Turn "On" Your DNA Repair

LIFE EXTENSION.com

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

September 2017

Age Reversal Impact of NAD+

Improve Sleep Via Gene Repair

Probiotics Improve Immune Health

Relieve Nighttime Urinary Distress

Novel Method to Prevent Glaucoma



BEN STILLER

His Triumph Over Prostate Cancer





Compare TWO-PER-DAY to the Leading Brand CENTRUM®



The **Two-Per-Day** multinutrient formula is superior to commercial multivitamins because it provides vastly *higher* potencies of **vitamins**, **minerals** and **plant extracts**.

TWO-PER-DAY

provides:

50 times the vitamin B1 25 times the vitamin B6

12 times the vitamin B12

10 times the biotin

10 times the selenium

8 times the vitamin C

2 times the vitamin D

2 times the vitamin E

2 times the vitamin B3

3 times the zinc

Two-Per-Day Capsules

Item #02114 • 120 capsules • Non-GMO

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

Two-Per-Day Tablets

Item #02115 • 120 tablets • Non-GMO

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

Each bottle provides a two-month supply.





For full product description and to order call 1-800-544-4440 or visit Life Extension.com

* Rated based on results of the 2017 ConsumerLab.com Survey of Supplements Users. More information at www.consumerlab.com/survey2017



Volume Twenty Three / Number Nine • September 2017

REPORTS

7 NEW RESVERATROL (AND NAD+) DOSING PROTOCOL

Resveratrol promotes **longevity** by activating proteins called **sirtuins**. For **sirtuins** to **function**, they *require* the coenzyme **NAD+**. Maturing people can derive considerable benefit by taking more NAD+ precursors with modest dose resveratrol. NAD+ is vital for youthful cell function including **DNA repair**.

40 IMPEDE UPPER RESPIRATORY INFECTIONS

Scientists have identified a targeted **probiotic cocktail** that can boost the body's immune defenses, in particular mucosal secretory **IgA**, thereby reducing colds, flu, and upper respiratory complications.

52 RELIEVE URINARY-TRACT SYMPTOMS

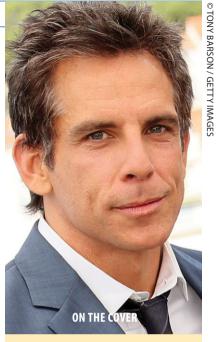
Urinary discomforts related to prostate enlargement are epidemic in aging men. Common symptoms include nighttime urinary frequency, urgency, and weak stream. Several **natural** extracts help alleviate urinary discomforts in men.

64 REVERSING A ROOT CAUSE OF GLAUCOMA

Glaucoma is a common cause of blindness. A human study demonstrates eye pressure reduced by 24% using two plant extracts. When combined with standard glaucoma therapy, intraocular **pressure** was lowered up to 40% with these natural extracts.

84 2017 CARDIOVASCULAR DISEASE PREVENTION SYMPOSIUM

Organized by Dr. Michael Ozner, the Cardiovascular Disease Prevention Symposium is an annual gathering of visionary cardiologists focused on conquering heart disease. Topics included PCSK9 to reduce cholesterol, benefits of fish oil and biomarkers of coronary artery disease.



30 BEN STILLER ADVOCATES PROSTATE CANCER SCREENING

Prostate cancer is curable when detected before it metastasizes. Yet millions of men are skipping annual PSA blood tests. Actor **Ben Stiller** is a beneficiary of early diagnosis and has become a passionate advocate for **PSA screening**.





21 IN THE NEWS

Meat increases mortality among cancer survivors; strict diabetes management extends life; NAD+ precursors slow aging; aspirin inhibits cancer; vitamin D relieves back pain; and updated disease protocols.

93 SUPERFOODS: CELERY

Celery has been cultivated for thousands of years and is a rich source of fiber, vitamin K, potent anti-inflammatory phytonutrients, and important minerals, including iron, zinc, copper, magnesium, calcium, selenium, and especially potassium.

75 RESEARCH UPDATE:

POMEGRANATE IMPROVES MITOCHONDRIA

Swiss researchers have identified a molecule produced in the body from pomegranates known as urolithin A that reduces a pathologic mechanism of aging called mitochondrial dysfunction.

95 WELLNESS PROFILE: **HEART TRANSPLANT SUCCESS STORY**

After Humberto Fasano's staggering 2001 diagnosis of severe cardiomyopathy and congestive heart failure (CHF), doctors told him that 50% of CHF patients usually die within five years of diagnosis. Over 15 years later, Fasano is thriving after he managed to outlast a chronic, seemingly hopeless disease with the help of firstrate cardiac care and Life Extension°.





Volume Twenty Three / Number Nine

Connect with Life Extension on the Web!



find us on facebook

Facebook.com/LifeExtension

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



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

Publisher • LE Publications, Inc.

