professor, Washington State University, Spokane, Washington) has conducted experiments to determine the effects



of sleep deprivation on cognitive performance.⁴⁰ He has found that sleep deprivation hampers decision making under conditions of uncertainty and unexpected change,⁴¹ but that the hampered decisions were more due to reduced reaction time than to reduced judgement.⁴² He has also found evidence that an increase

in the inflammatory cytokine TNF-alpha contributes to the impaired vigilance associated with sleep deprivation.⁴³ Other studies have shown not only that inflammatory cytokines increase after sleep loss,⁴⁴ but that cytokines (including TNF-alpha) released during illness increase sleepiness (to induce recovery-promoting sleep).⁴⁵

Objective Measures of Insomnia

Julio Fernandez-Mendoza, PhD, (Assistant Professor, Pennsylvania State University, Hershey, Pennsylvania) seeks objective measures of insomnia. He cited studies



that **8%-10%** of people suffer from chronic insomnia and that **20%-30%** of people will be suffering from temporary insomnia at any given time.⁴⁶ He notes that insomnia patients have higher levels of adrenal-like compounds (*catecholamines*), higher oxygen consumption, and higher blood pressure than good sleepers.⁴⁶ He has also found elevated stress hormone

(cortisol) in the saliva of children aged 5-12 reported by parents to have insomnia.⁴⁷ He has found that patients with severe insomnia were several times more likely to also be suffering from depression.⁴⁸ And Dr. Fernandez-Mendoza reported that excessive daytime sleepiness (affecting up to **30%** of people) is associated with obesity and weight gain.⁴⁹

Kai Spiegelhalter, MD, PhD, (Insomnia researcher, Freiberg Institute for Advanced Studies, Freiberg, Germany) also has sought objective measures of insomnia.



Chronic insomnia is generally defined based on subjective complaints of difficulty falling asleep or difficulty remaining asleep, along with daytime sleepiness and fatigue. For diagnosis of chronic insomnia, these complaints must persist for at least four weeks.⁵⁰ Overall, women complain of insomnia more than men, with the difference rising after age 45.⁵¹

Electronic monitoring of brain, heart, skeletal muscle, and eye movements (polysomnography) shows less difference between good sleepers and those complaining of insomnia than the complaints would indicate.⁵⁰ Nonetheless, there is evidence that those complaining of insomnia have higher heart rates, elevated cortisol (stress hormone), and greater brain glucose metabolism (indicating greater brain activity) during sleep.⁵² Inherited insomnia is associated with increased cortisol and decreased melatonin secretion.⁵¹ These quantitative indications of arousal in insomniacs form the

basis of Dr. Spiegelhalter's "hyper-arousal model of insomnia."^{50,53} Dr. Spiegelhalter has shown that patients complaining of insomnia put much greater effort into trying to fall asleep, which worsens the problem.⁵⁴ He does not recommend benzodiazepine tranquilizers such as **Librium®** or **Valium®** for anything but short-term use because of declining effectiveness, increasing dependency, and memory impairment.⁵⁵ Antihistamines and antidepressants are less addictive, but can have worse side effects, including liver and heart damage. Alcohol and barbiturates reduce the amount of time to fall asleep, but also reduce sleep quality.^{56,57} Prolonged-release melatonin supplementation has been shown to reduce insomnia in elderly patients without causing side effects.^{58,59} The main treatment Dr. Spiegelhalter recommends is Cognitive Behavioral Therapy for Insomnia (CBT-I).⁵¹

Cognitive Behavioral Therapy for Insomnia

Although Cognitive Behavioral Therapy for Insomnia (**CBT-I**) is often recommended by insomnia specialists as being the most safe and effective treatment, Simon



Kyle, PhD, (Lecturer, University of Manchester, Manchester, England) has found that the therapy is not standardized, is poorly documented when used, and often fails to work.⁶⁰ CBT-I can involve sleep diaries to identify thoughts or habits that cause sleep problems, relaxation training, using the bedroom only for sleep or



sex, avoiding caffeine, nicotine, or alcohol, leaving the bedroom if sleep does not occur within ten minutes, and sleep restriction therapy. Dr. Kyle has concentrated his attention on sleep restriction therapy. Interviewing patients, he was told that the suffering patients experienced when implementing sleep restriction subsided within a couple of weeks, and that the **craving** they developed for sleep led to sleep that was deeper and more efficient.⁶¹ Conducting objective measures of sleep restriction therapy, Dr. Kyle found that although sleep efficiency improved within four weeks, daytime attention and reaction speed deteriorated, not returning to starting values until after three months of the therapy.⁶²

Sleep Loss and Sex Hormones

Fiona Baker, PhD, (Senior Program Director, Human Sleep Research, SRI International, Menlo Park, California) studies sleep disturbances associated with menstruation and menopause. **Hot flashes** refer to episodes of sweating followed by chills.⁶³ Hot flashes are reported for **12.5%** of women just before menopause, **79.0%** during menopause, and **50.7%** following menopause. Chronic insomnia increases with increased severity of hot flashes.⁶⁴ Dr. Baker found that heart rate increased when hot flashes occurred.⁶⁵ Dr. Baker studied young women (average age 21) who did not have menstrual complaints, and found reduced sleep quality for three days before menstruation and for four days during menstruation.⁶⁶



Katherine Sharkey, MD, PhD, (Internist, Rhode Island Hospital, West Warwick, Rhode Island) studies the effects of aging and hormonal changes on sleep. In men between the ages of 64 and 74 the time by which nighttime sleep is reduced corresponded with the amount by which testosterone is reduced.⁶⁷ In women, menstrual irregularities begin at an average age of 46, lasting two to eight years.⁶⁸ Dr. Sharkey has shown that the greater incidence of waking during sleep is associated with higher levels of progesterone near the time of menstruation.⁶⁹



Sleep Disturbance with Shift Work

Karl Doghramji, MD, (Director, Sleep Disorder Center, Jefferson University Hospitals, Philadelphia, Pennsylvania) reported on the health consequences of shift work. Worldwide, about **20%** of the labor force works outside the hours of 7 A.M. to 6 P.M.⁷⁰ Shift workers suffer more obesity, cardiovascular disease, and cancer as well as experiencing nearly triple the incidence of occupational accidents as day workers.⁷⁰ Industrial productivity has been found to be about **5%** lower during night shifts.⁷¹ Caffeine and naps could reduce the accident rate and increase productivity.



A study in Japan found that prostate cancer among rotating-shift workers was triple normal, but that fixed-shift night workers had a regular prostate cancer incidence.⁷² The authors speculated that disruption of the circadian rhythm of melatonin secretion was responsible. A study of American nurses found that rotating-shift work was associated with a **4%** increase in ischemic stroke for every five years worked.⁷³ These authors also speculated that disruption of the circadian rhythm of melatonin secretion could be responsible.

A study of Iranian long-distance night-shift drivers found an increased incidence of metabolic syndrome.⁷⁴ In this case, the authors speculated that physical inactivity and roadside restaurant food was responsible. But healthy adults subjected to three weeks of combined sleep restriction (5-6 hours sleep per 24 hours) and circadian disruption (28 hour "days" in a laboratory setting) showed a **32%** decrease in insulin secretion after a standardized meal. Elevated blood glucose rose to prediabetic levels in some cases. The subjects showed an **8%** drop in resting metabolic rate, which would result in a weight gain of approximately **12.5** pounds per year.⁷⁵





Obstructive Sleep Apnea (OSA)

Ina Djonlagic, MD, (Instructor, Harvard Medical School, Boston, Massachusetts) has studied the effects of obstructive sleep apnea (OSA). Persons with obstructive sleep apnea suffer bouts of halted breathing (**apnea**) lasting 20 to 40 seconds several times per hour during sleep, resulting in disrupted sleep quality. Victims are often unaware of their condition, although they experience excessive daytime sleepiness.^{76,77}



Djonlagic

Obstructive sleep apnea victims typically have elevated blood pressure, and a four-fold increase in heartbeat irregularities in the atrium of the heart.⁷⁸ For people without obstructive sleep apnea, sudden cardiac death risk is minimal between midnight and 6 A.M., but for obstructive sleep apnea victims the risk of sudden cardiac death during that time is elevated more than two-and-a-half times.⁷⁹ Up to **95%** of patients with obstructive sleep apnea snore.⁷⁶ Obstructive sleep apnea is twice as common in men as it is in women.⁸⁰ Other risk factors include being overweight, being over age 40, and having an enlarged neck, tongue, or tonsils. Alcohol and other sedatives contribute to obstructive sleep apnea by relaxing throat muscles. Dr. Djonlagic has found that untreated obstructive sleep apnea caused cognitive deficits and increases the likelihood of becoming demented.⁸¹ Sleeping on one's back greatly increases the risk of obstructive sleep apnea, whereas sleeping on the side reduces it.⁸² Sleeping with an appliance that keeps the airways open with *continuous positive airway pressure* (CPAP) is the "gold standard" treatment for obstructive sleep apnea.⁸³ CPAP reduces the risk of cardiovascular death.⁸⁴ Dr. Djonlagic has found immediate benefits for attention, vigilance, and well-being after the first night of CPAP in obstructive sleep apnea patients.⁸⁵

Sleep Disorders in Soldiers

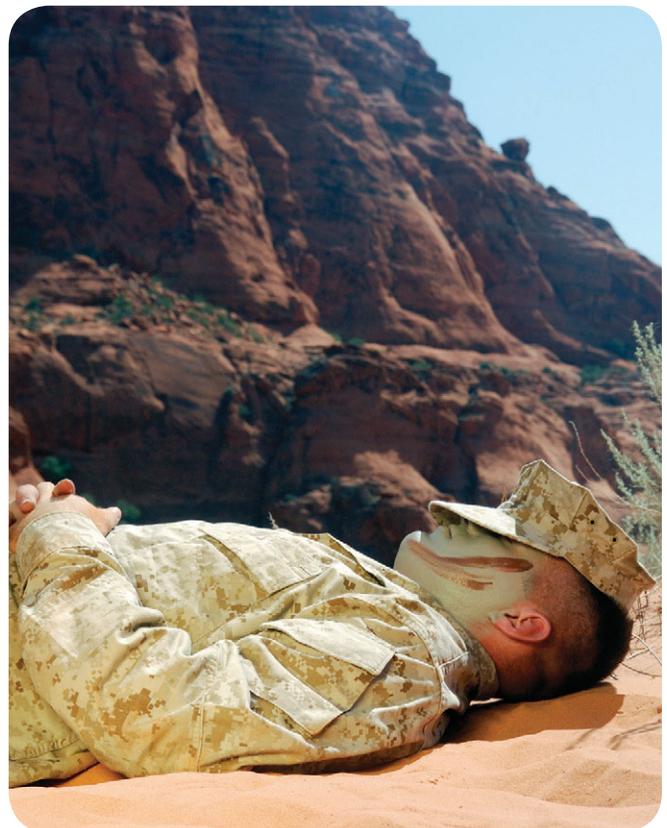
Vincent Mysliwec, MD, (Internist, Madigan Army Medical Center, Tacoma, Washington) has studied the sleep disturbances of soldiers returning from combat.

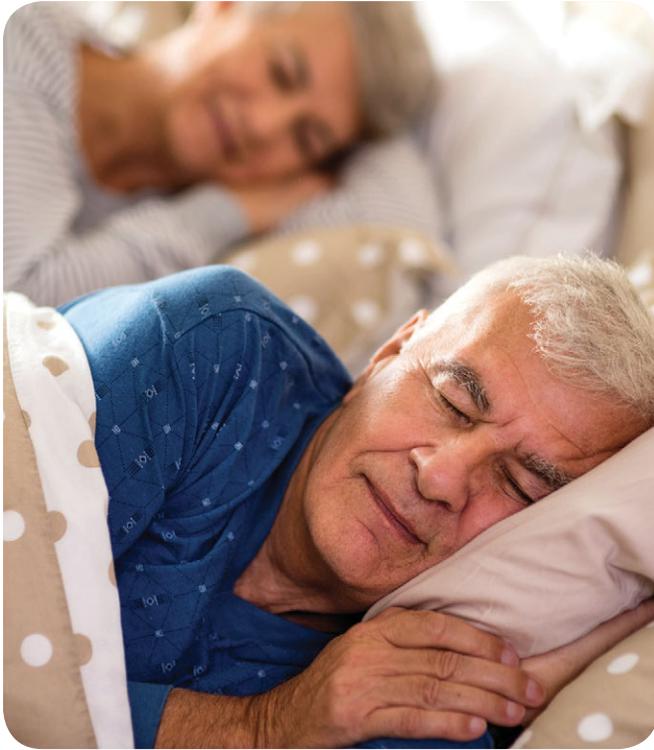


Mysliwec

Soldiers on deployment often sleep at odd hours, disrupting circadian rhythm.⁸⁶ According to one report, sleep time for soldiers averages 6.5 hours, but soldiers suffering from post-traumatic stress disorder (PTSD) sleep significantly less.⁸⁶ In the military, there is a stigma associated with mental health problems, which may account for the fact that soldiers are

about five times more likely to seek help for insomnia than for a mental health problem.⁸⁷ About one quarter of soldiers deployed to combat stations during the war on terror have experienced traumatic brain injury (TBI) from improvised explosive devices.⁸⁸ Compared to soldiers without traumatic brain injury, soldiers with a single traumatic brain injury had about four times the incidence of insomnia, whereas soldiers with multiple traumatic brain injuries had close to ten times the incidence of insomnia.⁸⁹ Soldiers with traumatic brain injury from blast injuries were more likely to develop anxiety and insomnia, whereas blunt trauma more often resulted in obstructive sleep apnea.⁹⁰





Age, Cognitive Ability, and Sleep

Anne Richards, MD, (Assistant Professor, University of California School of Medicine, San Francisco, California) used data from the online cognitive training program Lumosity to assess the relationship between age, cognitive ability, and reported sleep time. Other studies have indicated that consolidation of learning occurs during sleep for young adults, but not for healthy elderly adults,⁹¹ and that those who sleep between 6 and 9 hours per night do better on cognitive tests than those who sleep more or less.⁹² Dr.



Richards

Richards found that for young Lumosity participants, cognitive scores were highest for those reporting 7 hours of sleep per night. But, unexpectedly, for the oldest participants, those sleeping 7 hours per night had worse scores than those sleeping more or less.⁹³

Sleep and the Circadian Rhythm

Derk-Jan Dijk, PhD, (Professor of Sleep and Physiology, University of Surrey, Surrey, England) has been studying the effects of the day-night circadian rhythm on sleep and health.⁹⁴ The **circadian rhythm** is characterized by melatonin release in the evening (which promotes sleep) and cortisol release before awakening (which promotes arousal).¹⁵ Dr. Dijk has shown that the efficiency of sleep for replenishment of brain function



Dijk

for humans is highest when sleep times coincide with the 24-hour circadian rhythm, and is greatly reduced when circadian rhythm is disrupted by being in a sleep laboratory with an artificial 28-hour day.⁹⁵ He has also shown that the human “circadian clock” is disrupted by insufficient sleep.⁹⁶ Dr. Dijk has reported that the study of three geographically-separated modern hunter-gatherer communities without access to electric lighting showed an average sleep time of 7.7 hours, estimated to be about a half-hour longer than in modern industrial society.⁹⁷ Comparing three modern industrial countries (Singapore, Norway, and the United Kingdom) he found no seasonal variation in sleep time between the three countries, despite the fact that two of the countries are at high latitude and Singapore is near the equator (lacking much seasonal variation).⁹⁸

Concluding Remarks

With many sessions at the conference running in parallel, I may have missed presentations about two important sleep problems which I will now mention.

A study in Europe found that more than **5%** of the population suffered from **restless legs syndrome** (more often women than men), which causes insomnia.⁹⁹ This affliction is largely inherited, and is characterized by uncomfortable leg sensations when at rest, causing victims to move their legs to stop the discomfort.¹⁰⁰





The urge to urinate at night is a frequent source of sleep disturbance. Although roughly **10%** of those between age 20 and 40 urinate at least twice per night, nearly half of women and more than two-thirds of men over age 70 urinate at least twice per night.¹⁰¹ Treatment of benign prostatic hyperplasia (BPH) in men only has a mild benefit.¹⁰² Drug treatments include antidepressants, diuretics,¹⁰³ and drugs like *finasteride* that shrink enlarged prostate glands¹⁰⁴ as well as *tamsulosin* (**Flo-max**®) that facilitate more complete emptying of the bladder.¹⁰⁵ Patients are advised to avoid drinking too many liquids at night.¹⁰³

Concerning the conference, it seems that most people would be healthier if they would get more sleep. Apparently, modern life increasingly tempts people to stay awake longer so as to be more productive or more entertained. Melatonin and the omega-3 fatty acid DHA may help some people to sleep better, but clearly more is needed to address this widespread problem.

A major problem with sleep research is that it too often relies on people reporting on their sleep rather than objective data. Laboratory measures of sleep (polysomnography) are often at variance with subjective experience. And sleeping in a laboratory can disrupt sleep. Based on the presentations I heard, I could imagine many of the consequences of poor sleep: inflammation, liver DNA damage, blood pressure, or insulin resistance; blood cortisol or immune function; poor performance on standardized tests for cognition and reaction time. “Personalized medicine” has not yet arrived in sleep research because only averages of results for many people are used. The variation in sleep-need from one individual to another might be very large.¹⁰⁶ ●

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

References

1. Herculano-Houzel S. Sleep it out. *Science*. 2013;342(6156):316-7.
2. Xie L, Kang H, Xu Q, et al. Sleep drives metabolite clearance from the adult brain. *Science*. 2013;342(6156):373-7.
3. Lee H, Xie L, Yu M, et al. The effect of body posture on brain lymphatic transport. *J Neurosci*. 2015;35(31):11034-44.
4. Huang Y, Potter R, Sigurdson W, et al. Effects of age and amyloid deposition on Abeta dynamics in the human central nervous system. *Arch Neurol*. 2012;69(1):51-8.
5. Mackiewicz M, Shockley KR, Romer MA, et al. Macromolecule biosynthesis: a key function of sleep. *Physiol Genomics*. 2007;31(3):441-57.
6. Stickgold R, Walker MP. Memory consolidation and reconsolidation: what is the role of sleep? *Trends Neurosci*. 2005;28(8):408-15.
7. Rauchs G, Desgranges B, Foret J, et al. The relationships between memory systems and sleep stages. *J Sleep Res*. 2005;14(2):123-40.
8. Raichle ME, Gusnard DA. Appraising the brain's energy budget. *Proc Natl Acad Sci USA*. 2002;99(16):10237-9.
9. Reichert CF, Maire M, Schmidt C, et al. Sleep-wake regulation and its impact on working memory performance: the role of adenosine. *Biology (Basel)*. 2016;5(1).
10. Snel J, Lorist MM. Effects of caffeine on sleep and cognition. *Prog Brain Res*. 2011;190:105-17.
11. Mednick SC, Cai DJ, Kanady J, et al. Comparing the benefits of caffeine, naps and placebo on verbal, motor and perceptual memory. *Behav Brain Res*. 2008;193(1):79-86.
12. Ferrara M, De Gennaro L. How much sleep do we need? *Sleep Med Rev*. 2001;5(2):155-79.
13. Foley DJ, Monjan A, Simonsick EM, et al. Incidence and remission of insomnia among elderly adults: an epidemiologic study of 6,800 persons over three years. *Sleep*. 1999;22 Suppl 2:S366-72.
14. Duffy JF, Willson HJ, Wang W, et al. Healthy older adults better tolerate sleep deprivation than young adults. *J Am Geriatr Soc*. 2009;57(7):1245-51.

15. Adan A, Archer SN, Hidalgo MP, et al. Circadian typology: a comprehensive review. *Chronobiol Int*. 2012;29(9):1153-75.
16. Vgontzas AN, Liao D, Pejovic S, et al. Insomnia with short sleep duration and mortality: the Penn State cohort. *Sleep*. 2010;33(9):1159-64.
17. Van Dongen HP, Maislin G, Mullington JM, et al. The cumulative cost of additional wakefulness: dose-response effects on neurobehavioral functions and sleep physiology from chronic sleep restriction and total sleep deprivation. *Sleep*. 2003;26(2):117-26.
18. Weber M, Webb CA, Deldonno SR, et al. Habitual 'sleep credit' is associated with greater grey matter volume of the medial prefrontal cortex, higher emotional intelligence and better mental health. *J Sleep Res*. 2013;22(5):527-34.
19. Stamatakis KA, Punjabi NM. Long sleep duration: a risk to health or a marker of risk? *Sleep Med Rev*. 2007;11(5):337-9.
20. Hale L, Parente V, Dowd JB, et al. Fibrinogen may mediate the association between long sleep duration and coronary heart disease. *J Sleep Res*. 2013;22(3):305-14.
21. Brooks A, Lack L. A brief afternoon nap following nocturnal sleep restriction: which nap duration is most recuperative? *Sleep*. 2006;29(6):831-40.
22. Montgomery P, Burton JR, Sewell RP, et al. Fatty acids and sleep in UK children: subjective and pilot objective sleep results from the DOLAB study--a randomized controlled trial. *J Sleep Res*. 2014;23(4):364-88.
23. Ford ES, Cunningham TJ, Croft JB. Trends in self-reported sleep duration among US adults from 1985 to 2012. *Sleep*. 2015;38(5):829-32.
24. Ford ES, Cunningham TJ, Giles WH, et al. Trends in insomnia and excessive daytime sleepiness among U.S. adults from 2002 to 2012. *Sleep Med*. 2015;16(3):372-8.
25. Basner M, McGuire S, Goel N, et al. A new likelihood ratio metric for the psychomotor vigilance test and its sensitivity to sleep loss. *J Sleep Res*. 2015;24(6):702-13.
26. Spaeth AM, Dinges DF, Goel N. Phenotypic vulnerability of energy balance responses to sleep loss in healthy adults. *Sci Rep*. 2015;5:14920.
27. Killick R, Banks S, Liu PY. Implications of sleep restriction and recovery on metabolic outcomes. *J Clin Endocrinol Metab*. 2012;97(11):3876-90.
28. Spiegel K, Tasali E, Penev P, et al. Brief communication: Sleep curtailment in healthy young men is associated with decreased leptin levels, elevated ghrelin levels, and increased hunger and appetite. *Ann Intern Med*. 2004;141(11):846-50.
29. Banks S, Van Dongen HP, Maislin G, et al. Neurobehavioral dynamics following chronic sleep restriction: dose-response effects of one night for recovery. *Sleep*. 2010;33(8):1013-26.
30. Leproult R, Van Cauter E. Role of sleep and sleep loss in hormonal release and metabolism. *Endocr Dev*. 2010;17:11-21.
31. Cappuccio FP, D'Elia L, Strazzullo P, et al. Quantity and quality of sleep and incidence of type 2 diabetes: a systematic review and meta-analysis. *Diabetes Care*. 2010;33(2):414-20.
32. Leproult R, Holmback U, Van Cauter E. Circadian misalignment augments markers of insulin resistance and inflammation, independently of sleep loss. *Diabetes*. 2014;63(6):1860-9.
33. Waterhouse J, Reilly T, Atkinson G, et al. Jet lag: trends and coping strategies. *Lancet*. 2007;369(9567):1117-29.
34. Leproult R, Deliens G, Gilson M, et al. Beneficial impact of sleep extension on fasting insulin sensitivity in adults with habitual sleep restriction. *Sleep*. 2015;38(5):707-15.
35. Everson CA. Clinical assessment of blood leukocytes, serum cytokines, and serum immunoglobulins as responses to sleep deprivation in laboratory rats. *Am J Physiol Regul Integr Comp Physiol*. 2005;289(4):R1054-63.
36. Everson CA, Toth LA. Systemic bacterial invasion induced by sleep deprivation. *Am J Physiol Regul Integr Comp Physiol*. 2000;278(4):R905-16.
37. Cohen S, Doyle WJ, Alper CM, et al. Sleep habits and susceptibility to the common cold. *Arch Intern Med*. 2009;169(1):62-7.
38. Everson CA, Szabo A. Repeated exposure to severely limited sleep results in distinctive and persistent physiological imbalances in rats. *PLoS One*. 2011;6(8):e22987.
39. Everson CA, Henchen CJ, Szabo A, et al. Cell injury and repair resulting from sleep loss and sleep recovery in laboratory rats. *Sleep*. 2014;37(12):1929-40.
40. Jackson ML, Gunzelmann G, Whitney P, et al. Deconstructing and reconstructing cognitive performance in sleep deprivation. *Sleep Med Rev*. 2013;17(3):215-25.
41. Whitney P, Hinson JM, Jackson ML, et al. Feedback blunting: total sleep deprivation impairs decision making that requires updating based on feedback. *Sleep*. 2015;38(5):745-54.
42. Tucker AM, Whitney P, Belenky G, et al. Effects of sleep deprivation on dissociated components of executive functioning. *Sleep*. 2010;33(1):47-57.
43. Satterfield BC, Wisor JP, Field SA, et al. TNFalpha G308A polymorphism is associated with resilience to sleep deprivation-induced psychomotor vigilance performance impairment in healthy young adults. *Brain Behav Immun*. 2015;47:66-74.
44. Mullington JM, Simpson NS, Meier-Ewert HK, et al. Sleep loss and inflammation. *Best Pract Res Clin Endocrinol Metab*. 2010;24(5):775-84.
45. Imeri L, Opp MR. How (and why) the immune system makes us sleep. *Nat Rev Neurosci*. 2009;10(3):199-210.
46. Vgontzas AN, Fernandez-Mendoza J, Liao D, et al. Insomnia with objective short sleep duration: the most biologically severe phenotype of the disorder. *Sleep Med Rev*. 2013;17(4):241-54.
47. Fernandez-Mendoza J, Vgontzas AN, Calhoun SL, et al. Insomnia symptoms, objective sleep duration and hypothalamic-pituitary-adrenal activity in children. *Eur J Clin Invest*. 2014;44(5):493-500.
48. Fernandez-Mendoza J, Shea S, Vgontzas AN, et al. Insomnia and incident depression: role of objective sleep duration and natural history. *J Sleep Res*. 2015;24(4):390-8.
49. Fernandez-Mendoza J, Vgontzas AN, Kritikou I, et al. Natural history of excessive daytime sleepiness: role of obesity, weight loss, depression, and sleep propensity. *Sleep*. 2015;38(3):351-60.
50. Riemann D, Spiegelhalder K, Feige B, et al. The hyperarousal model of insomnia: a review of the concept and its evidence. *Sleep Med Rev*. 2010;14(1):19-31.
51. Riemann D, Nissen C, Palagini L, et al. The neurobiology, investigation, and treatment of chronic insomnia. *Lancet Neurol*. 2015;14(5):547-58.
52. Basta M, Chrousos GP, Vela-Bueno A, et al. Chronic insomnia and stress system. *Sleep Med Clin*. 2007;2(2):279-91.
53. Spiegelhalder K, Riemann D, Loring sleep. *Lancet Neurol*. 2015;14(6):571.
54. Hertenstein E, Nissen C, Riemann D, et al. The exploratory power of sleep effort, dysfunctional beliefs and arousal for insomnia severity and polysomnography-determined sleep. *J Sleep Res*. 2015;24(4):399-406.
55. Chouinard G. Issues in the clinical use of benzodiazepines: potency, withdrawal, and rebound. *J Clin Psychiatry*. 2004;65 Suppl 5:7-12.
56. Ebrahim IO, Shapiro CM, Williams AJ, et al. Alcohol and sleep I: effects on normal sleep. *Alcohol Clin Exp Res*. 2013;37(4):539-49.
57. Oswald I. Sleep, dreaming and drugs. *Proc R Soc Med*. 1969;62(2):151-3.
58. Lemoine P, Nir T, Laudon M, et al. Prolonged-release melatonin improves sleep quality and morning alertness in insomnia patients aged 55 years and older and has no withdrawal effects. *J Sleep Res*. 2007;16(4):372-80.
59. Luthringer R, Muzet M, Zisapel N, et al. The effect of prolonged-release melatonin on sleep measures and psychomotor performance in elderly patients with insomnia. *Int Clin Psychopharmacol*. 2009;24(5):239-49.
60. Kyle SD, Aquino MR, Miller CB, et al. Towards standardisation and improved understanding of sleep restriction therapy for insomnia disorder: A systematic examination of CBT-I trial content. *Sleep Med Rev*. 2015;23:83-8.
61. Kyle SD, Morgan K, Spiegelhalder K, et al. No pain, no gain: an exploratory within-subjects mixed-methods evaluation of the patient experience of sleep restriction therapy (SRT) for insomnia. *Sleep Med*. 2011;12(8):735-47.

62. Kyle SD, Miller CB, Rogers Z, et al. Sleep restriction therapy for insomnia is associated with reduced objective total sleep time, increased daytime somnolence, and objectively impaired vigilance: implications for the clinical management of insomnia disorder. *Sleep*. 2014;37(2):229-37.
63. Nelson HD. Menopause. *Lancet*. 2008;371(9614):760-70.
64. Ohayon MM. Severe hot flashes are associated with chronic insomnia. *Arch Intern Med*. 2006;166(12):1262-8.
65. de Zambotti M, Colrain IM, Sassoan SA, et al. Vagal withdrawal during hot flashes occurring in undisturbed sleep. *Menopause*. 2013;20(11):1147-53.
66. Baker FC, Driver HS. Self-reported sleep across the menstrual cycle in young, healthy women. *J Psychosom Res*. 2004;56(2):239-43.
67. Penev PD. Association between sleep and morning testosterone levels in older men. *Sleep*. 2007;30(4):427-32.
68. Lord C, Sekerovic Z, Carrier J. Sleep regulation and sex hormones exposure in men and women across adulthood. *Pathol Biol (Paris)*. 2014;62(5):302-10.
69. Sharkey KM, Crawford SL, Kim S, et al. Objective sleep interruption and reproductive hormone dynamics in the menstrual cycle. *Sleep Med*. 2014;15(6):688-93.
70. Wright KP, Jr., Bogan RK, Wyatt JK. Shift work and the assessment and management of shift work disorder (SWD). *Sleep Med Rev*. 2013;17(1):41-54.
71. Folkard S, Tucker P. Shift work, safety and productivity. *Occup Med (Lond)*. 2003;53(2):95-101.
72. Kubo T, Ozasa K, Mikami K, et al. Prospective cohort study of the risk of prostate cancer among rotating-shift workers: findings from the Japan collaborative cohort study. *Am J Epidemiol*. 2006;164(6):549-55.
73. Brown DL, Feskanich D, Sanchez BN, et al. Rotating night shift work and the risk of ischemic stroke. *Am J Epidemiol*. 2009;169(11):1370-7.
74. Mohebbi I, Shateri K, Seyedmohammadzad M. The relationship between working schedule patterns and the markers of the metabolic syndrome: comparison of shift workers with day workers. *Int J Occup Med Environ Health*. 2012;25(4):383-91.
75. Buxton OM, Cain SW, O'Connor SP, et al. Adverse metabolic consequences in humans of prolonged sleep restriction combined with circadian disruption. *Sci Transl Med*. 2012;4(129):129ra43.
76. Azagra-Calero E, Espinar-Escalona E, Barrera-Mora JM, et al. Obstructive sleep apnea syndrome (OSAS). Review of the literature. *Med Oral Patol Oral Cir Bucal*. 2012;17(6):e925-9.
77. Torelli F, Moscufo N, Garreffa G, et al. Cognitive profile and brain morphological changes in obstructive sleep apnea. *Neuroimage*. 2011;54(2):787-93.
78. Floras JS. Sleep apnea and cardiovascular risk. *J Cardiol*. 2014;63(1):3-8.
79. Gami AS, Howard DE, Olson EJ, et al. Day-night pattern of sudden death in obstructive sleep apnea. *N Engl J Med*. 2005;352(12):1206-14.
80. Morris LG, Kleinberger A, Lee KC, et al. Rapid risk stratification for obstructive sleep apnea, based on snoring severity and body mass index. *Otolaryngol Head Neck Surg*. 2008;139(5):615-8.
81. Djonlagic I, Guo M, Matteis P, et al. Untreated sleep-disordered breathing: links to aging-related decline in sleep-dependent memory consolidation. *PLoS One*. 2014;9(1):e85918.
82. Menon A, Kumar M. Influence of body position on severity of obstructive sleep apnea: a systematic review. *ISRN Otolaryngol*. 2013;2013:670381.
83. Spicuzza L, Caruso D, Di Maria G. Obstructive sleep apnoea syndrome and its management. *Ther Adv Chronic Dis*. 2015;6(5):273-85.
84. Ludka O, Konecny T, Somers V. Sleep apnea, cardiac arrhythmias, and sudden death. *Tex Heart Inst J*. 2011;38(4):340-3.
85. Djonlagic I, Guo M, Matteis P, et al. First night of CPAP: impact on memory consolidation attention and subjective experience. *Sleep Med*. 2015;16(6):697-702.
86. Luxton DD, Greenburg D, Ryan J, et al. Prevalence and impact of short sleep duration in redeployed OIF soldiers. *Sleep*. 2011;34(9):1189-95.
87. Livingston WS, Rusch HL, Nersesian PV, et al. Improved Sleep in Military Personnel is Associated with Changes in the Expression of Inflammatory Genes and Improvement in Depression Symptoms. *Front Psychiatry*. 2015;6:59.
88. Heinzelmann M, Reddy SY, French LM, et al. Military personnel with chronic symptoms following blast traumatic brain injury have differential expression of neuronal recovery and epidermal growth factor receptor genes. *Front Neurol*. 2014;5:198.
89. Bryan CJ. Repetitive traumatic brain injury (or concussion) increases severity of sleep disturbance among deployed military personnel. *Sleep*. 2013;36(6):941-6.
90. Collen J, Orr N, Lettieri CJ, et al. Sleep disturbances among soldiers with combat-related traumatic brain injury. *Chest*. 2012;142(3):622-30.
91. Scullin MK, Bliwise DL. Sleep, cognition, and normal aging: integrating a half century of multidisciplinary research. *Perspect Psychol Sci*. 2015;10(1):97-137.
92. Gildner TE, Liebert MA, Kowal P, et al. Associations between sleep duration, sleep quality, and cognitive test performance among older adults from six middle income countries: results from the Study on Global Ageing and Adult Health (SAGE). *J Clin Sleep Med*. 2014;10(6):613-21.
93. Available at: <http://www.neurologyreviews.com/specialty-focus/sleep/article/how-much-sleep-is-required-for-peak-cognitive-performance/fed5d4521ed235985885ea521d11eccc.html>. Accessed April 29, 2016.
94. Laing EE, Johnston JD, Moller-Levet CS, et al. Exploiting human and mouse transcriptomic data: Identification of circadian genes and pathways influencing health. *Bioessays*. 2015;37(5):544-56.
95. Lazar AS, Lazar ZI, Dijk DJ. Circadian regulation of slow waves in human sleep: Topographical aspects. *Neuroimage*. 2015;116:123-34.
96. Archer SN, Laing EE, Moller-Levet CS, et al. Mistimed sleep disrupts circadian regulation of the human transcriptome. *Proc Natl Acad Sci U S A*. 2014;111(6):E682-91.
97. Dijk DJ, Skeldon AC. Biological rhythms: Human sleep before the industrial era. *Nature*. 2015;527(7577):176-7.
98. Lo JC, Leong RL, Loh KK, et al. Young Adults' Sleep Duration on Work Days: Differences between East and West. *Front Neurol*. 2014;5:81.
99. Ohayon MM, Roth T. Prevalence of restless legs syndrome and periodic limb movement disorder in the general population. *J Psychosom Res*. 2002;53(1):547-54.
100. Ekboom K, Ulfberg J. Restless legs syndrome. *J Intern Med*. 2009;266(5):419-31.
101. Bosch JL, Weiss JP. The prevalence and causes of nocturia. *J Urol*. 2013;189(1 Suppl):S86-92.
102. Yoshimura K. Correlates for nocturia: a review of epidemiological studies. *Int J Urol*. 2012;19(4):317-29.
103. Yazici CM, Kurt O. Combination therapies for the management of nocturia and its comorbidities. *Res Rep Urol*. 2015;7:57-63.
104. Tacklind J, Fink HA, Macdonald R, et al. Finasteride for benign prostatic hyperplasia. *Cochrane Database Syst Rev*. 2010(10):Cd006015.
105. Fusco F, Palmieri A, Ficarra V, et al. Alpha1-blockers improve benign prostatic obstruction in men with lower urinary tract symptoms: A systematic review and meta-analysis of urodynamic studies. *Eur Urol*. 2016.
106. Blunden S, Galland B. The complexities of defining optimal sleep: empirical and theoretical considerations with a special emphasis on children. *Sleep Med Rev*. 2014;18(5):371-8.

EUROPEAN MILK THISTLE

Ultimate Protection For Your Liver

Milk thistle extract—rich in *silymarin*—is one of nature's most powerful weapons to support liver health. Scientific studies demonstrate silymarin's ability to provide potent protection for your liver.^{1,2}

Life Extension®'s European Milk Thistle contains standardized, top-grade potencies of *silymarin*, *silybin*, *isosilybin A*, and *isosilybin B*, providing a full spectrum of liver-supportive compounds. This unique formula includes *phosphatidylcholine*, a nutrient that promotes better absorption of milk thistle extract.³

The **silymarin** contained in **European Milk Thistle** is absorbed nearly **5 times** better than silymarin alone, and its bioavailability to the liver is **10 times** better.

References

1. *Mol Nutr Food Res.* 2009 Apr;53(4):460-6.
2. *Environ Toxicol.* 2007 Oct;22(5):472-9.
3. *Altern.Med Rev.* 2009;14(3):226-46.

SILIPHOS® is a registered trademark of Indena S.p.A., Italy.

European Milk Thistle Advanced Phospholipid Delivery

Item #01922 • 60 Softgels

	Retail Price	Your Price
1 bottle	\$28	\$21
4 bottles		\$18.75 each

Non-GMO



To order **European Milk Thistle Advanced Phospholipid Delivery**, call 1-800-544-4440 or visit www.LifeExtension.com

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



Sweet DREAMS

Choose the Melatonin That's Right For You

Healthy sleep is one of the best ways to feel revitalized and maintain optimal health. Increasingly, research has shown the health benefits of a good night's sleep. Melatonin is one of the most popular supplements for supporting sleep.

Now, pick the right melatonin for your needs with doses ranging from **300 mcg** to **10 mg**. For optimal results, melatonin should be taken within 30-60 minutes of going to sleep.



Melatonin Timed Release 300 mcg
100 vegetarian tablets
Retail: \$12
Your Price: \$9
Item # 01787



Melatonin 500 mcg
200 vegetarian capsules
Retail: \$18
Your Price: \$13.50
Item # 01083



Melatonin Timed Release 750 mcg
60 vegetarian tablets
Retail: \$8
Your Price: \$6
Item # 01788



Melatonin 1 mg
60 capsules
Retail: \$5
Your Price: \$3.75
Item # 00329



Melatonin 10 mg
60 vegetarian capsules
Retail: \$28
Your Price: \$21
Item # 00331



Melatonin 300 mcg
100 vegetarian capsules
Retail: \$5.75
Your Price: \$4.31
Item # 01668



Melatonin 3 mg
60 vegetarian capsules
Retail: \$8
Your Price: \$6
Item # 00330



Melatonin 3 mg
60 vegetarian lozenges
Retail: \$8
Your Price: \$6
Item # 00332



Melatonin Timed Release 3 mg
60 vegetarian tablets
Retail: \$12
Your Price: \$9
Item # 01786



Natural Sleep® 3 mg
60 vegetarian capsules
Retail: \$13
Your Price: \$9.75
Item # 01444



Natural Sleep® Melatonin 5 mg
60 vegetarian capsules
Retail: \$18
Your Price: \$13.50
Item # 01445

ChromeMate®, a patented, biologically active oxygen-coordinated niacin-bound chromium complex, is a registered trademark of Interhealth Nutritionals Inc.

Caution: Consult your health care provider before taking this product if you are being treated for a medical condition (especially autoimmune or depressive disorders). Use caution if combining with alcohol. This product is not intended for children, pregnant or lactating women, or women trying to become pregnant. Do not attempt to drive or operate heavy machinery after taking this product.

To order any of these premium-grade Melatonin supplements, call **1-800-544-4440** or visit www.LifeExtension.com

Boost Mental *and* Physical Energy with **RHODIOLA EXTRACT**

Restore vital cellular energy with Life Extension®'s Rhodiola.

Prized for its potent adaptogenic properties, **Rhodiola** has long been used by elite athletes, soldiers, and even cosmonauts to enhance mental and physical stamina.

Scientific research shows that **Rhodiola**, an optimizing, adaptogenic herb supports:

- Cellular energy production¹
- Enhanced physical endurance²
- Clarity and mental performance under stress³

Life Extension®'s **Rhodiola** is uniquely standardized for adaptogenic compounds *rosavins* and *salidroside*s.