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 Writer • Garry Messick
Creative Director • Robert Vergara
Art Director • Alexandra Maldonado

Chief Medical Officer

Steven Joyal, MD

Scientific Advisory Board

Örn Adalsteinsson, PhD • John Boik, PhD • Aubrey de Grey, PhD Frank Eichhorn, 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

Director of Scientific Affairs

Anita Boddie, PhD, FACN, RDN

Contributors

Michael Downey • Leslie Hunter • Garry Messick Alma Ross • Jon VanZile

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



Ratings based on results of the 2017 ConsumerLab.com Survey of Supplement Users. More information at www.consumerlab.com/survey2017.



LIFE EXTENSION (ISSN 1524-198X) Vol. 23, No. 9 ©2017 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 advertis

Support Youthful Brain Function

LifeExtension MEMORY PROTECT



Defends Against Memory Loss*

12 Colostrinin-Lithium (C-Li) Capsules | 24 Lithium (Li) Capsules 36 DAY SUPPLY | Dietary Supplement

Human studies demonstrate robust neurological benefits in response to low-dose **lithium** and colostrum-derived **proline-rich polypeptides**.

Memory Protect has been formulated with these <u>two</u> nutrients to support healthy structure of brain cells, normal memory, and recall function.

Each box is designed to enable you to take these nutrients on the identical dosage schedule used in successful clinical trials.

Memory Protect

Item #02101 • 36 capsules

	Retail Price	Your Price
1 box	\$24	\$18
4 boxes		\$16 each

For full product description and to order **Memory Protect**, call **1-800-544-4440** or visit **www.LifeExtension.com**

Non-GMO. Contains milk.

CAUTION: Consult your healthcare provider before use if you are taking medication. Do not use if pregnant, lactating, or trying to become pregnant. Rare, mild and temporary anxiety and/or sleep disturbance may be experienced.





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 boardcertified 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.

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 hemapoetic 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 Montenapoleone 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

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 boardcertified 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.

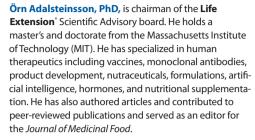
Michael D. Seidman, MD, FACS, is the director of skull base surgery and wellness for the Adventist Health System in Celebration, FL.

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.

Scientific Advisory Board







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.



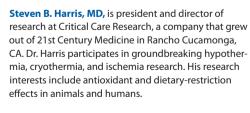
Aubrey de Grey, PhD, is a biomedical gerontologist and Editor-in-Chief of Rejuvenation Research, the world's highest-impact peer-reviewed journal focused on intervention in aging. He received his BA and PhD from the University of Cambridge in 1985 and 2000 respectively. Dr. de Grey is a Fellow of both 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 Eichhorn, 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.





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



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

NICOTINAMIDE RIBOSIDE

(NAD+ Precursor)



NAD+

(Nicotinamide Adenine Dinucleotide)



YOUTHFUL CELL FUNCTION

Normal **aging** results in marked <u>decline</u> of cellular **NAD+**.

Oral ingestion of nutrients like **nicotinamide riboside** boosts **NAD+** in cells throughout the body.



and NAD+ Suggestions

BY WILLIAM FALOON

NAD+ is a coenzyme found in all living cells. It is essential for **energy** production and **DNA repair.**¹⁻³

In **2014**, we introduced a **NAD+** precursor called **nicotinamide riboside**.⁴ When you take **nicotinamide riboside**, it converts to **NAD+** in your cells, where it facilitates **regenerative** processes.⁵

The most important **NAD+** benefit is promoting rapid **DNA repair** and fueling beneficial **longevity proteins.**^{2,3,6,7}

Resveratrol favorably enhances the <u>activation</u> of cellular **sirtuin proteins**.⁸ **NAD+** is required for our **sirtuins** to function.^{7,9,10}

As we age, **NAD+** levels plummet, which impedes the ability of **resveratrol** to deliver its benefits.¹¹⁻¹⁴

In **2003**, we introduced **resveratrol** based on its ability to <u>activate</u> **sirtuin longevity** factors.^{15,16}

New Resveratrol

Genetic research we conducted back then indicated that biologically meaningful **resveratrol** dosing for humans might be as low as **20 mg** a day. ^{16,17} Subsequent studies suggested *higher* resveratrol intake. ^{18,19}

Based on our interpretation of emerging evidence, we have reformulated our premium supplements to provide <u>more NAD+</u> precursor (**nicotinamide riboside**) with <u>lower</u> **resveratrol**.

You're going to learn about potential **age-reversal** benefits of boosting cellular **NAD+** later this year. This article will provide a summary of what's been uncovered in recent published studies.



NAD+ is required for healthy cellular functions including **DNA** repair.1-3

The amount of damage inflicted to cellular **DNA** is grossly underestimated. Be it background radiation or normal metabolic processes, our DNA is constantly "broken" and then "repaired" using specialized *coenzymes* like **NAD+**.

Failure to **repair** damaged DNA can result in cell death or transformation into malignant or senescent states.

NAD+ levels markedly decline with age.11-14 NAD+ deficit manifests clinically in the form of degenerative disorders of the brain, 20-23 heart, 24-26 and other tissues. 27,28

In animal studies, regenerative effects have been observed in the **brain** when NAD+ is restored.^{29,30}

Sleep quality deteriorates with normal aging in many people. Restoring youthful **NAD+** levels in the brain may support a healthy circadian rhythm.31

Loss of NAD+ activity is linked to type II diabetes. In mice, administration of an NAD+ precursor restores insulin sensitivity and protects against the diabetic impact of a high-fat diet. 6,32,33

Resveratrol and NAD+

Resveratrol has become a popular dietary supplement because of its ability to activate sirtuin proteins in our cells.34

When **sirtuins** are activated. the effect is delayed aging, which has been demonstrated in a wide spectrum of experimental models. including mammals.35-44

Sirtuins that are activated by **resveratrol** require **NAD+** as their energy substrate. Loss of NAD+ impedes beneficial sirtuin function.7,10,45

Younger people have high **NAD+** levels that enable them to benefit from the **sirtuin-boosting** effects of resveratrol.

To improve the <u>functionality</u> of **sirtuin** proteins, it makes sense for maturing individuals to boost their NAD+ levels.

The good news is that a precursor to NAD+ can be found in nicotinamide riboside supplements. New dosage recommendations can enable older people to restore cellular NAD+ to more youthful profiles.

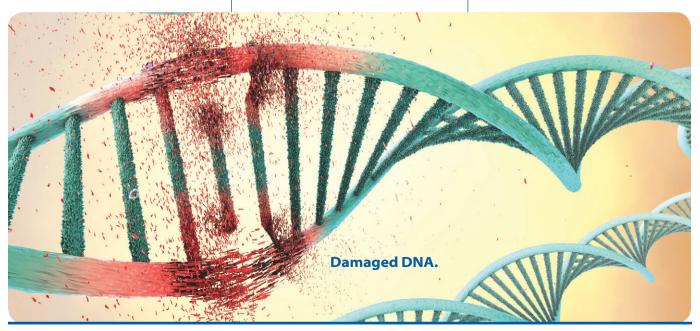
NAD+ Benefits More Than Just Sirtuins

The favorable effect of resveratrol in promoting sirtuin activity is well established. For **sirtuins** to function properly, they must have sufficient NAD+ to fuel their activity.7,10

Protecting against pathological aging, however, requires more than securing sirtuin structurefunction. We must also ensure the following two types of **DNA** damage are repaired:

- **Single-strand** DNA breaks occur often and are usually fixed by nutrients that most of you supplement with today.
- Double-strand DNA breaks are more difficult to restore. Left unrepaired, doublestrand breaks create cellular havoc that can lead to systemic degeneration.

A critical *enzyme* that repairs double-strand DNA breaks is PARP1.46-48 For the PARP1 enzyme to <u>function</u> it requires lots of NAD+. 49,50



When it comes to protecting against **cancer**, a tumor suppressor called **p53** protects against runaway cell propagation.

NAD+ supports **p53** activation to help thwart malignant transformation. ⁵¹⁻⁵³

Magnitude of Daily DNA Damage

Few people understand the degree of daily damage inflicted to their cellular **DNA**.

To put this into perspective, a study analyzed how many double-stranded **DNA breaks** occur **per cell** each **day**. The number turned out to be **10** DNA breaks **per cell** every day.⁵⁴

Your cells require **NAD+** to facilitate repair of **DNA breaks**. Sufficient NAD+ is needed for the **PARP1** repair enzyme to function. ^{49,50,55}

Imagine every <u>dividing</u> cell in your body undergoing <u>ten</u> double-stranded **DNA breaks** per day and NOT being repaired because your **NAD+** is depleted from aging, or from outside abuse such as excess alcohol and toxic food ingestion.

It explains many degenerative pathologies that occur as aging cells lose their **NAD+**.

Repairing **DNA** breaks will probably go a long way towards preventing cells from turning **malignant**. That's because **NAD+** helps maintain **activity** of cell division regulators like **p53**.⁵¹⁻⁵³

Restoring Youthful Cell Functionality

As we age, beneficial **genes** that support cell health "turn off" while detrimental genes overexpress.

Nutrients like **curcumin** help suppress genes that generate system-wide **inflammation**.⁵⁶⁻⁵⁹

Likewise, **omega-3s**⁶⁰⁻⁶⁶ and **vitamin D**⁶⁷⁻⁷⁰ favorably impact hundreds of **genes** that protect against degenerative illnesses.

To <u>reverse</u> the accumulation of damage inflicted to cellular **DNA**, we should support the efficient function of PARP1 enzymes.

PARP1 facilitates **DNA repair** via multiple mechanisms.

Higher **NAD+** cell levels enable PARP1 to function properly.^{49,71,72}

Aging creates a chaotic environment in the brain that can make sound **sleep** difficult.⁷³ As **DNA** is **repaired**, we regain youthful cell **functionality** that can result in **improved** overall health.