Caution: Individuals with manic or bipolar disorder should not use rhodiola. Take early in the day if Rhodiola Extract interferes with your sleep.

Life Extension® has formulated a **Rhodiola Extract** that provides **250 mg** of *Rhodiola rosea* extract in each capsule.

Rhodiola

Item #00889 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$14	\$10.50
4 bottles		\$9 each

Non-GMO

References

1. Proceedings Institute of Cytology of Russian Academy of Science. 1971;89-92.
2. *Int J Sport Nutr Exerc Metab.* 2004 Jun;14(3):298-307.
3. *Phytomedicine.* 2000 Oct;7(5):365-71.



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

BY MAUREEN FIONA

Newly Discovered Benefits of **LUTEIN**

The carotenoid **lutein** has long been studied for its vision-protective properties. It has been shown to reduce the risk of two of the leading causes of blindness: age-related macular degeneration and cataracts.¹

A recent study has revealed that we've only scratched the surface of lutein's health-promoting benefits. In a first-of-its-kind analysis, lutein has been associated with a reduced risk for **cardiometabolic diseases**.

A large meta-analysis involving **71 published papers** and representing more than **387,000 individuals** showed that people with **higher** lutein intake, or higher blood concentrations of lutein, have a reduced risk of coronary heart disease, stroke, and metabolic syndrome.²

The reason lutein provides such wide-reaching effects is because of its ability to protect tissues from oxidative stress and inflammation—two factors that play a major role in cardiovascular and metabolic diseases.

This study will likely change the way we think of lutein, broadening its appeal to everyone who wants to optimally protect their blood vessels, heart, and brain against the ravages of oxidative stress and chronic inflammation.

Beyond Eye Health

Lutein is a type of *carotenoid*, a natural compound found in vegetables and other plants. Although it is yellow, it is especially abundant in dark green, leafy vegetables such as kale.³

Lutein has received a great deal of attention for its ability to help reduce the incidence of age-related macular degeneration and cataracts, two of the leading causes of blindness.¹ These benefits arise from lutein's ability to provide structural support for the macula pigment while protecting the eye from the oxidative stresses caused by exposure to light and oxygen. It also helps reduce inflammation.^{2,4,5}

Researchers from ErasmusAGE, a center for aging research, hypothesized that these mechanisms of action could make lutein beneficial for cardiometabolic health as well, since oxidative stress and inflammation play a role in both cardiovascular and metabolic diseases.

That's why Dr. Elizabeth Leermakers and her research team decided to conduct an expansive search of the medical literature for studies involving either lutein intake or blood concentrations of lutein in the context of cardiometabolic diseases.²

In this type of study, called a **systematic review and meta-analysis**, researchers review multiple studies on a single topic, and then combine the findings mathematically for analysis. The results of these types of studies have the potential to provide strong indicators of clinical evidence.⁶

And in fact, the results of this particular study were so impactful that they could open the door to new therapeutic benefits of lutein. Instead of being viewed solely as a vision-saving nutrient, lutein could

be seen as one that enhances total-body health and longevity.

After examining 4,377 individual studies, the researchers narrowed their review to **71 relevant articles** that included a total of more than **387,000 participants**.² They found that in all of these studies there were some very powerful and significant associations between lutein and cardiometabolic diseases.

Compared to people in the group with the lowest 1/3 of intake or blood concentration of lutein, those in the top 1/3 were found to have:²

- A **12% reduction** in the risk of having **coronary heart disease** (atherosclerosis of the heart's own blood vessels, which may lead to angina and ultimately to heart attacks)
- An **18% reduction** in the risk of having a **stroke**
- A **25% reduction** in the risk of having **metabolic syndrome**, the cluster of medical conditions that includes abdominal obesity, elevated blood pressure, elevated fasting glucose levels, elevated blood triglycerides, and low levels of protective high-density lipoprotein (HDL) cholesterol

This study is significant because it is the first meta-analysis to show that higher intake or blood levels of lutein have such prominent cardiometabolic benefits.

And ultimately, it validates the use of lutein for reducing the risk for stroke, heart attack, and metabolic syndrome, which itself is associated with an increased risk for developing cardiovascular disease, diabetes, and death from multiple causes.^{7,8}

How It Works

Numerous studies have shown that lutein combats many of the underlying factors in both cardiovascular and metabolic diseases.

Prior studies have shown significant associations between higher lutein levels and improvements in blood lipid levels, especially with higher HDL (good) cholesterol.⁹⁻¹¹

Inflammation is another underlying factor in the development of cardiovascular disease and atherosclerosis. Studies have shown that lutein may activate branches of the immune system that suppress chronic inflammation.^{2,4} In addition, previous studies have shown that lutein is effective in reducing arterial wall thickening, which contributes to poor blood flow and raises the risk of heart attack and stroke.^{2,12,13}





Together, these factors may help to explain the reduced risks of heart attack, stroke, and metabolic syndrome that were seen in the recent meta-analysis.

Summary

Lutein has long been associated with vision protection, which it provides both by reducing oxidative stresses in the eye and by lowering chronic inflammation that can contribute to cataracts and age-related macular degeneration.

Oxidative stress and chronic inflammation also contribute to cardiometabolic diseases such as heart disease, stroke, and diabetes.

In this first-of-its-kind analysis, scientists discovered that higher lutein intake and blood concentrations are strongly and significantly related to reductions in cardiometabolic risks. This analysis showed that lutein reduces the risk of stroke, heart attack, and metabolic syndrome.

Based on the results of this study, people interested in supporting their cardiometabolic health should consider boosting their lutein intake.

Most readers of this publication have been obtaining potent doses of lutein in the supplements they have been using for the past several decades. ●

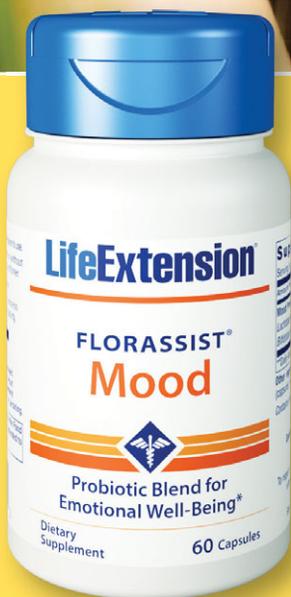
If you have any questions on the scientific content of this article, please call a Life Extension® Wellness Specialist at 1-866-864-3027.

References

1. Scripsema NK, Hu DN, Rosen RB. Lutein, zeaxanthin, and meso-zeaxanthin in the clinical management of eye disease. *J Ophthalmol*. 2015;2015:865179.
2. Leermakers ET, Darweesh SK, Baena CP, et al. The effects of lutein on cardiometabolic health across the life course: a systematic review and meta-analysis. *Am J Clin Nutr*. 2016;103(2):481-94.
3. Abdel-Aal el SM, Akhtar H, Zaheer K, et al. Dietary sources of lutein and zeaxanthin carotenoids and their role in eye health. *Nutrients*. 2013;5(4):1169-85.
4. Kijlstra A, Tian Y, Kelly ER, et al. Lutein: more than just a filter for blue light. *Prog Retin Eye Res*. 2012;31(4):303-15.
5. Sujak A, Gabrielska J, Grudzinski W, et al. Lutein and zeaxanthin as protectors of lipid membranes against oxidative damage: the structural aspects. *Arch Biochem Biophys*. 1999;371(2):301-7.
6. McNamara ER, Scales CD. Role of systematic reviews and meta-analysis in evidence-based clinical practice. *Indian J of Urol*. 2011;27(4):520-4.
7. Kaur J. A comprehensive review on metabolic syndrome. *Cardiol Res Pract*. 2014;2014:943162.
8. Micucci C, Valli D, Matacchione G, et al. Current perspectives between metabolic syndrome and cancer. *Oncotarget*. 2016.
9. Renzi LM, Hammond BR, Jr, Dengler M, et al. The relation between serum lipids and lutein and zeaxanthin in the serum and retina: results from cross-sectional, case-control and case study designs. *Lipids Health Dis*. 2012;11:33.
10. Sugiura M, Nakamura M, Ogawa K, et al. Associations of serum carotenoid concentrations with the metabolic syndrome: interaction with smoking. *Br J Nutr*. 2008;100(6):1297-306.
11. Wang Y, Chung SJ, McCullough ML, et al. Dietary carotenoids are associated with cardiovascular disease risk biomarkers mediated by serum carotenoid concentrations. *J Nutr*. 2014;144(7):1067-74.
12. Kowluru RA, Menon B, Gierhart DL. Beneficial effect of zeaxanthin on retinal metabolic abnormalities in diabetic rats. *Invest Ophthalmol Vis Sci*. 2008;49(4):1645-51.
13. Izumi-Nagai K, Nagai N, Ohgami K, et al. Macular pigment lutein is anti-inflammatory in preventing choroidal neovascularization. *Arterioscler Thromb Vasc Biol*. 2007;27(12):2555-62.



Probiotic Blend Supports Positive **MOOD** and **RELAXATION**



Research suggests specific **probiotics** positively influence biochemical signaling between the gastrointestinal tract and the nervous system—resulting in positive effects on mood.¹

FLORASSIST® Mood provides **3 billion** colony-forming units of *Lactobacillus helveticus* strain R0052 and *Bifidobacterium longum* strain R0175.

Human clinical research conducted on this dual-probiotic combination demonstrated improvements in mood, reduction in perceived stress, and promotion of relaxation.^{2,3}

Two daily capsules of **FLORASSIST® Mood** naturally promote relaxation and improve mood.

FLORASSIST® Mood

Item #02000 • 60 capsules

	Retail Price	Your Price
1 bottle	\$33	\$24.75
4 bottles		\$22.50 each

Non-GMO

Contains milk and soybeans.

References

1. *Cell*. 2015 Apr 9;161(2):193-4.
2. *Gut Microbes*. 2011 Jul-Aug;2(4):256-61.
3. *Br J Nutr*. 2011 Mar;105(5):755-64.

To order **FLORASSIST® Mood**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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

Tremendous Value in One Softgel

HEALTH BOOSTER

Provides Important Nutrients Missing from Multi-Vitamins

• **Saffron** and **alpha-carotene** to promote eye health.



• **Vitamin K2** is the active form that keeps calcium in bones and out of arteries.



• **Gamma tocopherols** to quench oxidation missed by conventional **vitamin E**.



• **Vitamin K1** is rapidly absorbed and partially converts to vitamin K2 in the body.

• **Lycopene** to promote prostate health.



• **Zeaxanthin, lutein, and meso-zeaxanthin** to support healthy vision.

• **Chlorophyllin** to protect against environmental DNA damage.



• **Sesame lignans** to boost gamma tocopherol activity.

• **Black currant extract** to promote eye health.



• **Blueberry** extract to sustain healthy **neurological** function.

• **Methylcobalamin** form of **B12** to better support healthy nerve function.



Once-Daily Health Booster

Item #01991 • 60 softgels (two-month supply)

	Retail Price	Your Price
1 bottle	\$54	\$40.50
4 bottles		\$38 each

Non-GMO

Caution: Avoid use during pregnancy. Consult your healthcare practitioner before using this product if you are taking anti-coagulant or anti-platelet medications or have a bleeding disorder.

Lyc-O-Mato® is a registered trademark of LycoRed, LTD. **LuteinPlus**® and **Mz**® are registered trademarks of NutriProducts Ltd., UK, licensed under U.S. Patent 8,623,428.

*The same nutrients sold separately would cost **2-3 times** more money!*

To order **Once-Daily Health Booster**, call **1-800-544-4440** or visit **www.LifeExtension.com**

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

Super Selenium *Supports* HEART *and* BRAIN HEALTH



Linked to Longevity

Selenium's longevity power comes from its ability to offer powerful protection throughout the body, including heart support, brain support, and healthy cell division.^{1,2}

Super Selenium Complex has three different forms of selenium—each of which uniquely acts along a different pathway to support healthy cell division.

- Sodium selenite
- L-selenomethionine
- Selenium-Methyl L-Selenocysteine

References

1. *Biol Trace Elem Res.* 2004 Oct;101(1):73-86.
2. *Biol Trace Elem Res.* 2011 Sep;142(3):274-83

Super Selenium Complex

Item #01778 • 100 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$14	\$10.50
4 bottles		\$9 each



Each bottle provides a supply that lasts more than three months.

Non-GMO

SelenoPure™ is a trademark of Nutrition 21.

Caution: If you are taking anticoagulant or antiplatelet medications, or have a bleeding disorder, consult your healthcare provider before taking this product.

To order **Super Selenium Complex**, call **1-800-544-4440** or visit **www.LifeExtension.com**

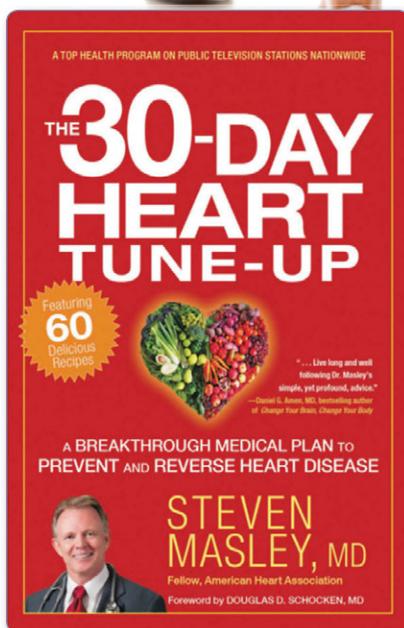
BY ASTRID DERFLER KESSLER

AUTHOR INTERVIEW



The 30-Day Heart Tune-Up

An Interview with Dr. Steven Masley



Steven Masley, MD, a practicing physician from St. Petersburg, Florida, is a fellow with the American Heart Association, the American Academy of Family Physicians, and the American College of Nutrition. He wrote *The 30-Day Heart Tune-Up* because he was becoming increasingly frustrated with the current state of affairs when it comes to America's number one killer, cardiovascular disease. His goal, and the purpose of the book, he says, is to prevent heart disease and stroke by making treatment options available to patients long before they develop the disease. The book provides the tools that are needed—as well as 60 heart-healthy recipes—to accomplish this goal. Following this plan for one month will help prevent plaque from forming, reduce plaque if it already exists, and potentially save your life, says Dr. Masley.

LE: What makes your approach to heart disease different from other plans?

SM: Most doctors focus on lowering cholesterol and blood pressure to prevent heart disease. Their approach relies on drug therapy and laboratory testing, which have been a tremendous boon to the pharmaceutical and medical laboratory industry, but not necessarily to patients. Statins...still have an appropriate use in some patients, yet they increase the risk of diabetes and unfortunately, their testosterone-lowering effect may cause weight gain, reduce sexual enjoyment, and lower a person's drive to stay fit. Managing cholesterol and blood pressure have helped reduce the rate of heart attacks and strokes, but they don't do enough. That's because we're focusing on the wrong issue.

The 30-day heart tune-up is about shrinking arterial plaque, improving circulation, and strengthening your heartbeat. I recommend neither an extreme vegan diet, which few people can maintain for more than a week or

two, nor deprivation, nor expensive medications that merely treat symptoms rather than the cause (although I agree that some medications, when indicated, do help make the transition to optimal health).

So how can you accomplish this goal in such a short time? Here are your tools:

- Incorporate five easy-to-remember categories of heart-healing foods into your diet.
- Engage in exercise that strengthens your heart and arteries.
- Learn to better manage stress.
- Follow a customized, heart-friendly supplement plan.

LE: What is the number-one risk factor people face when it comes to cardiovascular disease?

SM: We used to believe that high cholesterol was the number-one risk factor. But we must look at the whole picture. Having metabolic syndrome, appropriately called "diabesity" by Dr. Mark Hyman in his book *The Blood Sugar Solution*

and also known as prediabetes, is a greater risk factor than having high cholesterol. In fact, it's the number one risk factor of cardiovascular disease. (To read an interview with Dr. Hyman, see the May 2013 issue of *Life Extension Magazine*®.)

More than **30%** of adults (**50%** of baby boomers) have this condition. The bad news is metabolic syndrome can kill you before you ever develop diabetes. Because it changes your cholesterol profile, increases inflammation, and raises your blood pressure levels in ways that are similar to those of diabetes, it can cause a heart attack or stroke.

LE: Your book offers the tools people need to, as you put it, "make a U-turn on the road to heart disease in just 30 days." Can you elaborate on your 30-day plan?

SM: It's about shrinking arterial plaque, improving circulation, and strengthening your heartbeat. You will be using four important tools: heart-healing foods, exercise that strengthens your heart and arteries, stress management, and a customized heart-friendly supplement plan. The [plan is] based on my life's work. I've devoted every hour of every day striving to make my patients' lives and hearts better without their having to resort to surgery or other invasive procedures. The losses we suffer due to cardiovascular disease are great—personal pain, decreased income, reduced productivity, physical suffering, and even premature death. So much of this anguish is unnecessary. Instead of muddling through as a helpless, hopeless victim of heart disease, with this plan, you will have all the tools you need to attack and defeat this scourge. It is possible to reverse cardiovascular disease and certainly to prevent it.



Tune Up Your Heart with Supplements

LE: Feeding the heart the nutrients it needs for total health isn't easy to do from diet alone. Supplements are key to good health. What supplements do you recommend for total heart health?

SM: Everyone needs a personalized eating plan to meet their key nutrient needs because no one eats well all the time. Just bear in mind that...vitamins, minerals, herbals, and other valuable treatments...can only enhance a healthy eating plan and an optimal lifestyle. They can't replace it. They will never make up for snacking on junk food or being stressed or inactive.

Nevertheless, supplements can benefit your heart health substantially even if you eat well, exercise regularly, and manage stress. They can also provide some nutrients you may otherwise be lacking. At least **80%** of Americans are nutrient deficient. Yes, we get plenty of calories, but we don't consume enough vitamins and minerals. Nutrient content has been dropping in our highly processed food supply. That's why supplements are an integral part of your 30-day heart tune-up. And, they may boost your heart health in ways that food alone cannot.

LE: What should a high-quality multivitamin contain?

SM: A multivitamin is a good place to start. Keep in mind that a multivitamin won't supply fiber, fish oil, magnesium, and potassium in a significant way. These are critical heart nutrients.

To begin with [you need at least] **1,500 to 2,000 IUs** of vitamin D. You'll also want a pill that has mixed carotenoids, not just beta-carotene, as well as mixed forms of folate, especially 5-MTHF (5-methyltetrahydrofolate), not just

folate. If you're over 50, you'll need adequate B12—at least **100 mcg** daily, but if you take an acid blocker, a minimum of **500 mcg** is recommended. Vitamin K should come as a mixture of vitamin K1 and K2, as I'll discuss later. And finally, you should take **75 to 150 mcg** of iodine.

LE: Vitamin D is critical to so many aspects of overall good health, from diabetes and bone health to cancer and cognitive health, but how does it relate to heart health?

SM: Vitamin D is called a fat-soluble vitamin, but once converted to its active form, it functions like a hormone. Every cell in the body appears to have vitamin D receptors, so having adequate amounts in your bloodstream is absolutely critical to your health. Studies show that people with higher blood levels of vitamin D have lower rates of heart attacks and strokes and better blood pressure and weight control. Clearly, vitamin D is essential for optimal heart and general health. As a hormone, vitamin D communicates directly with your genes, influencing how they function, and tells the cells which protein to produce.