Combining **resveratrol** with <u>more</u> **nicotinamide riboside** supports healthy cellular **NAD**+ levels,⁷³ which are important to support **anti-aging** enymes like **PARP1** and **BubR1**.^{71,74,75}

$$O = P - O$$

$$O = P$$

Nutrients That Facilitate DNA Repair

People seeking to extend their lifespans today avoid toxins (such as tobacco smoke and overcooked food) that damage DNA.

Vitamin D has been shown to play an important role in **DNA repair**, which helps explain why people with higher levels of vitamin D show lower rates of most degenerative diseases. 70,76-78

Folic acid also plays a role in maintaining certain **DNA repair** mechanisms.⁷⁹⁻⁸⁵

Many of the supplements we take daily help facilitate DNA repair. The box below provides a partial listing of these nutrients.

Up until now, no nutrient could accelerate **DNA** repair to the magnitude needed to induce possible agereversal benefits.

That may have changed based on data showing remarkable **DNA repair** occurring when the amount of NAD+ (*nico-tinamide adenine dinucleotide*) is <u>increased</u> in our cells.^{3,86}

DNA Repair Activators					
Fish Oil ⁸⁷⁻⁸⁹	Vitamin B12	⁹⁰ Vitamin E ⁹	1	Vitamin C92	Nicotinamide86
CoQ10 ⁹³	Zinc ⁹⁴	Magnesium ⁹	95,96	Selenium ⁹⁷	Polyphenols98-100
Grape Seed Ex	ktract ^{101,102}	Curcumin ^{103,104}	Carote	noids ¹⁰⁵⁻¹⁰⁷	Vitamin B6 ^{108,109}

BubR1 is an *enzyme* that protects against **chromosome instability**. According to one study, sustained high-level expression of **BubR1** "provides a unique opportunity to extend healthy lifespan".⁷⁵

Some of you may find these new acronyms like **PARP1** a bit confusing.

I hope you appreciate (as I do) how rapidly our understanding of **aging** is expanding, along with accessible ways to <u>reverse</u> many degenerative changes.

Take Control by Boosting Your NAD+

George Church, PhD, is a Harvard professor pioneering **CRISPR/Cas9** gene editing technology.^{110,111}

Once perfected, Dr. Church has publicly stated that this will enable aging humans to "edit" their genes in a way that will empower them to regain **youth**.

We've reported on Dr. Church's research in past issues of *Life Extension Magazine*®. 110,111 This **age-reversal** gene-editing technology is predicted to be perfected in the next **5-10** years.

In the meantime, we can exert significant control over cellular health factors by taking <u>more</u> **nicotinamide riboside.** This will boost **NAD+** blood levels several fold.⁵

How our **genes** are **expressed** and their **stability** determines whether we retain healthy vitality or suffer relentless degeneration.

Nutrients like **curcumin**, ^{112,113} **fish oil**, ⁶⁰⁻⁶³ **folate**, ¹¹⁴⁻¹¹⁶ and **vitamin D**^{70,117-119} promote youthful genomic stability.

The box below displays additional benefits one can obtain by boosting cellular **NAD+**.

This can be accomplished by supplementing with **250 mg** each day of **nicotinamide riboside** that converts to **NAD+** in your body.

We may recommend higher **nicotinamide riboside** doses in coming months as scientific data emerges.

For longer life,

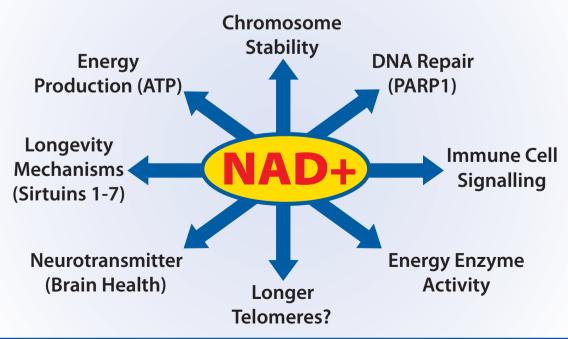
William Faloon, Co-Founder Life Extension Foundation Buyers Club

References

- Busso N, Karababa M, Nobile M, et al. Pharmacological inhibition of nicotinamide phosphoribosyltransferase/visfatin enzymatic activity identifies a new inflammatory pathway linked to NAD. PLoS One. 2008;3(5):e2267.
- Canto C, Menzies KJ, Auwerx J. NAD(+) Metabolism and the Control of Energy Homeostasis: A Balancing Act between Mitochondria and the Nucleus. *Cell Metab.* 2015;22(1):31-53.

(References continued on page 12.)

What is NAD+ Used For?





What Should Cancer Patients Being Treated With Chemo or Radiation Therapy Do?

Cancer chemotherapy drugs function via several destructive mechanisms, but the ultimate objective is to inflict massive damage to DNA so that cancer cells are destroyed. Radiation does this by directly breaking **DNA** strands.

One way cancer cells escape complete eradication after exposure to chemotherapy or radiation is to repair damaged DNA via a wide range of survival mechanisms.

Some studies suggest adding "DNA repair inhibitor" drugs might enable conventional chemo/radiation therapies to kill more cancer cells.

The downside to **DNA repair inhibitors** is they might increase the **toxicity** of chemo/radiation therapy to healthy cells and thus create more serious side effects. To cite a conclusion from a published study on this topic:

"With the addition of DNA repair inhibitors, standard chemotherapy could become more effective but also more toxic."120

What the above conclusion alludes to is that adding drugs that impede **DNA repair** might make **chemo**therapy more effective, but in the process make the chemotherapy more toxic. A major limiting factor to chemotherapy is **toxicity** so <u>severe</u> that patients are forced to discontinue therapy even when it is demonstrating efficacy.

For example, one of many **toxic** side effects of chemotherapy is painful neuropathy. 121-124 Cancer patients who take steps to boost their NAD+ levels have experienced relief from fatigue.125 Animals given nicotinamide riboside experienced reductions in chemotherapy-induced neuropathy.126

Another side effect of certain chemotherapy drugs and radiation to the chest is **heart failure**. 127-132 Nutrients like coenzyme Q10 have been shown to protect against this cardiac damage 133-135 and improve survival in cancer patients. 136-139 Conventional oncologists are largely unaware of this clinical research.

As it relates to supplementation with higher-dose nicotinamide riboside, we've reviewed numerous published studies and it is not possible to reach a rational consensus as to what actively-treated cancer patients should do as it relates to boosting their cellular **NAD+**.

Out of an abundance of caution, we suggest cancer patients undergoing chemotherapy or radiation avoid higher-dose NAD+ during therapy and for a reasonable period after.

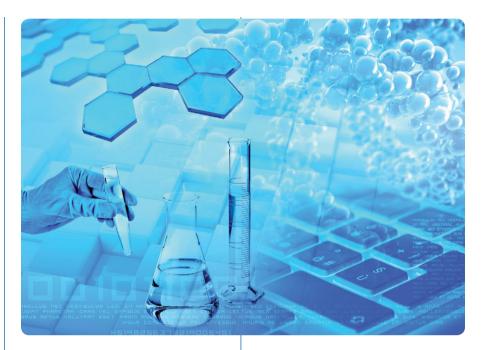
The conundrum cancer patients have faced for decades are arguments from many oncologists to take no supplements during chemo/radiation therapy. The concern is the nutrients might protect malignant cells from destruction.

Opposing this conventional view are numerous studies demonstrating that nutritional and herbal supplements do not interfere with the effectiveness of chemotherapy or radiation therapy. Furthermore, a myriad of controlled studies show marked survival **improvements** when cancer patients supplement with **nutrients** that boost immune function and protect against treatment side effects. 140-159

Stated simply, malignant cells can preferentially "hijack" many of the same factors healthy cells require for survival. There is thus a delicate theoretical balance as to what cancer patients should do during conventional treatment. For updated guidance on nutrients, hormones and off-label drugs that actively-treated cancer patients should consider, refer to our updated protocols at:

> www.LifeExtension.com/Chemotherapy and www.LifeExtension.com/Radiation

- Satoh MS, Poirier GG, Lindahl T. NAD(+)-dependent repair of damaged DNA by human cell extracts. J Biol Chem. 1993;268(8):5480-7.
- Available at: http://www.lifeextension. com/Magazine/2014/11/The-Youth-Restoring-Benefits-Of-NAD/Page-01. Accessed June 15, 2017.
- Trammell SA, Schmidt MS, Weidemann BJ, et al. Nicotinamide riboside is uniquely and orally bioavailable in mice and humans. Nat Commun. 2016;7:12948.
- Canto C, Houtkooper RH, Pirinen E, et al. The NAD(+) precursor nicotinamide riboside enhances oxidative metabolism and protects against high-fat diet-induced obesity. Cell Metab. 2012;15(6):838-47.
- Canto C, Auwerx J. Targeting sirtuin 1 to improve metabolism: all you need is NAD(+)? Pharmacol Rev. 2012;64(1): 166-87.
- Lekli I, Ray D, Das DK. Longevity nutrients resveratrol, wines and grapes. Genes Nutr. 2010;5(1):55-60.
- Landry J, Sutton A, Tafrov ST, et al. The silencing protein SIR2 and its homologs are NAD-dependent protein deacetylases. Proc Natl Acad Sci U S A. 2000;97(11):5807-11.
- 10. Villalba JM, Alcain FJ. Sirtuin activators and inhibitors. Biofactors. 2012;38(5):349-59.
- 11. Guest J, Grant R, Mori TA, et al. Changes in oxidative damage, inflammation and [NAD(H)] with age in cerebrospinal fluid. PLoS One. 2014;9(1):e85335.
- 12. Massudi H, Grant R, Braidy N, et al. Ageassociated changes in oxidative stress and NAD+ metabolism in human tissue. PLoS One. 2012;7(7):e42357.
- Stein LR, Imai S. Specific ablation of Nampt in adult neural stem cells recapitulates their functional defects during aging. Embo j. 2014;33(12):1321-40.
- 14. Braidy N, Guillemin GJ, Mansour H, et al. Age related changes in NAD+ metabolism oxidative stress and Sirt1 activity in wistar rats. PLoS One. 2011;6(4):e19194.
- 15. Available at: http://www.lifeextension. com/Magazine/2008/2/Living-Longer-Healthier-Lives-With-Resveratrol/Page-01. Accessed June 15, 2015.
- 16. Available at: http://www.lifeextension. com/magazine/2007/3/report_resveratrol/ page-01. Accessed June 15, 2017.
- 17. Available at: http://www.lifeextension. com/magazine/2006/7/report_longevity/ Page-01. Accessed July 5, 2017.
- 18. Dash S, Xiao C, Morgantini C, et al. High-dose resveratrol treatment for 2 weeks inhibits intestinal and hepatic lipoprotein production in overweight/ obese men. Arterioscler Thromb Vasc Biol. 2013;33(12):2895-901.