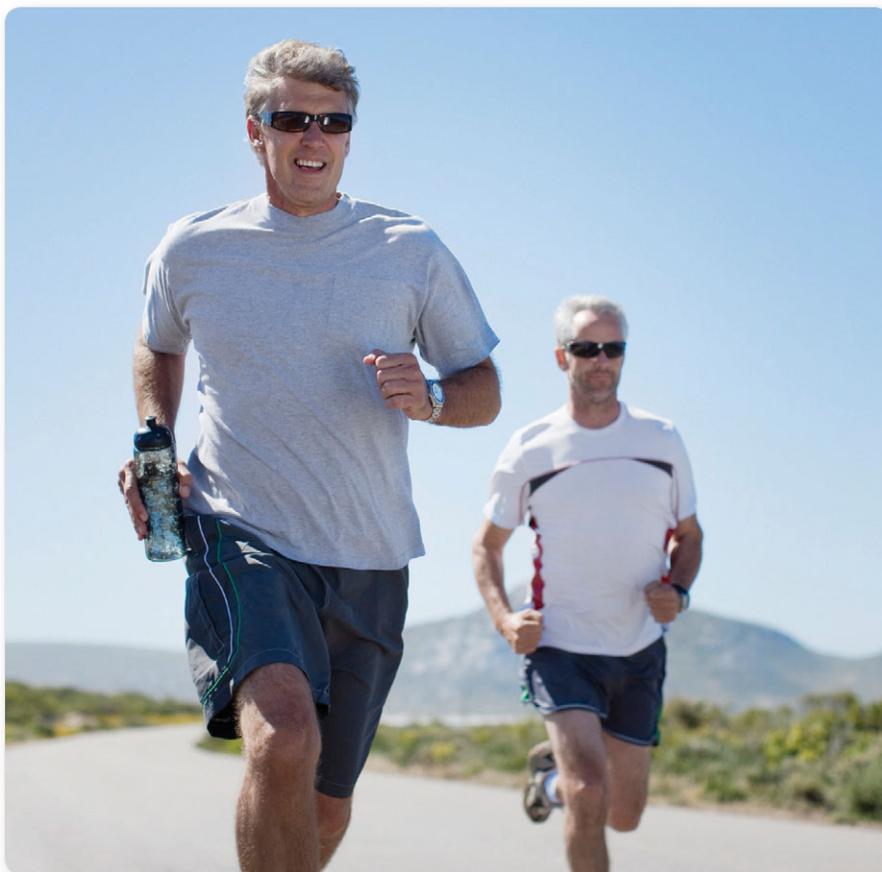
How much vitamin D [you need] would depend on your current blood levels. For most people, optimal dosages vary from **1,500 to 3,000 IU** daily, with the goal to achieve a blood level of **40 to 70 ng/mL**. (**Life Extension**[®] recommends a blood level of **50 to 80 ng/mL**.) Less than **32 ng/mL** is deficient. If your level is very low, then taking **5,000 IU** daily for two to three months will probably bring you back to normal. (**Life Extension** recommends that healthy adults supplement each day with at least **5,000 IU** of vitamin D.)

LE: Vitamin K, in the form of vitamin K1 and vitamin K2, is critical to cardiovascular health, as well as offering protection against arterial calcification, bone loss, cancer, metabolic syndrome, and aging. You call it an essential supplement.

SM: Vitamin K is essential for cardiovascular health because it prevents calcium from shifting from your bones to your arteries. Not surprisingly, then, vitamin K is essential to keeping bones strong. With increasing vitamin K deficiency, artery walls calcify, blood pressure increases, and the lining of your arteries grows more plaque. The minimum dietary intake of vitamin K for proper clotting is around **100 mcg** per day—**90 mcg** for women and **120 mcg** for men. Yet bones and arteries function much better with **250 mcg** and preferably **1,000 mcg** of vitamin K daily. (**Life Extension** recommends **2,700 mcg** of vitamins K1 and K2 together.)

There are two primary forms of vitamin K—K1 and K2. K1 comes from leafy green vegetables and provides most of the dietary intake in the US. K2 comes from fermented soy products and most Americans [don't] consume enough. Keep in mind that calcium supplements may already provide some of the K your body needs. Calculate how much vitamin K you get from your diet and other vitamins, and supplement accordingly.

LE: Several large studies have shown benefits of vitamin E intake in reducing cardiovascular disease and death from heart attacks. What do you recommend regarding vitamin E for cardiovascular health?



SM: Vitamin E is made up of several molecules—four tocopherols (alpha, beta, gamma, delta) and four tocotrienols (alpha, beta, gamma, delta). Gamma and delta tocopherols and tocotrienols improve the advanced lipid profile, so if you're taking...a vitamin E supplement, make sure it has mixed forms of vitamin E—in particular, delta and gamma tocopherols.

For Those With Congestive Heart Failure and Heart Disease

LE: Your book is focused on the prevention of heart disease and keeping the heart healthy. But what if someone has congestive heart failure or heart disease—what do

you recommend as far as supplements are concerned?

SM: With heart failure, the heart is starved for energy. Without it, your heart can't pump efficiently, and the fluid from your blood backs up into your lungs. One of the many keys to restoring function, then, is restoring heart cell energy.

Mitochondria produce most of the energy your cells use. Your heart cells have more mitochondria than any others in the body. Dr. Stephen Sinatra, a nationally known integrative cardiologist, was one of the pioneers in advocating therapy to enhance heart cell mitochondrial function—something he called metabolic cardiology. He focused on three key supplements: CoQ10, ribose, and carnitine. Clinical experi-

ence using these agents has shown they're excellent in improving the quality of life for some people who have heart failure.

- **CoQ10—50 to 200 mg** daily. I suggest you ask your doctor to confirm that you have a blood level of at least **2 to 2.5 mcg/mL** of CoQ10. (**Life Extension** recommends a level of **>4 mcg/mL** for those with congestive heart failure.)
- **Ribose—5 grams** twice daily.
- **Carnitine—2,000 to 3,000 mg**. (Some people take **3,000 mg** of lower cost **taurine** in lieu of carnitine.)

LE: What other supplements do you recommend for those who currently have heart disease?

SM: Two other agents have great promise to enhance mitochondrial function and energy production. These are curcumin and resveratrol. [These] compounds are essential to controlling artery inflammation and oxidative stress, but are required in such large amounts that it is unrealistic to get an adequate dosage from food (or red wine, in the case of resveratrol).

Curcumin is such an outstanding anti-inflammatory that I even prescribe it to my patients with arthritis, those who want to limit cognitive decline, and for cancer prevention. Curcumin absorption is highly variable, so it is critical to take a high-quality form with proven absorption rates.

As [stated] earlier, vitamin K prevents artery calcification. If you have advanced heart disease/failure, ask about adding extra vitamin K. Increase magnesium to **300 to 500 mg** twice daily. It will help relax blood vessels and enhance blood pressure control, making pumping easier for the heart. At some point on this dosage, ask your doctor to

measure your blood cell magnesium levels to confirm you're in the normal range.

Arginine is also an excellent nutrient to enhance artery function and improve blood pressure control. It is the building block that the body uses to make nitric oxide, which is essential to arterial health and wellness. Nitric oxide induces the arteries to dilate... [I suggest] a supplement with **1,000 mg** twice daily.

If You're On a Cholesterol-Lowering Medication

LE: Statin medications are taken by millions of Americans. You recommend additional supplements for those on statins.

SM: Statin medications decrease not only cholesterol but also compounds in the body derived from cholesterol, such as testosterone, CoQ10, and other substances that repair muscle. Yes, cholesterol-lowering medications help reduce heart attacks and strokes in people with known heart disease and for those at high risk, but they can produce a variety of symptoms. Most people assume the benefit from taking a statin is related to its cholesterol-lowering effects, but other important benefits are due to the fact that statins also decrease artery inflammation and make your blood less sticky, so it's less likely to clot.

CoQ10 is essential for energy production, especially for heart cells. Every cell has energy-generating organelles called mitochondria...which require CoQ10 to produce energy. Studies show that the ability to produce CoQ10 drops by **20%** in people who take a statin...and a big concern is that it might decrease your energy and

mental sharpness and interfere with quality of life. Most holistic and integrative physicians recommend that patients on a statin should take CoQ10 with it...to bring the body back to its normal, pre-statin state. I find **50 to 100 mg** of high-quality and well-absorbed product will bring the CoQ10 level back to pre-statin levels.

Summary

LE: You've integrated the 30-day heart tune-up in many of your patients. What do they tell you after that one-month mark?

SM: The combination of eating heart-friendly foods, building your fitness level, managing your stress, and meeting your heart's nutritional needs will not only help you dramatically reduce your risk for heart disease and stroke, but it will also turn back the clock on aging—allowing you to regain energy and vitality you haven't known in years. The greatest thanks I get from my patients have been their frequent observation after the first 30 days [when they tell me] "I forgot how much better I could feel."

LE: Thank you, Dr. Masley, for sharing this important information with our readers. ●

Steven Masley, MD, is a Fellow with the American Heart Association, the American Academy of Family Physicians, and with the American College of Nutrition. He has devoted his medical career to the study of heart disease and aging, and has published significant research on these subjects in leading medical journals. Currently he is the president of the Masley Optimal Health Center in St. Petersburg, Florida, and has a clinical appointment with the University of South Florida.

For more information or to contact Dr. Masley, visit drmasley.com or email info@drmasley.com

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

To order a copy of *The 30-Day Heart Tune-Up*, call 1-800-544-4440 or visit LifeExtension.com

Item #34002
Retail Price: \$17.59
Your Price: \$13.19



Jarrow
FORMULAS®

BROCCO**MAX**® THE SULFORAPHANE GENERATOR™



AN ULTIMATE LIVER & CELLULAR DETOXIFIER & PROTECTOR*

Broccoli (*Brassica oleracea spp.*) is associated with antioxidant and cellular protection due to its components glucoraphanin (GR), aka sulforaphane glucosinolate (SGS), and myrosinase which together produce sulforaphane, a powerful, beneficial electrophile and inducer of the liver's potent Phase 2 Detoxification System.*

Jarrow Formulas' BroccoMax® contains a standardized concentration of SGS and the myrosinase enzyme from broccoli seeds that have been processed using ultra clean super critical CO₂ technology. Vitamin C (as calcium ascorbate) catalyzes the sulforaphane generating action of the myrosinase enzyme.

BroccoMax® is delivered in delayed release vegetarian capsules.

To order Jarrow Formulas' BroccoMax® 60 veggie caps, Item # 26576

Call 1-800-544-4440 or visit www.LifeExtension.com

Retail price \$26.95 Your price \$20.21

MADE IN USA

BroccoMax® is protected by
U.S. Patent # 9,017,666

DRcaps
PROTECTED

**VEGGIE
CAPS**

**GLUTEN
FREE**

vegan

**NON
GMO**

DRcaps® is a registered trademark of Capsugel®

*These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

VISIT US AT WWW.JARROW.COM FOR MORE PRODUCT INFORMATION

© 2016 Jarrow FORMULAS®

RAISING THE STANDARD.

We have always been about Trust.

Scan the QR Code to view test results for every lot.



For provable potency, purity and identity, Master Supplements continues to raise the standard.



We recently patented the ability to scan a QR code on a probiotic, enzyme, or fiber supplement that leads to product quality information because it is important to us that what is on the label, is in the bottle. Our products help support digestive and immune health. They are backed with years of efficacy and scientific research.* If you try them and are not satisfied with the results, we offer a **100% money back guarantee**.



WHAT SETS US APART?

- ✓ **VALUE:** Two probiotic capsules per week benefit the microbiome.
- ✓ **TECHNOLOGY:** Sixteen U.S. Patents ensure effectiveness.
- ✓ **QUALITY:** Online test results for every bottle.
- ✓ **RESULTS:** Fast-acting digestive support.*

Call your *Life Extension*® Wellness Specialist to learn more.

Call Life Extension to place your order today.
www.LifeExtension.com

1-800-544-4440



30 capsules
Item# 01038
Retail: \$47.95
Your Price: \$35.96



32 capsules
Item# 01389
Retail: \$42.95
Your Price: \$32.21



6.2 OZ
Item# 01386
Retail: \$32.95
Your Price: \$24.71

*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.



master-supplements.com
Master Supplements
INCORPORATED

2016 PROSTATE CANCER CONFERENCE

The #1 Conference for Patients

*a collaborative
learning experience for*
Patients + Caregivers



**Los Angeles Airport Marriott
September 9-11, 2016**

[Featuring]

- Presentations from practicing physicians and leading clinical researchers on new developments, treatments, pharmaceuticals, and clinical trials for every stage of prostate cancer
- Presentations on reducing treatment side effects and maximizing quality of life
- Interaction with Experts in extensive Q+A and “Ask The Experts” breakout sessions
- Support groups
- Informational sessions for caregivers
- Exhibit hall featuring resources and products from our industry partners
- Explore Los Angeles by signing up for our Malibu Winery excursion

[Topics]

- Adjuvant Chemotherapy for High-Risk PC
- Focal Therapy for Early Stage PC
- Correcting Urinary Incontinence
- Body Scans for the Detection of Early Metastases
- Diet & Nutrition
- Xtandi, Zytiga & Xofigo

Registration and more info at:
www.pcri.org/2016-conference
info@pcri.org | 800.641.7274

Registration is \$120
Use code LIFEEXTENSION for 50% discount!



THE WELLNESS CENTER AT

POST HASTE

COMPOUNDING PHARMACY

Quality | Reliability | Integrity

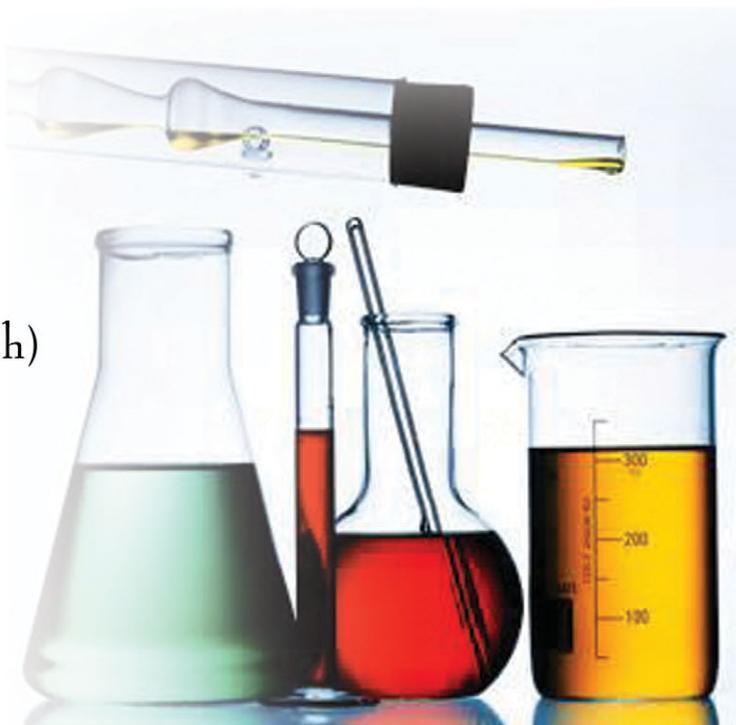
Lowest Prices: Ask for Customer Discount Pricing

1-866-892-3432

Providing Trusted Prescription Compounding for Over 40 Years

- * Full Compounding Lab
- * Full Retail Pharmacy
- * Bio-identical Hormone Replacement Therapy
- * Free Standard Delivery/Shipping
- * Durable Medical Equipment
- * Trilingual (English, Spanish, French)
- * Liscensed to ship into 42 States

Did you know we carry a full line of PET products? Mention this AD and recieve 10% off your first order!



Proud members of PCCA &



IACP
INTERNATIONAL ACADEMY OF
COMPOUNDING PHARMACISTS

PH:877-877-9700 FAX:866-892-3432

Renew Rx's online to receive \$1 off each Rx
(cash RX's only)

4401 Sheridan St. | Hollywood, Fl 33021

www.POSTHASTEPHARMACY.com



Blood testing provides the ultimate information regarding correctable risk factors that may predispose you to disorders such as cancer, diabetes, cardiovascular disease, and more. Information about general health and nutritional status can also be gained through standard blood analysis. Standing behind the belief that blood testing is an essential component of any program designed to attain optimal health and longevity, **Life Extension®** offers this innovative and convenient service at a very affordable price. Not only is comprehensive blood testing an important step in safeguarding your health, it is a simple process from virtually anywhere in the United States.

Five Easy Steps:

1. Call 1-800-208-3444 to discuss and place your order with one of our knowledgeable Wellness Specialists. (This order form can also be faxed to 1-866-728-1050 or mailed). Online orders can also be placed at www.lifeextension.com.
2. After your order is placed, you will be mailed either a requisition form to take to your local LabCorp Patient Service Center or a Blood Draw Kit; whichever is applicable (Please note: If a blood draw kit is used, an additional local draw fee may be incurred.)
3. Have your blood drawn.
4. Your blood test results will be sent directly to you by Life Extension.
5. Take the opportunity to discuss the results with one of our knowledgeable wellness specialists by calling 1-800-226-2370; or review the results with your personal physician.

It's that simple! Don't delay—call today!

For Our Local Customers:

For those residing in the Ft. Lauderdale, Florida, area, blood-draws are also performed at the Life Extension Nutrition Center from 9:00 am to 2:00 pm Monday through Saturday. Simply purchase the blood test and have it drawn with no wait! Our address is 5990 North Federal Highway, Ft. Lauderdale, FL, 33308-2633.

Blood Testing The Ultimate Information

MOST POPULAR PANELS

- | | |
|--|--|
| <p>COMPREHENSIVE PANELS</p> <p>MALE LIFE EXTENSION PANEL (LC322582) \$269
CBC/Chemistry Profile (See description)
C-Reactive Protein Homocysteine
DHEA-S Free Testosterone
TSH for thyroid function Total Testosterone
Estradiol PSA (prostate-specific antigen)
Vitamin D 25- hydroxy Hemoglobin A1c</p> | |
| <p>FEMALE LIFE EXTENSION PANEL (LC322535) \$269
CBC/Chemistry Profile (See description)
C-Reactive Protein Homocysteine
DHEA-S Free Testosterone
TSH for thyroid function Total Testosterone
Estradiol Vitamin D 25-hydroxy
Progesterone Hemoglobin A1c</p> | |
| <p>FEMALE HORMONE REPLACEMENT PANEL (LC100023) \$189
CBC/Chemistry Profile (see description), Estradiol, Estrone, Free and Total Testosterone, DHEA-S, Progesterone, TSH, and Insulin</p> | |
| <p>WEIGHT LOSS PANEL-COMPREHENSIVE (LC100028) \$275
CBC/Chemistry profile (see description), DHEA-S, Free and Total Testosterone, Estradiol, Progesterone, Cortisol, TSH, Free T3, Free T4, Reverse T3, Insulin, Hemoglobin A1c, Vitamin D 25-hydroxy, C-Reactive Protein (high sensitivity), and Ferritin.</p> | |
| <p>MALE ELITE PANEL* (LC100016) \$575
CBC/Chemistry Profile (See description), Free and Total Testosterone, Total Estrogens, Estradiol, DHEA-S, Progesterone, Pregnenolone, DHT, FSH, LH, TSH, Free T3, Free T4, Reverse T3, Free and Total PSA, IGF-1, SHBG, Vitamin D 25-OH, hs-CRP, Ferritin, Homocysteine</p> | |
| <p>FEMALE ELITE PANEL* (LC100017) \$575
CBC/Chemistry Profile (See description), Free and Total Testosterone, Total Estrogens, Estradiol, Estrone, DHEA-S, Progesterone, Pregnenolone, DHT, FSH, LH, TSH, Free T3, Free T4, Reverse T3, IGF-1, SHBG, Vitamin D 25-OH, hs-CRP, Ferritin, Homocysteine</p> | |
| <p>MALE HORMONE ADD-ON PANEL (LCADDM)* Pregnenolone and Dihydrotestosterone (DHT) \$120
To provide an even more in-depth analysis of a man's hormone status, Life Extension has created this panel as an addition to the Male Life Extension Panel.</p> | |
| <p>FEMALE HORMONE ADD-ON PANEL (LCADDF)* Pregnenolone and Total Estrogens \$125
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.</p> | |
| <p>COMPREHENSIVE THYROID PANEL (LC100018) \$199
TSH, T4, Free T4, Free T3, Reverse T3, TPO, ATA</p> | |
| <p>LIFE EXTENSION THYROID PANEL (LC304131) \$75
TSH, T4, Free T3, Free T4.</p> | |
| <p>THYROID PANEL WITH REVERSE T3 (LC100044) \$120
TSH, T4, Free T3, Free T4, Reverse T3</p> | |
| <p>THE CBC/CHEMISTRY PROFILE (LC381822) \$35
Note: This CBC/Chemistry Profile is included in many Life Extension panels. Please check panel descriptions.</p> <p>CARDIOVASCULAR RISK PROFILE
Total Cholesterol Cholesterol/HDL Ratio
HDL Cholesterol Estimated CHD Risk
LDL Cholesterol Glucose
Triglycerides Iron</p> <p>LIVER FUNCTION PANEL
AST (SGOT) Total Bilirubin
ALT (SGPT) Alkaline Phosphatase
LDH</p> <p>KIDNEY FUNCTION PANEL
BUN BUN/Creatinine Ratio
Creatinine Uric Acid</p> <p>BLOOD PROTEIN LEVELS
Total Protein Globulin
Albumin Albumin/Globulin Ratio</p> <p>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</p> <p>BLOOD MINERAL PANEL
Calcium Sodium
Potassium Chloride
Phosphorus Iron</p> | |
| <p>MALE COMPREHENSIVE HORMONE PANEL* (LC100010) \$299
CBC/Chemistry Profile, DHEA-S, Estradiol, DHT, PSA, Pregnenolone, Total and Free Testosterone, SHBG, TSH, Free T3, Free T4, Cortisol.</p> | |
| <p>FEMALE COMPREHENSIVE HORMONE PANEL* (LC100011) \$299
CBC/Chemistry Profile, DHEA-S, Estradiol, Total Estrogens, Progesterone, Pregnenolone, Total and Free Testosterone, SHBG, TSH, Free T3, Free T4, Cortisol.</p> | |
| <p>FOOD SAFE ALLERGY TEST** (LCM73001) \$198
This test measures delayed (IgG) food allergies for 95 common foods.</p> | |
| <p>STRESS MANAGEMENT PROFILE (LC100043) \$125
Cortisol AM/PM, DHEA-S, Glucose, Insulin, Progesterone, Free T3, Lipid Panel.</p> | |
| <p>ADRENAL STRESS PROFILE-SALIVA** (LC100046) \$175
Cortisol X4, DHEA-S, Cortisol AM/DHEA-S ratio, Secretory IgA.</p> | |
| <p>BASIC CORTISOL PROFILE-SALIVA** (LC100047) \$129
Cortisol X4 to measure cortisol rhythm over time.</p> | |
| <p>SLEEP HORMONES PROFILE-SALIVA** (LC100048) \$175
Cortisol and Melatonin plus ratio.</p> | |
| <p>MTHFR/COMT GENETIC METHYLATION PROFILE** (LC100045) \$149
Tests for genetic mutations in MTHFR and COMT.</p> | |