- 19. Kennedy DO, Wightman EL, Reay JL, et al. Effects of resveratrol on cerebral blood flow variables and cognitive performance in humans: a double-blind. placebo-controlled, crossover investigation. Am J Clin Nutr. 2010;91(6):1590-7.
- 20. Imai S, Guarente L. NAD+ and sirtuins in aging and disease. Trends Cell Biol. 2014:24(8):464-71.
- 21. Liu D, Pitta M, Mattson MP. Preventing NAD(+) depletion protects neurons against excitotoxicity: bioenergetic effects of mild mitochondrial uncoupling and caloric restriction. Ann NY Acad Sci. 2008;1147:275-82.
- 22. Min SW, Sohn PD, Cho SH, et al. Sirtuins in neurodegenerative diseases: an update on potential mechanisms. Front Aging Neurosci. 2013;5:53.
- 23. Anekonda TS, Reddy PH. Neuronal protection by sirtuins in Alzheimer's disease. J Neurochem. 2006:96(2):305-13.
- 24. Borradaile NM, Pickering JG. NAD(+), sirtuins, and cardiovascular disease. Curr Pharm Des. 2009;15(1):110-7.
- Hsu CP, Oka S, Shao D, et al. Nicotinamide phosphoribosyltransferase regulates cell survival through NAD+ synthesis in cardiac myocytes. Circ Res. 2009;105(5):481-91.
- 26. Pillai VB. Sundaresan NR. Kim G. et al. Exogenous NAD blocks cardiac hypertrophic response via activation of the SIRT3-LKB1-AMP-activated kinase pathway. J Biol Chem. 2010;285(5):3133-
- 27. Frederick DW, Loro E, Liu L, et al. Loss of NAD Homeostasis Leads to Progressive and Reversible Degeneration of Skeletal Muscle. Cell Metab. 2016;24(2):269-82.

- 28. Zhou CC, Yang X, Hua X, et al. Hepatic NAD(+) deficiency as a therapeutic target for non-alcoholic fatty liver disease in ageing. Br J Pharmacol. 2016;173(15):2352-68.
- 29. Zhao Y, Guan YF, Zhou XM, et al. Regenerative Neurogenesis After Ischemic Stroke Promoted by Nicotinamide Phosphoribosyltransferase-Nicotinamide Adenine Dinucleotide Cascade. Stroke. 2015;46(7):1966-74.
- Wang S, Xing Z, Vosler PS, et al. Cellular NAD replenishment confers marked neuroprotection against ischemic cell death: role of enhanced DNA repair. Stroke. 2008;39(9):2587-95.
- 31. Imai S. "Clocks" in the NAD World: NAD as a metabolic oscillator for the regulation of metabolism and aging. Biochim Biophys Acta. 2010;1804(8):1584-90.
- 32. Yoshino J, Mills KF, Yoon MJ, et al. Nicotinamide mononucleotide, a key NAD(+) intermediate, treats the pathophysiology of diet- and age-induced diabetes in mice. Cell Metab. 2011;14(4):528-36.
- Trammell SA, Weidemann BJ, Chadda A, et al. Nicotinamide Riboside Opposes Type 2 Diabetes and Neuropathy in Mice. Sci Rep. 2016;6:26933.
- 34. Higashida K, Kim SH, Jung SR, et al. Effects of resveratrol and SIRT1 on PGC-1alpha activity and mitochondrial biogenesis: a reevaluation. PLoS Biol. 2013;11(7):e1001603.
- 35. Rehan L, Laszki-Szczachor K, Sobieszczanska M, et al. SIRT1 and NAD as regulators of ageing. Life Sci. 2014;105 $(1-2)\cdot 1-6$
- 36. Imai S-i, Guarente L. It takes two to tango: NAD+ and sirtuins in aging/ longevity control. NPJ Aging Mech Dis. 2016;2:16017.

- 37. Mouchiroud L, Houtkooper RH, Moullan N, et al. The NAD(+)/Sirtuin Pathway Modulates Longevity through Activation of Mitochondrial UPR and FOXO Signaling. Cell. 2013;154(2):430-41.
- Kaeberlein M, McVey M, Guarente L. The SIR2/3/4 complex and SIR2 alone promote longevity in Saccharomyces cerevisiae by two different mechanisms. Genes Dev. 1999;13(19):2570-80.
- 39. Rogina B, Helfand SL. Sir2 mediates longevity in the fly through a pathway related to calorie restriction. Proc Natl Acad Sci U S A. 2004;101(45):15998-6003.
- 40. Giblin W. Skinner MF. Lombard DB. Sirtuins: guardians of mammalian healthspan. Trends Genet. 2014;30(7):271-86.
- 41. Satoh A, Brace CS, Rensing N, et al. Sirt1 extends life span and delays aging in mice through the regulation of Nk2 homeobox 1 in the DMH and LH. Cell Metab. 2013;18(3):416-30.
- 42. Satoh A, Imai S. Systemic regulation of mammalian ageing and longevity by brain sirtuins. Nat Commun. 2014:5:4211.
- 43. Araki T, Sasaki Y, Milbrandt J. Increased nuclear NAD biosynthesis and SIRT1 activation prevent axonal degeneration. Science. 2004;305(5686):1010-3.
- 44. Schmeisser K, Mansfeld J, Kuhlow D, et al. Role of sirtuins in lifespan regulation is linked to methylation of nicotinamide. Nat Chem Biol. 2013;9(11):693-700.
- 45. Haigis MC, Sinclair DA. Mammalian sirtuins: biological insights and disease relevance. Annu Rev Pathol. 2010;5:253-95.
- 46. Mao Z, Hine C, Tian X, et al. SIRT6 promotes DNA repair under stress by activating PARP1. Science. 2011;332(6036):1443-6.
- 47. Hochegger H, Dejsuphong D, Fukushima T, et al. Parp-1 protects homologous recombination from interference by Ku and Ligase IV in vertebrate cells. Embo j. 2006:25(6):1305-14.
- 48. Paddock MN, Buelow BD, Takeda S, et al. The BRCT domain of PARP-1 is required for immunoglobulin gene conversion. PLoS Biol. 2010;8(7):e1000428.
- 49. Kim MY, Zhang T, Kraus WL. Poly(ADPribosyl)ation by PARP-1: 'PAR-laying' NAD+ into a nuclear signal. Genes Dev. 2005;19(17):1951-67.
- 50. Siegel C. McCullough LD. NAD+ depletion or PAR polymer formation: which plays the role of executioner in ischaemic cell death? Acta Physiol (Oxf). 2011;203(1):225-34.
- 51. Langley E, Pearson M, Faretta M, et al. Human SIR2 deacetylates p53 and antagonizes PML/p53-induced cellular senescence. Embo j. 2002;21(10):2383-96.
- 52. Pfister NT, Yoh KE, Prives C. p53, DNA damage, and NAD+ homeostasis. Cell Cycle. 2014;13(11):1661-2.
- 53. McLure KG, Takagi M, Kastan MB. NAD+ modulates p53 DNA binding specificity and function. Mol Cell Biol. 2004;24(22):9958-67.

- 54. Lieber MR. The mechanism of doublestrand DNA break repair by the nonhomologous DNA end-joining pathway. Annu Rev Biochem. 2010;79:181-211.
- 55. El Ramy R, Magroun N, Messadecq N, et al. Functional interplay between Parp-1 and SirT1 in genome integrity and chromatin-based processes. Cell Mol Life Sci. 2009;66(19):3219-34.
- Singh S, Aggarwal BB. Activation of transcription factor NF-kappa B is suppressed by curcumin (diferuloylmethane). J Biol Chem. 1995;270(42):24995-5000.
- 57. Ranjan D, Chen C, Johnston TD, et al. Curcumin inhibits mitogen stimulated lymphocyte proliferation, NFkappaB activation, and IL-2 signaling. J Surg Res. 2004;121(2):171-7.
- 58. Aggarwal BB, Shishodia S. Suppression of the nuclear factor-kappaB activation pathway by spice-derived phytochemicals: reasoning for seasoning. Ann NY Acad Sci. 2004;1030:434-41.
- 59. Kumar A, Dhawan S, Hardegen NJ, et al. Curcumin (Diferulovlmethane) inhibition of tumor necrosis factor (TNF)-mediated adhesion of monocytes to endothelial cells by suppression of cell surface expression of adhesion molecules and of nuclear factorkappaB activation. Biochem Pharmacol. 1998;55(6):775-83.
- 60. Bouwens M, van de Rest O, Dellschaft N, et al. Fish-oil supplementation induces antiinflammatory gene expression profiles in human blood mononuclear cells. Am J Clin Nutr. 2009;90(2):415-24.
- 61. Weaver KL, Ivester P, Seeds M, et al. Effect of dietary fatty acids on inflammatory gene expression in healthy humans. J Biol Chem. 2009;284(23):15400-7.
- 62. Clarke SD. Polyunsaturated fatty acid regulation of gene transcription: a molecular mechanism to improve the metabolic syndrome. J Nutr. 2001;131(4):1129-32.
- 63. Gillies PJ, Bhatia SK, Belcher LA, et al. Regulation of inflammatory and lipid metabolism genes by eicosapentaenoic acid-rich oil. J Lipid Res. 2012;53(8):1679-89.
- 64. Thomas J, Thomas CJ, Radcliffe J, et al. Omega-3 Fatty Acids in Early Prevention of Inflammatory Neurodegenerative Disease: A Focus on Alzheimer's Disease. Biomed Res Int. 2015;2015:172801.
- 65. Dyall SC, Michael GJ, Michael-Titus AT. Omega-3 fatty acids reverse agerelated decreases in nuclear receptors and increase neurogenesis in old rats. JNeurosci Res. 2010;88(10):2091-102.
- 66. Kitajka K, Puskas LG, Zvara A, et al. The role of n-3 polyunsaturated fatty acids in brain: modulation of rat brain gene expression by dietary n-3 fatty acids. Proc Natl Acad Sci U S A. 2002;99(5):2619-24.
- Chakraborti CK. Vitamin D as a promising anticancer agent. Indian J Pharmacol. 2011;43(2):113-20.