Other Popular Tests and Panels

- | | |
|--|--|
| <p><input type="radio"/> HEALTHY AGING PANEL-COMPREHENSIVE* (LC100026) \$249
CBC/Chemistry profile, C-Reactive Protein (high sensitivity), Vitamin B12, Folate, Homocysteine, Vitamin D 25-hydroxy, Hemoglobin A1c, TSH, Free T3, Free T4, Ferritin, Urinalysis, Fibrinogen, and Insulin.</p> <p><input type="radio"/> HEALTHY AGING PANEL-BASIC* (LC100025) \$149
CBC/Chemistry profile, C-Reactive Protein (high sensitivity), Vitamin B12, Folate, Vitamin D 25-hydroxy, Hemoglobin A1c, TSH, Ferritin, and Insulin.</p> <p><input type="radio"/> NMR LIPOPROFILE® (LC123810) \$99
The NMR LipoProfile® directly measures LDL particle size and number as well as HDL particle number, total cholesterol, and triglycerides. It also provides a calculation of one's risk of insulin resistance by assessing abnormalities in lipoprotein markers.</p> <p><input type="radio"/> ANEMIA PANEL* (LC100006) \$79
CBC/Chemistry Profile, Ferritin, Total Iron Binding Capacity (TIBC), Vitamin B12, Folate</p> <p><input type="radio"/> AUTOIMMUNE DISEASE SCREEN* (L100041) \$199
ANA screen, hs-CRP, TNF-alpha, Immunoglobulins, IgA, IgG, IgM</p> <p><input type="radio"/> DIABETES MANAGEMENT PROFILE – COMPREHENSIVE (LC100040) \$129
Hemoglobin A1C, Glucose, Insulin, Lipid Panel, Glycomark</p> <p><input type="radio"/> DIABETES MANAGEMENT PROFILE – BASIC (LC100039) \$39
Hemoglobin A1C, Glucose, Insulin</p> <p><input type="radio"/> ADVANCED CARDIAC BIOMARKERS
ADVANCED OXIDIZED LDL PANEL* (LC100035) \$285
This panel looks at vascular inflammatory biomarkers, beginning with lifestyle choices to the development of metabolic as well as cardiovascular disease and the formation of vulnerable plaque. The panel contains the following tests: F2-Isoprostanes, Myeloperoxidase, and Oxidized LDL.</p> <p><input type="radio"/> OXIDIZED LDL PANEL* (LC100034) \$175
This panel looks at vascular inflammatory biomarkers, beginning with the development of metabolic as well as cardiovascular disease and the formation of vulnerable plaque. The panel contains the following tests: Myeloperoxidase and Oxidized LDL.</p> <p><input type="radio"/> OMEGA CHECK™ (LCOMEGA) \$131.50
Provides valuable information on your risk of developing heart disease, sudden heart attack and cardiac death. The Omega Check™ also includes your AA:EPA ration allowing you to determine and track a major factor in total body inflammation.</p> | <p><input type="radio"/> HORMONES
DHEA-SULFATE (LC004020) \$61
This test shows if you are taking the proper amount of DHEA.</p> <p><input type="radio"/> MALE BASIC HORMONE PANEL (LC100012) \$75
DHEA-S, Estradiol, Free and Total Testosterone, PSA</p> <p><input type="radio"/> FEMALE BASIC HORMONE PANEL (LC100013) \$75
DHEA-S, Estradiol, Free and Total Testosterone, Progesterone</p> <p><input type="radio"/> DIHYDROTESTOSTERONE (DHT)* (LC500142) \$50
Measures serum concentrations of DHT.</p> <p><input type="radio"/> ESTRADIOL (LC004515) \$33
For men and women. Determines the proper amount in the body.</p> <p><input type="radio"/> INSULIN FASTING (LC004333) \$29.90
Can predict those at risk of diabetes, obesity, heart and other diseases.</p> <p><input type="radio"/> PREGNENOLONE* (LC140707) \$116
Used to determine ovarian failure, hirsutism, adrenal carcinoma, and Cushing's syndrome.</p> <p><input type="radio"/> PROGESTERONE (LC004317) \$55
Primarily for women. Determines the proper amount in the body.</p> <p><input type="radio"/> SEX HORMONE BINDING GLOBULIN (SHBG) (LC082016) \$33
This test is used to monitor SHBG levels which are under the positive control of estrogens and thyroid hormones, and suppressed by androgens.</p> <p><input type="radio"/> GENERAL HEALTH
VITAMIN D (25OH) (LC081950) \$47
This test is used to rule out vitamin D deficiency.</p> <p><input type="radio"/> FERRITIN (LC004598) \$28
Ferritin levels reflect your body's iron stores and is also a biomarker for insulin resistance.</p> <p><input type="radio"/> PSA (PROSTATE SPECIFIC ANTIGEN) (LC010322) \$31
Screening test for prostate disorders and possible cancer.</p> |
|--|--|

Blood tests available in the continental United States only. Restrictions apply in NY, NJ, RI, and MA. Not available in Maryland. Kits not available in Pennsylvania.

This is NOT a complete listing of LE blood test services. Call 1-800-208-3444 for additional information.

ORDER LIFESAVING 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 Life Extension® 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 Life Extension® 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 _____

X _____

CUSTOMER NO. _____

Male

Female

Name _____

Date of Birth (required) / /

Address _____

City _____

State _____

Zip _____

Phone _____

Credit Card No. _____

Expiration Date /

Mail your order form to:

LifeExtension®
National Diagnostics, Inc.

3600 West Commercial Boulevard
Fort Lauderdale, FL 33309

Phone your order to: 1-800-208-3444

Fax your order to: 1-866-728-1050



With Your Healthy Rewards, you earn LE Dollars back on every purchase you make — including blood tests!

See www.LifeExtension.com/Rewards for details.

* 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.

** This test is packaged as a kit.

Amino Acids

Arginine/L-Ornithine Capsules
 Arginine Ornithine Powder
 Branched Chain Amino Acids
 D,L-Phenylalanine Capsules
 L-Arginine Caps
 L-Carnitine
 L-Glutamine
 L-Glutamine Powder
 L-Lysine
 L-Taurine Powder
 L-Tyrosine Powder
 Super Carnosine
 Taurine

Blood Pressure & Vascular Support

Advanced Olive Leaf Vascular Support with Celery Seed Extract
 Arterial Protect
 Blood Pressure Monitor Arm Cuff
 Endothelial Defense™ with Pomegranate Complete and CORDIART™
 Endothelial Defense™ with GliSODin®
 Natural BP Management
 NitroVasc with CORDIART™
 Pomegranate Complete
 Pomegranate Fruit Extract

Bone Health

Bone Restore
 Bone Restore with Vitamin K2
 Bone Strength Formula with KoAct®
 Bone-Up™
 Calcium Citrate with Vitamin D
 Dr. Strum's Intensive Bone Formula
 Strontium Caps

Brain Health

Acetyl-L-Carnitine
 Acetyl-L-Carnitine Arginate
 Blast
 Brain Shield® Gastrodin
 Cognitex® Basics
 Cognitex® with Brain Shield®
 Cognitex® with Pregnenolone & Brain Shield®
 Cognizin® CDP-Choline Caps
 DMAE Bitartrate (dimethylaminoethanol)
 Dopa-Mind™
 Ginkgo Biloba Certified Extract™
 Huperzine A
 Lecithin Granules
 Migra-Eeze™
 Neuro-Mag® Magnesium L-Threonate
 Neuro-Mag® Magnesium L-Threonate with Calcium and Vitamin D3
 Optimized Ashwagandha Extract
 Prevagen™
 PS (Phosphatidylserine) Caps
 Vinpocetine

Cholesterol Management

Advanced Lipid Control
 Cho-Less™
 CHOL-Support™
 Red Yeast Rice
 Theaflavins Standardized Extract
 Vitamin B3 Niacin Capsules

Digestion Support

Artichoke Leaf Extract
 Carnosoothe with PicroProtect™
 Digest RC®
 Effervescent Vitamin C - Magnesium Crystals
 Enhanced Super Digestive Enzymes
 Enhanced Super Digestive Enzymes w/Probiotics
 Esophageal Guardian
 Extraordinary Enzymes
 Fem Dophilus
 Fiber-Immune Support

Ginger Force®
 Organic Golden Flax Seed
 Pancreatin
 Regimint
 Tranquil Tract™
 TruFiber™
 WellBetX PGX plus Mulberry

Energy Management

Adrenal Energy Formula
 Asian Energy Boost
 D-Ribose Powder
 D-Ribose Tablets
 Forskolin
 Mitochondrial Basics with BioPQQ®
 Mitochondrial Energy Optimizer with BioPQQ®
 NAD+ Cell Regenerator™
 Peak ATP® with GlycoCam®
 PQQ Caps with BioPQQ®
 Rhodiola Extract
 RiboGen™ French Oak Wood Extract
 Triple Action Thyroid

Eye Health

Astaxanthin with Phospholipids
 Brite Eyes III
 Certified European Bilberry Extract
 Eye Pressure Support with Mirtogenol®
 MacuGuard® Ocular Support
 MacuGuard® Ocular Support with Astaxanthin
 Tear Support with MaquiBright®

Fish Oil & Omegas

OMEGA FOUNDATIONS® Mega EPA/DHA
 OMEGA FOUNDATIONS® Mega GLA with Sesame Lignans
 OMEGA FOUNDATIONS® Super Omega-3 EPA/DHA with Sesame Lignans & Olive Extract
 OMEGA FOUNDATIONS® Super Omega-3 Plus EPA/DHA with Sesame Lignans, Olive Extract, Krill & Astaxanthin
 Organic Golden Flax Seed
 OMEGA FOUNDATIONS® Provinal® Purified Omega-7
 OMEGA FOUNDATIONS® Vegetarian DHA

Food

California Estate Extra Virgin Olive Oil
 Rich Rewards® Breakfast Blend
 Rich Rewards® Breakfast Blend Natural Mocha Flavor
 Rich Rewards® Breakfast Blend Natural Vanilla Flavor
 Rich Rewards® Breakfast Blend Whole Bean Coffee
 Rich Rewards® Decaf Roast
 Stevia Sweetener

Glucose Management

CinSulin® with InSea2® and Crominex® 3+
 Mega Benfotiamine
 Natural Glucose Absorption Control
 Tri Sugar Shield®

Heart Health

Aspirin (Enteric Coated)
 BioActive Folate & Vitamin B12 Caps
 Cardio Peak™ with Standardized Hawthorn and Arjuna
 Fibrinogen Resist™ with Nattokinase
 Optimized Carnitine with GlycoCam®
 Super Ubiquinol CoQ10
 Super Ubiquinol CoQ10 with BioPQQ®
 Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™
 Super-Absorbable CoQ10 Ubiquinone with α-Limonene
 TMG Powder
 TMG Liquid Capsules

Hormone Balance

DHEA (Dehydroepiandrosterone)
 Inner Power
 Pregnenolone
 Triple Action Cruciferous Vegetable Extract with Resveratrol
 Triple Action Cruciferous Vegetable Extract

Immune Support

AHCC®
 Echinacea Extract
 Enhanced Zinc Lozenges
 i26 Hyperimmune Egg
 Immune Modulator with Tinofend®
 Immune Protect with PARACTIN®
 Immune Senescence Protection Formula™
 Kinoko® Gold AHCC
 Kyolic® Garlic Formula 102
 Kyolic® Garlic Formula 105
 Kyolic® Reserve
 Lactoferrin (apolactoferrin) Caps
 NK Cell Activator™
 Optimized Garlic
 Optimized Quercetin
 Peony Immune
 ProBoost Thymic Protein A
 Reishi Extract Mushroom Complex
 Standardized Cistanche
 Ten Mushroom Formula®
 Zinc Lozenges

Inflammation Management

5-LOX Inhibitor with AprèsFlex®
 Advanced Bio-Curcumin® with Ginger & Turmerones
 Black Cumin Seed Oil with Bio-Curcumin®
 Black Cumin Seed Oil
 Boswellia
 Cytokine Suppress™ with EGCG
 Nervia®
 Serraflazyme
 Specially-Coated Bromelain
 Super Bio-Curcumin®
 Zylamend® Whole Body

Joint Support

Arthro-Immune Joint Support
 ArthroMax® Advanced with UC-II® & AprèsFlex®
 ArthroMax® with Theaflavins & AprèsFlex®
 Bio-Collagen with Patented UC-II®
 Fast-Acting Joint Formula
 Glucosamine/Chondroitin Capsules
 Krill Healthy Joint Formula
 MSM (Methylsulfonylmethane)

Kidney & Bladder Support

Cran-Max® Cranberry Whole Fruit Concentrate
 Optimized Cran-Max® with Ellirose™
 Uric Acid Control
 Water-Soluble Pumpkin Seed Extract

Liver Health & Detoxification

Anti-Alcohol Antioxidants with HepatoProtection Complex
 Calcium D-Glucarate
 Chlorella
 Chlorophyllin
 European Milk Thistle
 Glutathione, Cysteine & C
 HepatoPro
 Liver Efficiency Formula
 N-Acetyl-L-Cysteine
 PectaSol-C®
 Silymarin
 SODzyme® with GliSODin® & Wolfberry

Longevity & Wellness

AMPK Activator
 AppleWise Polyphenol Extract
 Berry Complete
 Blueberry Extract
 Blueberry Extract with Pomegranate

CR Mimetic Longevity Formula
 DNA Protection Formula
 Enhanced Berry Complete with Acai
 Essential Daily Nutrients
 Grapeseed Extract with Resveratrol & Pterostilbene
 Mega Green Tea Extract (decaffeinated)
 Mega Green Tea Extract (lightly caffeinated)
 Optimized Fucoidan with Maritech® 926
 Optimized Resveratrol
 Optimized Resveratrol with Nicotinamide Riboside
 pTeroPure®
 Pycnogenol® French Maritime Pine Bark Extract
 Resveratrol with Pterostilbene
 RNA (Ribonucleic Acid)
 Super Alpha-Lipoic Acid
 Super R-Lipoic Acid
 X-R Shield

Men's Health

Mega Lycopene Extract
 PalmettoGuard® Saw Palmetto with Beta-Sitosterol
 PalmettoGuard® Saw Palmetto/Nettle Root Formula with Beta-Sitosterol
 Prelox® Natural Sex for Men®
 Super MiraForte with Standardized Lignans
 Triple Strength ProstaPollen™
 Ultra Natural Prostate

Minerals

Boron
 Iron Protein Plus
 Magnesium (Citrate)
 Magnesium Caps
 Only Trace Minerals
 Optimized Chromium with Crominex® 3+ Sea-Iodine™
 Se-Methyl L-Selenocysteine
 Super Selenium Complex
 Vanadyl Sulfate
 Zinc Caps

Miscellaneous

Solarshield® Sunglasses

Mood & Stress Management

5 HTP
 L-Theanine
 Natural Stress Relief
 SAME (S-Adenosyl-Methionine)

Multivitamins

Children's Formula Life Extension Mix™
 Comprehensive Nutrient Packs ADVANCED
 Life Extension Mix™ Capsules without Copper
 Life Extension Mix™ Capsules
 Life Extension Mix™ Powder without Copper
 Life Extension Mix™ Powder
 Life Extension Mix™ Tablets with Extra Niacin
 Life Extension Mix™ Tablets without Copper
 Life Extension Mix™ Tablets
 Once-Daily Health Booster
 One-Per-Day Tablets
 Two-Per-Day Capsules
 Two-Per-Day Tablets

Personal Care

Anti-Aging Rejuvenating Scalp Serum
 Biosil
 Dr. Proctor's Advanced Hair Formula
 Dr. Proctor's Shampoo
 European Leg Solution Featuring Certified Diosmin 95
 Face Master Platinum
 Facial Toning System
 Hair, Skin & Nail Rejuvenation Formula w/VERISOL®
 Hair Suppress Formula
 Life Extension Toothpaste

Sinus Cleanser
 Venotone
 Xylwhite Mouthwash

Pet Care

Cat Mix
 Dog Mix

Probiotics

Bifido GI Balance
 BroccoMax®
 FLORASSIST® Heart Health
 FLORASSIST® Oral Hygiene
 FLORASSIST® Balance
 FLORASSIST® Mood
 FLORASSIST® Throat Health
 Theralac® Probiotics
 TruFlora® Probiotics

Skin Care

Advanced Anti-Glycation Peptide Serum
 Advanced Lightening Cream
 Advanced Peptide Hand Therapy
 Advanced Triple Peptide Serum
 Advanced Under Eye Serum with Stem Cells
 Amber Self MicroDermAbrasion
 Anti-Aging Face Oil
 Anti-Aging Mask
 Anti-Aging Rejuvenating Face Cream
 Anti-Glycation Serum with Blueberry & Pomegranate Extracts
 Antioxidant Facial Mist
 Anti-Oxidant Rejuvenating Foot Cream
 Anti-Oxidant Rejuvenating Foot Scrub
 Anti-Oxidant Rejuvenating Hand Cream
 Anti-Redness & Adult Blemish Lotion
 Bioflavonoid Cream
 Broccoli Sprout Cream
 Collagen Boosting Peptide Serum
 Corrective Clearing Mask
 DNA Repair Cream
 Essential Plant Lipids Reparative Serum
 Face Rejuvenating Anti-Oxidant Cream
 Fine Line-Less
 Healing Formula
 Healing Mask
 Healing Vitamin K Cream
 Hyaluronic Facial Moisturizer
 Hyaluronic Oil-Free Facial Moisturizer
 Hydrating Anti-Oxidant Facial Mist
 Hydroderm
 Lifting & Tightening Complex
 Lycopene Cream
 Melatonin Cream
 Mild Facial Cleanser
 Multi Stem Cell Skin Tightening Complex
 Neck Rejuvenating Anti-Oxidant Cream
 Pigment Correcting Cream
 Rejuvenating Serum
 Rejuvenex® Body Lotion
 Rejuvenex® Factor Firming Serum
 Renewing Eye Cream
 Resveratrol Anti-Oxidant Serum
 Shade Factor
 Skin Lightening Serum
 Skin Restoring Phytoceramides with Lipowheat®
 Skin Stem Cell Serum
 Stem Cell Cream with Alpine Rose
 Tightening & Firming Neck Cream
 Triple-Action Vitamin C Cream
 Ultimate MicroDermabrasion
 Ultra Eyelash Booster
 Ultra Lip Plumper
 Ultra Rejuvenex®
 Ultra RejuveNight®
 Ultra Wrinkle Relaxer
 Under Eye Refining Serum
 Under Eye Rescue Cream
 Vitamin C Serum
 Vitamin D Lotion
 Vitamin E-ssential Cream
 Youth Serum

Sleep

Bioactive Milk Peptides
 Enhanced Natural Sleep® with Melatonin
 Enhanced Natural Sleep® without Melatonin
 Fast-Acting Liquid Melatonin
 Glycine
 L-Tryptophan
 Melatonin
 Optimized Tryptophan Plus

Sports Performance

Creatine Capsules
 Creatine Whey Glutamine Powder (Vanilla Flavor)
 New Zealand Whey Protein Concentrate (Natural Chocolate and Vanilla Flavor)
 Tart Cherry Extract
 Whey Protein Isolate (Chocolate and Vanilla Flavor)

Vitamins

Ascorbyl Palmitate
 Benfotiamine with Thiamine
 Beta-Carotene
 BioActive Complete B-Complex
 Biotin
 Buffered Vitamin C Powder
 Daily C+
 Fast-C® with Dihydroquercetin
 Gamma E Tocopherol with Sesame Lignans
 Gamma E Tocopherol/Tocotrienols
 High Potency Optimized Folate
 Inositol Caps
 Liquid Emulsified Vitamin D3
 Liquid Vitamin D3
 Low-Dose Vitamin K2
 Methylcobalamin
 MK-7
 Natural Vitamin E
 No Flush Niacin
 Optimized Folate (L-Methylfolate)
 Pantothenic Acid (Vitamin B-5)
 Pyridoxal 5'-Phosphate Caps
 Super Absorbable Tocotrienols
 Super Ascorbate C Capsules
 Super Ascorbate C Powder
 Super K with Advanced K2 Complex
 Vitamin B12
 Vitamin B6
 Vitamin C with Dihydroquercetin
 Vitamin D3 with Sea-Iodine™
 Vitamin D3
 Vitamins D and K with Sea-Iodine™

Weight Management

7-Keto® DHEA Metabolite
 Advanced Anti-Adipocyte Formula
 Advanced Natural Appetite Suppress
 CalReduce Selective Fat Binder
 DHEA Complete
 Garcinia HCA
 HCAActive™ Garcinia Cambogia Extract
 Integra-Lean®
 Mediterranean Trim with Sinetrol™-XPur
 Optimized Irvingia with Phase 3™ Calorie Control Complex
 Optimized Saffron with Satiereal®
 Super Citrimax®
 Super CLA Blend with Guarana and Sesame Lignans
 Super CLA Blend with Sesame Lignans
 Waist-Line Control™

Women's Health

Advanced Natural Sex for Women® 50+
 Breast Health Formula
 Femmenessence MacaPause®
 Natural Estrogen
 Progesta-Care®
 Super-Absorbable Soy Isoflavones
 Ultra Soy Extract

ITEM No.	PRODUCT	YOUR PRICE			QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each		
A						
01524	ACETYL-L-CARNITINE • 500 mg, 100 veg. caps	34.00	25.50	22.50		
01525	ACETYL-L-CARNITINE ARGINATE • 100 veg. caps	59.00	44.25	38.24		
01628	ADRENAL ENERGY FORMULA • 60 veg. caps	24.00	18.00	16.50		
01630	ADRENAL ENERGY FORMULA • 120 veg. caps	46.00	34.50	31.50		
01828	ADVANCED LIPID CONTROL • 60 veg. caps	30.00	22.50	20.25		
00681	AHCC® • 500 mg, 30 caps	59.98	44.99			
29727	AHCC® (KINOKO® GOLD) • 500 mg, 60 veg. caps	74.95	52.47			
00457	ALPHA-LIPOIC ACID W/BIOTIN (Super) • 250 mg, 60 caps	37.00	27.75	24.00		
01907	AMPK ACTIVATOR • 90 veg. caps	48.00	36.00	33.00		
01440	ANTI-ALCOHOL ANTIOXIDANTS W/HEPATOPRO • 100 caps	26.00	19.50	17.25		
01509	ANTI-ADIPOCYTE FORMULA W/MERATRIM® & INTEGRA LEAN® (Advanced) • 60 veg. caps	39.00	29.25	27.00		
01625	APPLEWISE POLYPHENOL EXTRACT 600 mg, 30 veg. caps	21.00	15.75	14.25		
01039	ARGININE/ORNITHINE • 500/250, 100 caps	17.99	13.49			
00038	ARGININE/ORNITHINE POWDER • 150 grams	22.95	17.21	14.25		
01624	(L)-ARGININE CAPS • 700 mg, 200 veg. caps	26.50	19.88	17.44		
02004	ARTERIAL PROTECT • 30 veg. caps	48.00	36.00	33.00		
01617	ARTHROMAX® W/THEAFLAVINS & APRÈSFLEX® 120 veg. caps	44.00	33.00	30.00		
01618	ARTHROMAX® ADVANCED W/UC-II® & APRÈSFLEX® 60 caps	36.00	27.00	24.00		
01404	ARTHO-IMMUNE JOINT SUPPORT • 60 veg. caps	32.00	24.00	21.00		
00919	ARTICHOKE LEAF EXTRACT • 500 mg, 180 veg. caps	30.00	22.50	21.00		
01533	ASCORBYL PALMITATE • 500 mg, 100 veg. caps	22.50	16.88	15.00		
00888	ASHWAGANDHA EXTRACT (Optimized) • 60 veg. caps	10.00	7.50	6.75		
01805	ASIAN ENERGY BOOST • 90 veg. caps	24.00	18.00	16.50		
01066	ASPIRIN • 81 mg, 300 enteric coated tablets	6.00	4.50	4.00		
01923	ASTAXANTHIN WITH PHOSPHOLIPIDS • 4 mg, 30 softgels	16.00	12.00	10.50		
B						
00920	BENFOTIAMINE W/ THIAMINE • 100 mg, 120 veg. caps	19.95	14.96	13.95		
00925	BENFOTIAMINE (Mega) • 250 mg, 120 veg. caps	30.00	22.50	20.25		
01206	BERRY COMPLETE • 30 veg. caps	21.00	15.75	14.00		
01496	BERRY COMPLETE W/ACAI (Enhanced) • 60 veg. caps	29.00	21.75	19.50		
00664	BETA-CAROTENE • 25,000 IU, 100 softgels	11.25	8.44			
01622	BIFIDO GI BALANCE • 60 veg. caps	20.00	15.00	13.50		
01873	BILBERRY EXTRACT • 100 mg, 90 veg. caps	36.00	27.00	24.00		
01512	BIOACTIVE MILK PEPTIDES • 30 caps	18.00	13.50	12.00		
01631	BIO-COLLAGEN W/PATENTED UC-II® • 40 mg, 60 small caps	36.00	27.00	24.00		
*01006	BIOSIL™ • 5 mg, 30 veg. caps	18.95	15.16			
*01007	BIOSIL™ • 1 fl oz	31.99	25.59			
00102	BIOTIN • 600 mcg, 100 caps	7.50	5.63	4.88		
01709	BLACK CUMIN SEED OIL • 60 softgels	16.00	12.00	10.50		
01710	BLACK CUMIN SEED OIL W/BIO-CURCUMIN® • 60 softgels	32.00	24.00	22.50		
01008	BLAST™ • 600 grams of powder	26.95	20.21			
70000	BLOOD PRESSURE MONITOR (ACCUFIT™) • med/lq cuff	79.99	49.99			
70004	BLOOD PRESSURE MONITOR • Digital wrist cuff	69.95	52.46			
SUBTOTAL OF COLUMN 1						

ITEM No.	PRODUCT	YOUR PRICE			QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each		
01214	BLUEBERRY EXTRACT • 60 veg. caps	22.50	16.88	15.00		
01438	BLUEBERRY EXTRACT W/ POMEGRANATE • 60 veg. caps	30.00	22.50	20.25		
01506	BONE FORMULA (DR. STRUM'S INTENSIVE) • 300 caps	56.00	42.00	37.50		
01726	BONE RESTORE • 120 caps	22.00	16.50	14.25		
01727	BONE RESTORE W/VITAMIN K2 • 120 caps	24.00	18.00	16.50		
01725	BONE STRENGTH FORMULA W/KOACT® • 120 caps	45.00	33.75	30.00		
00313	BONE-UP® • 240 caps	28.95	21.71	20.41		
01661	BORON • 3 mg, 100 veg. caps	5.95	4.46	3.94		
00202	BOSWELLA • 100 caps	38.00	28.50	22.50		
01802	BRAIN SHIELD® GASTRODIN • 300 mg, 60 veg. caps	33.00	24.75	22.50		
01253	BRANCHED CHAIN AMINO ACIDS • 90 caps	19.50	14.63	12.75		
01942	BREAST HEALTH FORMULA • 60 caps	34.00	25.50	22.50		
00893	BRITE EYES III • 2 vials, 5 ml each	34.00	25.50	24.00		
26576	BROCCO MAX® • 60 veg. caps	26.95	20.21			
01203	BROMELAIN (Specially-coated) 500 mg, 60 enteric coated tablets	21.00	15.75	14.25		
C						
01653	CALCIUM CITRATE W/VITAMIN D • 300 caps	24.00	18.00	15.94		
01651	CALCIUM D-GLUCARATE • 200 mg, 60 veg. caps	18.00	13.50	11.25		
*01823	CALREDUCE SELECTIVE FAT BINDER 120 mint chewable tablets	45.00	33.75	28.50		
01700	CARDIO PEAK™ W/STANDARDIZED HAWTHORN & ARJUNA 120 veg. caps	36.00	27.00	24.00		
00916	CARNITINE W/GLYCOCARN® (Optimized) • 60 veg. caps	36.00	27.00	24.00		
01532	L-CARNITINE • 500 mg, 30 veg. caps	15.00	11.25	9.90		
01258	CARNOSOOTHE W/PICROPROTECT™ • 60 veg. caps	30.00	22.50	20.25		
01829	CARNOSINE • 500 mg, 60 veg. caps	36.00	27.00	24.00		
01687	CARNOSINE (Super) • 500 mg, 90 veg. caps	66.00	49.50	45.00		
01932	CAT MIX • 100 grams powder	14.00	10.50	8.25		
01899	CHILDREN'S FORMULA LIFE EXTENSION MIX™ 100 chewable tablets	20.00	15.00	13.50		
00550	CHLORELLA • 500 mg, 200 tablets	23.50	17.63			
01571	CHLOROPHYLLIN • 100 mg, 100 veg. caps	24.00	18.00	15.00		
01359	CHO-LESS™ • 90 capsules	35.00	26.25			
01910	CHOL-SUPPORT™ • 60 liquid veg. caps	48.00	36.00	32.00		
01504	CHROMIUM W/CROMINEX® 3+ (Optimized) 500 mcg, 60 veg. caps	9.00	6.75	6.00		
01503	CINSULIN® W/INSEA2® AND CROMINEX® 3+ • 90 veg. caps	38.00	28.50	25.50		
01906	CISTANCHE (Standardized) • 30 veg. caps	20.00	15.00	12.00		
01818	CITRIMAX® (Super) • 180 veg. caps	40.00	30.00	28.50		
00818	CLA BLEND W/SESAME LIGNANS (Super) 1,000 mg, 120 softgels	36.00	27.00	24.75	19.75	
00819	CLA BLEND W/GUARANA & SESAME (Super) 1,000 mg, 120 softgels	42.00	31.50	28.75		
01896	COGNITEX® W/BRAIN SHIELD® • 90 softgels	60.00	45.00	39.00	36.00	
01897	COGNITEX® W/PREGNENOLONE & BRAIN SHIELD® 90 softgels	62.00	46.50	39.75	37.50	
01421	COGNITEX® BASICS • 60 softgels	38.00	28.50	26.25	24.00	
SUBTOTAL OF COLUMN 2						

ITEM No.	PRODUCT	Retail Each \$	YOUR PRICE			QTY	Total
			1 Unit Each	4 Unit Each	10 Unit Each		
01659	COGNIZIN® CDP CHOLINE CAPS • 250 mg, 60 veg. caps	36.00	27.00	25.50			
01945	COMPLETE B-COMPLEX (BioActive) • 60 veg. caps	12.00	9.00	8.00			
02098	COMPREHENSIVE NUTRIENT PACKS ADVANCED • 30 packs	90.00	67.50	61.50			
01949	COQ10 w/d-LIMONENE (Super-Absorbable) 50 mg, 60 softgels	25.00	18.75	16.50	15.00		
01948	COQ10 w/d-LIMONENE (Super-Absorbable) 100 mg, 100 softgels	46.00	34.50	28.00	26.25		
01929	COQ10 (Super ubiquinol) • 100 mg, 60 softgels	56.00	42.00	36.00	33.00		
01733	COQ10 w/BIOPIQQ® (Super ubiquinol) • 100 mg, 30 softgels	54.00	40.50	33.00	30.00		
01426	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) • 100 mg, 60 softgels	62.00	46.50	39.00	36.00		
01425	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) • 50 mg, 100 softgels	58.00	43.50	34.50	31.50		
01427	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) • 50 mg, 30 softgels	20.00	15.00	12.00			
01431	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) • 200 mg, 30 softgels	62.00	46.50	39.00	36.00		
00862	CRAN-MAX® • 500 mg, 60 veg. caps	17.50	13.13	11.25			
01424	CRAN-MAX® WITH ELLIROSE™ (Optimized) • 60 veg. caps	18.00	13.50	12.00			
01529	CREATINE CAPSULES • 120 veg. caps	10.95	8.21	6.94			
01746	CREATINE WHEY GLUTAMINE POWDER • 454 grams (vanilla)	30.00	22.50	20.25			
01429	CR MIMETIC LONGEVITY FORMULA • 60 veg. caps	39.00	29.25	27.00			
00407	CURCUMIN® (Super Bio) • 400 mg, 60 veg. caps	38.00	28.50	26.25			
01924	CURCUMIN® W/GINGER & TURMERONES (Advanced Bio) 30 softgels	30.00	22.50	20.25			
01804	CYTOKINE SUPPRESS™ W/EGCG • 30 veg. caps	30.00	22.50	20.25			
COSMESIS							
80157	ADVANCED ANTI-GLYCATION PEPTIDE SERUM • 1 oz	53.00	39.75	34.50			
80154	ADVANCED LIGHTENING CREAM • 1 oz	65.00	48.75	42.75			
80155	ADVANCED PEPTIDE HAND THERAPY • 4 oz	46.00	34.50	29.25			
80152	ADVANCED TRIPLE PEPTIDE SERUM • 1 oz	65.00	48.75	42.75			
80140	ADVANCED UNDER EYE SERUM W/STEM CELLS • .33 oz	49.00	36.75	31.50			
80139	AMBER SELF MICRODERMABRASION • 2 oz	49.00	36.75	31.50			
80158	ANTI-AGING FACE OIL • 1 oz	59.00	44.25	39.00			
80118	ANTI-AGING MASK • 2 oz	72.00	54.00	47.52			
80151	ANTI-AGING REJUVENATING FACE CREAM • 2 oz	65.00	48.75	42.75			
80153	ANTI-AGING REJUVENATING SCALP SERUM • 2 oz	46.00	34.50	29.25			
80134	ANTI-GLYCATION SERUM W/BLUEBERRY & POMEGRANATE EXTRACTS • 1 oz	33.00	24.75	23.51			
80133	ANTIOXIDANT FACIAL MIST • 2 oz	32.00	24.00	22.80			
80127	ANTIOXIDANT REJUVENATING FOOT CREAM • 2 oz	45.00	33.75	32.10			
80128	ANTIOXIDANT REJUVENATING FOOT SCRUB • 2 oz	59.00	44.25	38.94			
80117	ANTIOXIDANT REJUVENATING HAND CREAM • 2 oz	64.00	48.00	43.12			
80105	ANTI-REDNESS & ADULT BLEMISH LOTION • 1 oz	74.50	55.88	49.17			
80147	BIOFLAVONOID CREAM • 1 oz	46.00	34.50	29.25			
80144	BROCCOLI SPROUT CREAM • 1 oz	46.00	34.50	29.25			
80156	COLLAGEN BOOSTING PEPTIDE SERUM • 1 oz	59.00	44.25	39.00			
80120	CORRECTIVE CLEARING MASK • 2 oz	64.50	48.38	42.57			
80141	DNA REPAIR CREAM • 1 oz	49.00	36.75	31.50			
SUBTOTAL OF COLUMN 3							

RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

ITEM No.	PRODUCT	Retail Each \$	YOUR PRICE			QTY	Total
			1 Unit Each	4 Unit Each	10 Unit Each		
80108	ESSENTIAL PLANT LIPIDS REPARATIVE SERUM • 1 oz	74.95	56.21	49.46			
80123	FACE REJUVENATING ANTIOXIDANT CREAM • 2 oz	69.50	52.13	45.87			
80107	FINE LINE-LESS • 1 oz	74.50	55.88	49.17			
80131	HAIR SUPPRESS FORMULA • 4 oz	59.00	44.25	38.94			
80137	HEALING FORMULA ALL-IN-ONE CREAM • 1 oz	53.00	39.75	34.07			
80115	HEALING MASK • 2 oz	64.50	48.38	42.57			
80102	HEALING VITAMIN K CREAM • 1 oz	79.50	59.63	52.47			
80109	HYALURONIC FACIAL MOISTURIZER • 1 oz	58.00	43.50	38.28			
80110	HYALURONIC OIL-FREE FACIAL MOISTURIZER • 1 oz	58.00	43.50	38.28			
80138	HYDRATING ANTIOXIDANT FACE MIST • 4 oz	39.95	29.96	28.50			
80103	LIFTING & TIGHTENING COMPLEX • 1 oz	74.50	55.88	49.17			
80146	LYCOPENE CREAM • 1 oz	28.00	21.00	19.05			
80135	MELATONIN CREAM • 1 oz	33.00	24.75	20.33			
80114	MILD FACIAL CLEANSER • 8 fl. oz	59.00	44.25	38.94			
80159	MULTI STEM CELL SKIN TIGHTENING COMPLEX • 1 oz	59.00	44.25	39.00			
80122	NECK REJUVENATING ANTIOXIDANT CREAM • 2 oz	64.00	48.00	42.24			
80111	PIGMENT CORRECTING CREAM • 1/2 oz	74.00	55.50	48.84			
80106	REJUVENATING SERUM • 1 oz	74.50	55.88	49.17			
80150	RENEWING EYE CREAM • 1/2 oz	65.00	48.75	42.75			
80142	RESVERATROL ANTI-OXIDANT SERUM • 1 oz	46.00	34.50	29.25			
80112	SKIN LIGHTENING SERUM • 1/2 oz	85.00	63.75	56.10			
80130	SKIN STEM CELL SERUM • 1 oz	74.00	55.50	51.75			
80143	STEM CELL CREAM W/ALPINE ROSE • 1 oz	66.00	49.50	43.50			
80148	TIGHTENING & FIRMING NECK CREAM • 2 oz	39.00	29.25	26.25			
80161	TRIPLE ACTION VITAMIN C CREAM • 1 oz jar	59.00	44.25	39.00			
80162	ULTIMATE MICRODERMABRASION • 8 fl. oz	39.00	29.25	26.25			
80160	ULTRA EYELASH BOOSTER • 0.25 oz (2 units \$39)	59.00	44.25				
80116	ULTRA LIP PLUMPER • 1/3 oz	64.00	48.00	42.24			
80101	ULTRA WRINKLE RELAXER • 1 oz	89.95	67.46	59.82			
80113	UNDER EYE REFINING SERUM • 1/2 oz	74.50	55.88	49.17			
80104	UNDER EYE RESCUE CREAM • 1/2 oz	74.50	55.88	49.17			
80129	VITAMIN C SERUM • 1 oz	85.00	63.75	56.10			
80136	VITAMIN D LOTION • 4 oz	36.00	27.00	25.25			
80145	VITAMIN E-ESSENTIAL CREAM • 1 oz	28.00	21.00	19.50			
80149	YOUTH SERUM • 1 oz	65.00	48.75	42.75			
D							
01912	DAILY C+ CITRUS FLAVOR • 30 stick packs	21.00	15.75	14.25			
00658	7-KETO® DHEA METABOLITE • 25 mg, 100 caps	28.00	21.00	18.00			
01479	7-KETO® DHEA METABOLITE • 100 mg, 60 veg. caps	40.00	30.00	27.00			
01640	DHA (Vegetarian) • 30 veg. softgels	20.00	15.00	13.50			
00607	DHEA • 25 mg, 100 tablets (Dissolve in mouth)	14.00	10.50	8.81			
01478	DHEA COMPLETE • 60 veg. caps	48.00	36.00	32.40			
00335	DHEA • 25 mg, 100 caps	16.00	12.00	11.00			
00454	DHEA • 15 mg, 100 caps	14.00	10.50	9.00			
00882	DHEA • 50 mg, 60 caps	19.00	14.25	12.75			
SUBTOTAL OF COLUMN 4							

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
01689	DHEA • 100 mg, 60 veg. caps	24.00	18.00	16.50			
01358	DIGEST RC® • 30 tablets	19.95	14.96	12.75			
02021	DIGESTIVE ENZYMES (Enhanced Super) • 60 veg. caps	22.00	16.50	15.00			
02022	DIGESTIVE ENZYMES w/PROBIOTICS (Enhanced Super) • 60 veg. caps	28.00	21.00	18.00			
01671	D, L-PHENYLALANINE • 500 mg, 100 veg. caps	18.75	14.06	12.00			
01540	DMAE BITARTRATE • 150 mg, 200 veg. caps	18.00	13.50	11.25			
01570	DNA PROTECTION FORMULA • 60 veg. caps	34.00	25.50	24.00			
01931	DOG MIX • 100 grams powder	18.00	13.50	11.25			
02006	DOPA-MIND™ • 60 veg. tabs	48.00	36.00	32.00			
00321	DR. PROCTOR'S ADVANCED HAIR FORMULA • 2 oz	39.95	29.96	24.00			
00320	DR. PROCTOR'S HAIR SHAMPOO • 8 oz	24.95	18.71	16.50			
E							
01528	ECHINACEA EXTRACT • 250 mg, 60 veg. caps	14.35	10.76	9.38			
01997	ENDOTHELIAL DEFENSE™ w/POMEGRANATE COMPLETE AND CORDIART™ • 60 softgels	68.00	51.00	46.50			
00997	ENDOTHELIAL DEFENSE™ w/GLISODIN® • 60 veg. caps	54.00	40.50	36.00			
01937	EPA/DHA (Mega) • 120 softgels	20.00	15.00	13.50			
01737	ESOPHAGEAL GUARDIAN (Berry flavor) • 60 chewable tablets	36.00	27.00	24.00			
01042	EUROPEAN LEG SOLUTION DIOSMIN 95 600 mg, 30 veg. tabs	20.00	15.00	13.50			
01706	EXTRAORDINARY ENZYMES • 60 caps	26.00	19.50	18.00			
02008	CALIFORNIA ESTATE EXTRA VIRGIN OLIVE OIL • 500 ml (16.9 fl. oz)	33.00	24.75	22.50			
01514	EYE PRESSURE SUPPORT W/MIRTOGENOL® • 30 veg. caps	38.00	28.50	25.50			
F							
*01054	FACE MASTER® PLATINUM • Facial Toning System	199.00	199.00				
00965	FAST-ACTING JOINT FORMULA • 30 caps	39.00	29.25	27.00			
01717	FAST-C® W/DIHYDROQUERCETIN • 120 veg. tabs	26.00	19.50	18.00			
20053	FEM DOPHILUS® • 30 caps	25.95	19.46				
20055	FEM DOPHILUS® • 60 caps	39.95	29.96				
01064	FEMMENSENCE MACAPAUSE® • 120 veg. caps	34.99	26.24				
02007	FIBER-IMMUNE SUPPORT (Apple Cinnamon) • 235 grams	34.00	25.50	23.50			
00718	FIBRINOGEN RESIST™ • 30 veg. caps	49.00	36.75	33.00			
01749	FLAX SEED (Organic golden) • 14 oz	11.67	8.75				
01821	FLORASSIST® HEART HEALTH • 60 veg. caps	32.00	24.00	21.00			
02019	FLORASSIST® ORAL HYGIENE • 30 lozenges	18.00	13.50	12.75			
01825	FLORASSIST® BALANCE • 30 liquid veg. caps	32.00	24.00	21.00			
02000	FLORASSIST® MOOD • 60 caps	33.00	24.75	22.50			
01920	FLORASSIST® THROAT HEALTH • 30 lozenges	20.00	15.00	13.50			
01913	FOLATE HIGH POTENCY (Optimized) • 5,000 mcg, 30 veg. tablets	25.00	18.75	16.50			
01939	FOLATE (Optimized) • 1,000 mcg, 100 veg. tablets	19.00	14.25	12.75			
01842	FOLATE + VITAMIN B12 (BioActive) • 90 veg. caps	12.00	9.00	8.00			
01544	FORSKOLIN • 10 mg, 60 veg. caps	16.00	12.00	10.50			
01513	FUCOIDAN W/MARITECH® 926 (Optimized) • 60 veg. caps	36.00	27.00	24.75			
G							
00559	GAMMA E TOCOPHEROL/TOCOTRIENOLS • 60 softgels	42.00	31.50	27.75			
00759	GAMMA E TOCOPHEROL W/SESAME LIGNANS • 60 softgels	32.00	24.00	21.75			
01394	GARLIC (Optimized) • 200 veg. caps	24.95	18.71	15.75			
SUBTOTAL OF COLUMN 5							

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
*01122	GINGER FORCE® • 60 liquid caps	34.95	26.21				
01658	GINKGO BILOBA CERTIFIED EXTRACT™ 120 mg, 365 veg. caps	46.00	34.50	31.50			
00756	GLA WITH SESAME LIGNANS (Mega) • 60 softgels	19.50	14.63	13.50			
00345	(L-) GLUTAMINE CAPSULES • 500 mg, 100 veg. caps	14.95	11.21	10.13			
00141	(L-) GLUTAMINE POWDER • 100 grams	22.00	16.50	15.00			
00522	GLUCOSAMINE/CHONDROITIN CAPSULES • 100 caps	38.00	28.50	24.00			
01541	GLUTATHIONE, CYSTEINE & C • 100 veg. caps	20.00	15.00	13.50			
01669	GLYCINE • 1,000 mg, 100 veg. caps	12.00	9.00	8.10			
01411	GRAPE SEED EXTRACT W/RESVERATROL & PTEROSTILBENE 100 mg, 60 veg. caps	36.00	27.00	25.50			
01620	GREEN COFFEE EXTRACT COFFEEGENIC® 400 mg, 90 veg. caps	32.00	24.00	21.00			
00953	GREEN TEA EXTRACT (Mega) • lightly caffeinated, 100 veg. caps	30.00	22.50	18.00			
00954	GREEN TEA EXTRACT (Mega) • decaffeinated, 100 veg. caps	30.00	22.50	18.00			
H							
01074	5 HTP • 100 mg, 60 caps	27.95	20.96				
*02002	HAIR, SKIN & NAIL REJUVENATION FORM W/VERISON® 90 tabs	32.00	24.00	22.00			
01738	HCA (Garnicia) • 90 veg. caps	17.00	12.75	11.25			
29754	HCACTIVE™ GARCINIA CAMBOGIA EXTRACT • 90 caps	30.00	22.50				
01393	HEPATOPRO • 900 mg, 60 softgels	50.00	37.50	34.50			
01527	HUPERZINE A • 200 mcg, 60 veg. caps	40.00	30.00	27.00			
00661	HYDRODERM® • 1 oz	79.95	59.96	49.00			
I							
*01060	I26 HYPERIMMUNE EGG • 140 grams powder	54.99	46.75				
01704	IMMUNE MODULATOR W/TINOFEND® • 60 veg. caps	17.00	12.75	11.25			
00955	IMMUNE PROTECT W/PARACTIN® • 30 veg. caps	29.50	22.13	19.91			
02005	IMMUNE SENESCENCE PROTECTION FORMULA™ • 60 veg. tabs	40.00	30.00	27.00			
01049	INNERPOWER™ • 530 grams powder	42.00	31.50				
01674	INOSITOL CAPSULES • 1,000 mg, 360 veg. caps	62.00	46.50	43.50			
01292	INTEGRA-LEAN® AFRICAN MANGO IRVINGIA 150 mg, 60 veg. caps	28.00	21.00	18.00			
01677	IRON PROTEIN PLUS • 300 mg, 100 caps	28.00	21.00	19.50			
01492	IRVINGIA W/PHASE 3™ CALORIE CONTROL COMPLEX (Optimized African Mango) • 120 veg. caps	56.00	42.00	36.00			
J, K, L							
00056	JARRO-DOPHILUS EPS® • 60 veg. caps	22.95	17.21				
01834	K W/ADVANCED K2 COMPLEX (Super) • 90 softgels	30.00	22.50	20.25			
01600	KRILL HEALTHY JOINT FORMULA • 30 softgels	32.00	24.00	21.75			
01050	KRILL OIL • 60 softgels	33.95	25.46				
00316	KYOLIC® GARLIC FORMULA 102 • 200 veg. caps	27.45	20.59				
00214	KYOLIC® GARLIC FORMULA 105 • 200 caps	28.45	21.34				
00789	KYOLIC® RESERVE • 600 mg, 120 caps	28.95	21.71				
01681	LACTOFERRIN • 60 caps	52.00	39.00	36.00			
00020	LECITHIN • 16 oz granules	18.00	13.50	12.00			
02055	LIFE EXTENSION MIX™ • 315 tablets	80.00	60.00	52.00	43.75		
02057	LIFE EXTENSION MIX™ W/EXTRA NIACIN • 315 tablets	80.00	60.00	52.00	43.75		
02054	LIFE EXTENSION MIX™ • 490 caps	90.00	67.50	58.00	47.50		
SUBTOTAL OF COLUMN 6							

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
02056	LIFE EXTENSION MIX™ POWDER • 14.81 oz	80.00	60.00	52.00	43.75		
02065	LIFE EXTENSION MIX™ • 315 tablets w/o copper	80.00	60.00	52.00	43.75		
02064	LIFE EXTENSION MIX™ • 490 caps w/o copper	90.00	67.50	58.00	47.50		
02066	LIFE EXTENSION MIX™ POWDER • 14.81 oz w/o copper	80.00	60.00	52.00	43.75		
01608	LIVER EFFICIENCY FORMULA • 30 veg. caps	18.00	13.50	12.00			
01639	5-LOX INHIBITOR W/APRÈSFLEX® • 100 mg, 60 veg. caps	22.00	16.50	15.00			
01678	L-LYSINE • 620 mg, 100 veg. caps	9.00	6.75	6.00			
00455	LYCOPENE (Mega) • 15 mg, 90 softgels	35.00	26.25	22.50			
M							
01992	MACUGUARD® OCULAR SUPPORT • 60 softgels	25.00	18.75	17.50			
01993	MACUGUARD® OCULAR SUPPORT w/ASTAXANTHIN 60 softgels	44.00	33.00	30.00			
01459	MAGNESIUM CAPS • 500 mg, 100 veg. caps	12.00	9.00	7.50			
01682	MAGNESIUM (CITRATE) • 160 mg, 100 veg. caps	9.00	6.75	5.63			
01908	MEDITERRANEAN TRIM WITH SINETROL™-XPUR 60 veg. caps	18.00	13.50	12.00			
01668	MELATONIN • 300 mcg, 100 veg. caps	5.75	4.31	3.75			
01083	MELATONIN • 500 mcg, 200 veg. caps	18.00	13.50	12.00			
00329	MELATONIN • 1 mg, 60 caps	5.00	3.75	3.47			
00330	MELATONIN • 3 mg, 60 veg. caps	8.00	6.00	5.16			
00331	MELATONIN • 10 mg, 60 veg. caps	28.00	21.00	18.00			
00332	MELATONIN • 3 mg, 60 veg. lozenges	8.00	6.00	5.16			
01734	MELATONIN (Fast-Acting Liquid) • 2 fl. oz (Citrus-Vanilla)	12.00	9.00	8.25			
01787	MELATONIN TIMED RELEASE • 300 mcg, 100 veg. tabs	12.00	9.00	8.25			
01788	MELATONIN TIMED RELEASE • 750 mcg, 60 veg. tablets	8.00	6.00	5.25			
01786	MELATONIN TIMED RELEASE • 3 mg, 60 veg. tabs	12.00	9.00	8.25			
01536	METHYLCOBALAMIN • 1 mg, 60 veg. lozenges (vanilla)	9.95	7.46	6.00			
01537	METHYLCOBALAMIN • 5 mg, 60 veg. lozenges (vanilla)	32.00	24.00	18.75	17.25		
00709	MIGRA-EEZE™ (Butterbur) • 60 softgels	29.50	22.13	19.75			
01522	MILK THISTLE (European) • 60 veg. caps	34.00	25.50	22.50			
01922	MILK THISTLE (European) • 60 softgels	28.00	21.00	18.75			
01925	MILK THISTLE (European) • 120 softgels	44.00	33.00	30.00			
01940	MIRAFORTE w/STANDARDIZED LIGNANS (Super) • 120 veg caps	62.00	46.50	42.00			
01869	MITOCHONDRIAL BASICS W/BIOPQQ® • 30 caps	44.00	33.00	30.00			
01868	MITOCHONDRIAL ENERGY OPTIMIZER w/BIOPQQ® • 120 caps	72.00	54.00	48.00			
00065	MK-7 • 90 mcg, 60 softgels	28.00	21.00	18.75			
00451	MSM (Methylsulfonylmethane) • 1,000 mg, 100 caps	14.00	10.50	8.96			
N							
01534	N-ACETYL-L-CYSTEINE • 600 mg, 60 veg. caps	14.00	10.50	10.13			
01904	NAD+ CELL REGENERATOR™ • 100 mg, 30 veg. caps	34.00	25.50	19.50			
00066	NATTOKINASE • 60 softgels	25.50	19.13				
01807	NATURAL APPETITE SUPPRESS (Advanced) • 60 veg. caps	38.00	28.50	25.50			
00984	NATURAL BP MANAGEMENT • 60 tablets	44.00	33.00	30.00			
01892	NATURAL ESTROGEN • 60 veg. tabs	38.00	28.50	25.50			
01626	NATURAL SEX FOR WOMEN® 50+ (Advanced) • 90 veg. caps	59.00	44.25	34.00			
01444	NATURAL SLEEP® • 60 veg. caps	13.00	9.75	7.50			
01551	NATURAL SLEEP® w/ MELATONIN (Enhanced) • 30 caps	22.00	16.50	15.00			
SUBTOTAL OF COLUMN 7							

RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
01511	NATURAL SLEEP® W/O MELATONIN (Enhanced) • 30 caps	20.00	15.00	13.50			
01445	NATURAL SLEEP® MELATONIN • 5 mg, 60 veg. caps	18.00	13.50	12.00			
00987	NATURAL STRESS RELIEF • 30 veg. caps	28.00	21.00	18.00			
30741	NERVIA® • 90 softgels	53.95	40.46				
01603	NEURO-MAG® MAGNESIUM L-THREONATE • 90 veg. caps	40.00	30.00	27.00			
01602	NEURO-MAG® MAGNESIUM L-THREONATE w/CALCIUM & VITAMIN D3 • 25 grams • Lemon flavor	40.00	30.00	27.00			
01990	NITROVASC w/CORDIART™ • 30 veg. caps	18.00	13.50	12.00			
01903	NK CELL ACTIVATOR™ • 30 veg. tablets	45.00	33.75	31.50			
00373	NO-FLUSH NIACIN • 800 mg, 100 caps	19.00	14.25	12.75			
O							
01824	OLIVE LEAF VASCULAR SUPPORT w/CELERY SEED EXTRACT (Advanced) • 60 veg. caps	36.00	27.00	24.00			
01988	OMEGA-3 PLUS EPA/DHA w/SESAME LIGNANS, OLIVE EXTRACT, KRILL & ASTAXANTHIN (SUPER) • 120 softgels	45.00	33.75	31.50	24.75		
01983	OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 60 softgels	18.00	13.50	12.00	9.38		
01982	OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 120 softgels	32.00	24.00	21.00	17.05		
01984	OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 120 enteric coated softgels	34.00	25.50	23.25	18.00		
01985	OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 60 enteric coated softgels	20.00	15.00	13.50	10.50		
01986	OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 240 small softgels	32.00	24.00	21.00	17.25		
01991	ONCE-DAILY HEALTH BOOSTER • 60 softgels	54.00	40.50	38.00			
02001	ONE-PER-DAY • 60 tablets	22.00	16.50	15.00			
01328	ONLY TRACE MINERALS • 90 veg. caps	15.00	11.25	9.38			
P							
01789	PALMETTOGUARD® SAW PALMETTO W/BETA-SITOSTEROL 30 softgels	15.00	11.25	10.50	9.00		
01790	PALMETTOGUARD® SUPER SAW PALMETTO/NETTLE ROOT W/BETA-SITOSTEROL • 60 softgels	28.00	21.00	19.50	18.00		
01323	PEAK ATP® WITH GLYCO-CARN® • 60 veg. caps	54.00	40.50	37.50			
*00342	PECTA SOL-C® MODIFIED CITRUS PECTIN • 454 grams powder	109.95	93.46				
*01080	PECTA SOL-C® MODIFIED CITRUS PECTIN • 270 veg. caps	79.95	67.96				
01811	PEONY IMMUNE • 60 veg. caps	36.00	27.00	24.00			
00673	PGX® PLUS MULBERRY (WellBetX®) • 180 veg. caps	34.95	26.21				
01676	PHOSPHATIDYLSERINE CAPS • 100 mg, 100 veg. caps	54.00	40.50	36.00			
01953	POMEGRANATE COMPLETE • 30 softgels	24.00	18.00	15.75			
00956	POMEGRANATE FRUIT EXTRACT • 30 veg. caps	19.50	14.63	13.16			
01500	PQQ CAPS W/BIOPQQ® • 10 mg, 30 veg. caps	24.00	18.00	13.50	12.00		
01647	PQQ CAPS W/BIOPQQ® • 20 mg, 30 veg. caps	40.00	30.00	24.00	21.00		
00302	PREGNENOLONE • 50 mg, 100 caps	26.00	19.50	16.50			
00700	PREGNENOLONE • 100 mg, 100 caps	30.00	22.50	20.25			
**01373	PRELOX® NATURAL SEX FOR MEN® • 60 tablets	52.00	39.00	36.00			
01576	PREVAGEN® • 30 caps	60.00	45.00				
*01577	PREVAGEN® ES • 30 caps	70.00	60.00				
00525	PROBOOST™ THYMIC PROTEIN A • 30 packets	66.60	49.95				
01441	PROGESTA-CARE® • 4 oz cream	36.39	27.29	25.72			
SUBTOTAL OF COLUMN 8							

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
01928	PROSTATE FORMULA (Ultra NAT) • 60 softgels	38.00	28.50	26.25	24.00		
01909	PROSTAPOLLEN™ (Triple strength) • 30 softgels	28.00	21.00	18.75			
01742	PROTEIN-ISOLATE (Whey) Vanilla • 403 grams	30.00	22.50	20.25			
01743	PROTEIN-ISOLATE (Whey) Chocolate • 437 grams	30.00	22.50	20.25			
01770	PROTEIN CONCENTRATE (New Zealand Whey) Vanilla 500 grams	30.00	22.50	19.95			
01771	PROTEIN CONCENTRATE (New Zealand Whey) Chocolate 640 grams	30.00	22.50	19.95			
01812	PROVINAL® PURIFIED OMEGA-7 • 30 softgels	27.00	20.25	18.00			
01508	PTEROPURE® Pterostilbene • 50 mg, 60 veg. caps	32.00	24.00	22.50			
01209	PUMPKIN SEED EXTRACT (Water-soluble) • 60 veg. caps	20.00	15.00	13.50			
01637	PYCNOGENOL® FRENCH MARITIME PINE BARK EXTRACT 100 mg, 60 veg. caps	64.00	48.00	45.00			
01217	PYRIDOXAL 5'-PHOSPHATE • 100 mg, 60 veg. caps	22.00	16.50	14.85			
Q, R							
01309	QUERCETIN (Optimized) • 250 mg, 60 veg. caps	22.00	16.50	15.00			
01030	RED YEAST RICE (Bluebonnet) • 600 mg, 60 veg. caps	17.60	13.20				
00605	REGIMINT • 60 enteric-coated caps	19.95	14.96	14.00			
01708	REISHI EXTRACT MUSHROOM COMPLEX • 60 veg. caps	30.00	22.50	20.25			
01448	REJUVENEX® BODY LOTION • 6 oz	24.00	18.00	14.85	12.75		
01621	REJUVENEX® FACTOR FIRING SERUM • 1.7 oz	65.00	48.75	37.50			
01220	REJUVENEX® (Ultra) • 2 oz	52.00	39.00	33.00	29.25		
00676	REJUVENIGHT® (Ultra) • 2 oz	39.95	29.96	27.00			
01410	RESVERATROL W/PTEROSTILBENE • 100 mg, 60 veg. caps	36.00	27.00	24.00			
02031	RESVERATROL W/NICOTINAMIDE RIBOSIDE (Optimized) • 30 veg. caps	42.00	31.50	27.00			
02030	RESVERATROL (Optimized) • 60 veg. caps	46.00	34.50	31.00			
00889	RHODIOLA EXTRACT • 250 mg, 60 veg. caps	14.00	10.50	9.00			
01900	RIBOGEN™ FRENCH OAK WOOD EXTRACT 200 mg, 30 veg. caps	36.00	27.00	24.75			
00972	(D) RIBOSE POWDER • 150 grams	27.50	20.63	18.56			
01473	(D) RIBOSE TABLETS • 100 veg. tabs	32.00	24.00	21.00			
01609	RICH REWARDS® BREAKFAST GROUND COFFEE • 12 oz. bag	13.00	9.75				
01730	RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Mocha • 12 oz. bag	15.00	11.25	10.50			
01729	RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Vanilla • 12 oz. bag	15.00	11.25	10.50			
01612	RICH REWARDS® BREAKFAST BLEND WHOLE BEAN COFFEE 12 oz. bag	13.00	9.75				
01610	RICH REWARDS® DECAFFEINATED ROAST GROUND COFFEE 12 oz. bag	14.00	10.50				
01208	R-LIPOIC ACID (Super) • 240 mg, 60 veg. caps	49.00	36.75	33.75			
00070	RNA CAPSULES • 500 mg, 100 caps	17.95	13.46	12.12			
S							
01432	SAFFRON W/SATIAREAL® (Optimized) • 60 veg. caps	36.00	27.00	24.00			
01935	SAMe (S-ADENOSYL-METHIONINE) 200 mg, 30 enteric coated tablets	25.00	18.75	16.50			
01933	SAMe (S-ADENOSYL-METHIONINE) 400 mg, 30 enteric coated tablets	36.00	27.00	24.00			
01934	SAMe (S-ADENOSYL-METHIONINE) 400 mg, 60 enteric coated tablets	66.00	49.50	45.00			

SUBTOTAL OF COLUMN 9

ITEM No.	PRODUCT	YOUR PRICE				QTY	Total
		Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each		
01740	SEA-IODINE™ • 1,000 mcg, 60 veg. caps	8.00	6.00	5.40			
00046	SELENIUM • 2 fl. oz dropper	11.95	8.96				
01879	SE-METHYL L-SELENOCYSTEINE • 200 mcg, 90 veg. caps	11.00	8.25	7.50			
00318	SERRAFLAZYME • 100 tablets	18.00	13.50	12.00			
01938	SHADE FACTOR • 120 veg. caps	44.00	33.00	30.00			
01884	SILYMARIN • 100 mg, 90 veg. caps	14.00	10.50	9.50			
01249	SINUS CLEANSER • 4 oz. bottle	25.00	18.75				
01596	SKIN RESTORING PHYTCERAMIDES w/LIPOWHEAT® 30 liquid veg. caps	25.00	18.75	17.25			
00961	SODZYME® w/GLISODIN® & WOLFBERRY • 90 veg. caps	28.00	21.00	18.00			
00657	SOLARSHIELD® SUNGLASSES • Smoke color	12.99	9.74	8.63			
01097	SOY EXTRACT (Ultra) • 150 veg. caps	87.00	65.25	58.50			
00432	STEVIA™ (Better) • 100 packets, 1 gram each	9.95	7.46				
00438	STEVIA™ ORGANIC LIQUID SWEETENER (Better) • 2 oz	11.00	8.25				
01476	STRONTIUM • 750 mg, 90 veg. caps	20.00	15.00	13.50			
01649	SUPER ABSORBABLE SOY ISOFLAVONES • 60 veg. caps	28.00	21.00	18.75			
01778	SUPER SELENIUM COMPLEX • 200 mcg, 100 veg. caps	14.00	10.50	9.00	8.25		
T							
02023	TART CHERRY EXTRACT W/STANDARDIZED CHERRYPURE® 60 veg. caps	20.00	15.00	14.00			
01827	TAURINE • 1,000 mg, 90 veg. caps	13.00	9.75	9.00			
01918	TEAR SUPPORT w/MAQUIBRIGHT® • 60 mg, 30 veg. caps	18.00	13.50	12.00			
00133	L-TAURINE POWDER • 300 grams	20.00	15.00	12.66			
*13685	TEN MUSHROOM FORMULA® • 120 veg. caps	39.95	33.96				
01304	THEAFLAVIN STANDARDIZED EXTRACT • 30 veg. caps	18.00	13.50	12.00			
01683	(L) THEANINE • 100 mg, 60 veg. caps	24.00	18.00	15.38			
**01038	THERALAC® PROBIOTICS • 30 caps	47.95	35.96				
00668	THYROID FORMULA (Metabolic Advantage™) • 100 caps	21.95	16.46				
00349	TMG POWDER • 50 grams	14.00	10.50	8.25			
01859	TMG • 500 mg, 60 liquid veg. caps	13.00	9.75	9.00			
01400	TOCOTRIENOLS (Super-absorbable) • 60 softgels	30.00	22.50	21.00			
01278	TOOTHPASTE • 4 oz (Mint) tube	9.50	7.13	6.50			
01917	TRANQUIL TRACT™ • 60 veg. caps	52.00	39.00	34.50			
01468	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT 60 veg. caps	24.00	18.00	16.50			
01469	TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT w/RESVERATROL • 60 veg. caps	32.00	24.00	22.20			
02003	TRIPLE ACTION THYROID • 60 veg. caps	36.00	27.00	24.00			
01803	TRI SUGAR SHIELD® • 60 veg. caps	36.00	27.00	24.00			
01386	TRUFIBER™ • 180 grams	32.95	24.71				
01389	TRUFLOA® PROBIOTICS • 32 veg. caps	42.95	32.21				
01722	L-TRYPTOPHAN • 500 mg, 90 veg. caps	33.00	24.75	22.50			
01721	TRYPTOPHAN PLUS (Optimized) • 90 veg. caps	32.00	24.00	21.75			
02016	TWO-PER-DAY • 60 tablets	10.50	7.88	7.13			
02015	TWO-PER-DAY • 120 tablets	20.00	15.00	13.50			
02014	TWO-PER-DAY • 120 caps	22.00	16.50	15.00			
00326	L-TYROSINE • 500 mg, 100 tablets	12.98	9.74				

SUBTOTAL OF COLUMN 10

ITEM No.	PRODUCT	Retail Each \$	YOUR PRICE			QTY	Total
			1 Unit Each	4 Unit Each	10 Unit Each		
U, V							
01921	URIC ACID CONTROL • 60 veg. caps	24.00	18.00	16.50			
00213	VANADYL SULFATE • 7.5 mg, 100 veg. tablets	15.00	11.25	9.38			
00408	VENOTONE • 60 caps	18.95	14.21	12.00			
01327	VINPOCETINE • 10 mg, 100 veg. tablets	18.00	13.50	10.50			
00372	VITAMIN B3 NIACIN • 500 mg, 100 caps	7.65	5.74	4.99			
00098	VITAMIN B5 • 500 mg, 100 caps (Pantothenic Acid)	10.50	7.88	7.04			
01535	VITAMIN B6 • 250 mg, 100 veg. caps	12.50	9.38	8.25			
00361	VITAMIN B12 • 500 mcg, 100 lozenges	8.75	6.56	5.44			
01634	VITAMIN C w/DIHYDROQUERCETIN 1,000 mg, 60 veg. tablets	10.00	7.50	6.75			
00927	VITAMIN C w/DIHYDROQUERCETIN 1,000 mg, 250 veg. tablets	25.50	19.13	17.44			
00084	VITAMIN C POWDER (BUFFERED) • 454 grams	23.95	17.96	16.50			
01736	VITAMIN C-MAGNESIUM CRYSTALS (EFFERVESCENT) 180 grams	20.00	15.00	13.50			
01732	VITAMIN D3 • 2,000 IU, 1 fl. oz, Mint flavor	28.00	21.00	18.75			
01753	VITAMIN D3 • 1,000 IU, 90 softgels	7.00	5.25	4.50			
01751	VITAMIN D3 • 1,000 IU, 250 softgels	12.50	9.38	8.44			
01713	VITAMIN D3 • 5,000 IU, 60 softgels	10.00	7.50	6.50			
01718	VITAMIN D3 • 7,000 IU, 60 softgels	14.00	10.50	9.45			
01758	VITAMIN D3 W/SEA-IODINE™ • 5,000 IU, 60 caps	14.00	10.50	9.38			
00864	VITAMIN D3 LIQUID EMULSION • 2,000 IU, 1 fl. oz	28.00	21.00	18.75			
01840	VITAMINS D AND K W/SEA-IODINE™ • 60 caps	24.00	18.00	16.50			
01863	VITAMIN E (Natural) • 400 IU, 90 softgels	28.00	21.00	19.50	18.00		
01936	VITAMIN K2 (Low dose) • 45 mcg, 90 softgels	18.00	13.50	12.00			
W							
01902	WAIST-LINE CONTROL™ • 120 veg. caps	42.00	31.50	28.50			
X, Y							
01919	X-R SHIELD • 90 veg. caps	15.00	11.25	9.75			
00409	XYLIWHITE™ MOUTHWASH • 16 oz	10.00	7.50				
Z							
01813	ZINC HIGH POTENCY • 50 mg, 90 veg. caps	7.95	5.96	5.25			
01561	ZINC LOZENGES • 60 veg. lozenges	9.00	6.75	6.00			
01961	ZINC LOZENGES (Enhanced) • 30 veg. lozenges	12.00	9.00	6.00			
*01051	ZYFLAMEND® WHOLE BODY • 120 liquid veg. caps	72.95	54.71				
BOOKS							
34002	THE 30-DAY HEART TUNE-UP by Steven Masley, MD • 2016	17.59	13.19				
33999	THE MENOPAUSE CURE by Jill D. Davey & Sergey Dzugan, MD • 2016	17.32	12.99				
33998	THE RIGHT TO TRY by Darcy Olsen • 2016	26.99	20.24				
33840	THE CRWAY® TO GREAT GLUCOSE CONTROL CD by Paul McGlothlin and Meredith Averill • 2016	189.00	189.00				
33890	FORTIFY YOUR LIFE by Tieraona Low Dog, MD • 2016	28.89	21.67				
33885	THE BLUE ZONES SOLUTION by Dan Buettner • 2015	26.00	19.50				
33880	OUTSTANDING HEALTH: THE 6 ESSENTIAL KEYS TO MAXIMIZE YOUR ENERGY AND WELL BEING by Michael Galitzer, MD & Larry Trivieri Jr. • 2015	24.95	18.71				
SUBTOTAL OF COLUMN 11							

RECEIVE 25% OFF THE RETAIL PRICE OF ALL PRODUCTS

ITEM No.	PRODUCT	Retail Each \$	YOUR PRICE			QTY	Total
			1 Unit Each	4 Unit Each	10 Unit Each		
33878	TESTOSTERONE REPLACEMENT THERAPY by Dr. John Crisler • 2015	19.99	14.99				
33877	THE TRUTH ABOUT MEN AND SEX by Abraham Morgentaler, MD, FACS • 2015	16.99	12.74				
33876	TOX-SICK • by Suzanne Somers • 2015	26.00	19.50				
33875	DOCTORED: THE DISILLUSIONMENT OF AN AMERICAN PHYSICIAN • by Sandeep Jauhar • 2015	26.00	19.50				
33874	MISSING MICROBES • by Martin J. Blaser, MD • 2014	28.00	21.00				
33873	EATING ON THE WILD SIDE • by Jo Robinson • 2014	16.00	12.00				
33872	GET SERIOUS • by Brett Osborn, MD • 2014	24.95	18.71				
33868	TOXIN TOXOUT: GETTING HARMFUL CHEMICALS OUT OF OUR BODIES AND OUR WORLD • by Bruce Lourie and Rick Smith • 2014	25.99	19.49				
33867	THE COMPLETE MEDITERRANEAN DIET by Michael Ozner, MD • 2014	19.95	14.96				
33869	UNLEASH THE POWER OF THE FEMALE BRAIN by Daniel Amen, MD • 2014	16.00	12.00				
33870	MAGNIFICENT MAGNESIUM by Dennis Goodman, MD • 2014	14.95	11.21				
DPT05	DISEASE PREVENTION AND TREATMENT, EXPANDED FIFTH EDITION (Hardcover) • 2014	69.95	39.95	36.00			
33865	THE RESTORATION OF THE HUMAN BODY [IN 7 PARTS] by Sergey A. Dzugan, MD, PhD • 2014	29.95	22.46				
33862	I'M TOO YOUNG FOR THIS • by Suzanne Somers • 2013	26.00	19.50				
33835	PHARMOCRACY • by William Faloon • 2011	24.00	9.60	8.00			
33958	THE VITAMIN D SOLUTION by Michael F. Holick, PhD, MD (Paperback) • 2013	16.00	12.00				
33838	YOUR GUIDE TO HEALTHY SKIN THE NATURAL WAY by Gary Goldfaden, MD • 2012	26.00	15.00				
33815	KNOCKOUT • by Suzanne Somers • 2009	25.99	17.00				
33809	TESTOSTERONE FOR LIFE by Abraham Morgentaler, MD • 2008	16.95	11.87				
33696	LIFE EXTENSION REVOLUTION by Philip Lee Miller, MD (Paperback)	16.00	12.00				
33805	MIAMI MEDITERRANEAN DIET WITH 300 RECIPES by Michael D. Ozner, MD, FACC, FAHA (Hardcover) • 2008	24.95	16.25				
33906	THE MIGRAINE CURE • by Sergey Dzugan, MD, PhD • 2006	24.00	15.60				
33803	WHAT YOUR DOCTOR MAY NOT TELL YOU ABOUT DIABETES by Steven V. Joyal, MD • 2008	14.99	10.49				
SUBTOTAL OF COLUMN 12							

- * These products are not 25% off retail price.
- ** Due to license restrictions, this product is not for sale to customers outside of the USA.
- *** Due to license restrictions, this product is not for sale to Canada.
- † Due to license restrictions, this product is not for sale to customers outside of the USA and Canada.



Not sure exactly which supplements you need?
Talk to a Wellness Specialist toll-free at 1-800-226-2370



ORDER SUBTOTALS

SUBTOTAL COLUMN 1	
SUBTOTAL COLUMN 2	
SUBTOTAL COLUMN 3	
SUBTOTAL COLUMN 4	
SUBTOTAL COLUMN 5	
SUBTOTAL COLUMN 6	
SUBTOTAL COLUMN 7	
SUBTOTAL COLUMN 8	
SUBTOTAL COLUMN 9	
SUBTOTAL COLUMN 10	
SUBTOTAL COLUMN 11	
SUBTOTAL COLUMN 12	
ORDER TOTALS	
SUBTOTAL OF COLUMNS 1 - 12	
POSTAGE & HANDLING (Any size order, in the U.S., includes Alaska & Hawaii)	\$5.50
C.O.D.s (ADD \$7 FOR C.O.D. ORDERS)	
SHIPPING <small>UPS OVERNIGHT add \$16, UPS 2nd DAY AIR add \$7. For Puerto Rico, US Virgin Islands, add \$7. CANADA UPS EXPRESS Flat rate \$17.50, UK Flat rate \$25 USD. ALL OTHER INTERNATIONAL AIR WILL BE ADDED.</small>	
GRAND TOTAL (MUST BE IN U.S. DOLLARS)	

PLEASE MAIL TO: Life Extension
P.O. Box 407198 • Ft. Lauderdale, Florida 33340-7198
Or Call Toll Free 1-800-544-4440 • Fax: 866-728-1050

PREMIER Order Over \$50?

Get a year of unlimited
FREE shipping and
double rewards for only \$0.

There's no catch: just apply your instant \$50 bonus at checkout. Your orders ship free for 12 months, you earn double (4%) rewards back all year long, and the cost of enrollment is offset by your instant bonus.* This makes Premier the *only* rewards program that more than pays for itself!

Premier. Simply. More.

Just \$49.95 | \$59.95 for international customers

Call **1-888-224-8239** to enroll

Visit www.LifeExtension.com/Premier
for details • Mention code **YRX618D**

LifeExtension®



No auto-renewal, no commitments.
*Certain restrictions apply. For complete terms and conditions please visit www.LifeExtension.com/Premier

BILL TO ADDRESS

NAME _____ E-MAIL _____

ADDRESS _____

CITY/STATE/ZIP-POSTAL CODE _____ COUNTRY _____

PHONE _____ FAX _____

VISA/MASTERCARD/AMEX/DISCOVER # _____

EXP. DATE _____

SIGNATURE _____

SHIP TO ADDRESS

NAME _____ E-MAIL _____

ADDRESS _____

CITY/STATE/ZIP-POSTAL CODE _____

COUNTRY _____

PHONE _____ FAX _____

SIGNATURE _____

**Prices subject to change without notice.
Please notify Life Extension of any address change.**

The Most Advanced Probiotic On The Market



WITH UNIQUE
DUAL
ENCAPSULATION
TECHNOLOGY

Scientists are increasingly discovering that probiotics impact the health of the entire body.¹⁻⁶ Unfortunately, most commercial probiotics are destroyed by the stomach's natural digestive acids before they reach their destination.⁷

FLORASSIST® Balance with "dual encapsulation" technology delivers maximum probiotic protection to your small intestines.

FLORASSIST® Balance:

- Contains probiotic strains that are **stomach acid resistant**
- Has **dual encapsulation technology**, which keeps the capsule intact longer and ensures that the probiotic reaches the small intestine
- Provides **15 billion CFU**—Colony Forming Units—per capsule[†]
- Contains **6** varieties of beneficial bacteria

FLORASSIST® Balance contains the following bacterial strains:

1. *Lactobacillus acidophilus* LA-14
2. *Lactobacillus rhamnosus* LR-32
3. *Lactobacillus paracasei* LPC-37
4. *Bifidobacterium longum* BL-05
5. *Bifidobacterium lactis* BL-04
6. *Bifidobacterium bifidum/lactis* BB-02

These potent strains of probiotic bacteria adhere to the soft lining of the intestinal tract to help maintain a healthy surface and aid in support for the digestive system.

[†] Colony forming units at time of manufacture.

FLORASSIST® Balance

Item #01825 • 30 liquid vegetarian capsules

	Retail Price	Your Price
1 bottle	\$32	\$24
4 bottles		\$21 each

Non-GMO



To order **FLORASSIST® Balance**,
call **1-800-544-4440**
or visit **www.LifeExtension.com**

References

1. *Eur J Clin Nutr.* 2013 Feb;67(2):161-7.
2. *Curr Top Microbiol Immunol.* 2013;358:273-89.
3. *Br J Nutr.* 2013 May 28;109(10):1866-72.
4. *Nutr Hosp.* 2011 Jan-Feb;26(1):228-35.
5. *Eur J Cancer Prev.* 2013 Jan;22(1):46-51.
6. *Pediatr Int.* 2012 Oct;54(5):682-7.
7. *Microbiology.* 2007 Oct;153(Pt 10):3563-71.



WHAT'S INSIDE

Visit us at www.LifeExtension.com

LifeExtension®

Magazine



7 GOVERNMENT-INFLICTED COLON CANCER

Life Extension Magazine® publishes material the government prefers you not read. Government groups are now recommending *against* colonoscopies. This will cut their healthcare costs in the short-term, but the tragic result will be colorectal cancers that go undetected and metastasize.



24 NOVEL APPROACH TO ALLEVIATE DEPRESSION

A misunderstood cause of depression is high **homocysteine** blood levels. Researchers found that **5-MTHF**, a form of **folate**, lowered homocysteine levels and significantly improved depression.



46 YOUNGER-LOOKING SKIN

A new self-administered exfoliating formula harnesses the unique properties of microdermabrasion utilizing ultra-fine **amber crystals** and **jojoba beads**.



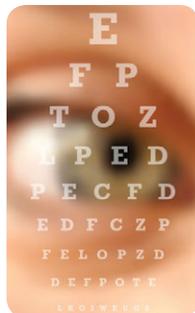
34 PROTECT AGAINST COLORECTAL CANCER

Colorectal cancer is the most preventable cause of cancer death. Healthy dietary practices and specific nutrients can help reduce risk of colon cancer development.



54 AGE-REVERSAL CONFERENCE

Life Extension Foundation® is cosponsoring a conference in San Diego on August 4-7, 2016. The focus will be to accelerate human **age-reversal** research.



87 NEWLY DISCOVERED LUTEIN BENEFITS

Lutein, well-known for protecting vision, can also help prevent *cardio-metabolic* disorders such as heart disease, stroke, and diabetes.