- 68. Li H, Stampfer MJ, Hollis JB, et al. A prospective study of plasma vitamin D metabolites, vitamin D receptor polymorphisms, and prostate cancer. PLoS Med. 2007;4(3):e103.
- Bao BY, Yao J, Lee YF. 1alpha, 25-dihydroxyvitamin D3 suppresses interleukin-8-mediated prostate cancer cell angiogenesis. Carcinogenesis. 2006;27(9):1883-93.
- 70. Hossein-nezhad A, Spira A, Holick MF. Influence of vitamin D status and vitamin D3 supplementation on genome wide expression of white blood cells: a randomized double-blind clinical trial. PLoS One. 2013;8(3):e58725.
- 71. Schiewer MJ, Knudsen KE, Transcriptional roles of PARP1 in cancer. Mol Cancer Res. 2014;12(8):1069-80.
- 72. 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.
- 73. Mouchiroud L, Houtkooper RH, Auwerx J. NAD(+) metabolism, a therapeutic target for age-related metabolic disease. Critical reviews in biochemistry and molecular biology. 2013;48(4):10.3109/10 409238.2013.789479.
- 74. North BJ, Rosenberg MA, Jeganathan KB, et al. SIRT2 induces the checkpoint kinase BubR1 to increase lifespan. The EMBO Journal. 2014;33(13):1438-53.
- 75. Baker DJ, Dawlaty MM, Wijshake T, et al. Increased expression of BubR1 protects against aneuploidy and cancer and extends healthy lifespan. Nat Cell Biol. 2013:15(1):96-102.
- 76. Halicka HD, Zhao H, Li J, et al. Attenuation of constitutive DNA damage signaling by 1,25-dihydroxyvitamin D3. Aging (Albany NY). 2012;4(4):270-8.
- 77. Available at: http://www.lifeextension. com/magazine/2013/8/The-Overlooked-Importance-of-Vitamin-D-Receptors/ Page-01. Accessed June 16, 2017.
- 78. Fleet JC, DeSmet M, Johnson R, et al. Vitamin D and cancer: a review of molecular mechanisms. Biochem L. 2012;441(1):61-76.
- 79. Ames BN. A role for supplements in optimizing health: the metabolic tune-up. Arch Biochem Biophys. 2004;423(1):
- 80. Wei O. Shen H. Wang LE, et al. Association between low dietary folate intake and suboptimal cellular DNA repair capacity. Cancer Epidemiol Biomarkers Prev. 2003;12(10):963-9.
- 81. Duthie SJ. Folate and cancer: how DNA damage, repair and methylation impact on colon carcinogenesis. J Inherit Metab Dis. 2011;34(1):101-9.
- 82. Basten GP. Duthie SJ. Pirie L. et al. Sensitivity of markers of DNA stability and DNA repair activity to folate supplementation in healthy volunteers. Br J Cancer. 2006;94(12):1942-7.

- 83. Choi SW, Kim YI, Weitzel JN, et al. Folate depletion impairs DNA excision repair in the colon of the rat. Gut. 1998:43(1):93-9.
- 84. Kruman, II, Kumaravel TS, Lohani A, et al. Folic acid deficiency and homocysteine impair DNA repair in hippocampal neurons and sensitize them to amyloid toxicity in experimental models of Alzheimer's disease. J Neurosci. 2002;22(5):1752-62.
- 85. Sadik NA, Shaker OG. Dietary folate suppresses DMH-induced colon carcinogenesis in a rat model and affects DMHinduced expression of four DNA repair enzymes. Nutr Cancer. 2012;64(8):1196-
- 86. Surjana D, Halliday GM, Damian DL. Role of Nicotinamide in DNA Damage, Mutagenesis, and DNA Repair. Journal of Nucleic Acids. 2010;2010:13.
- 87. Hong MY, Lupton JR, Morris JS, et al. Dietary fish oil reduces O6-methylguanine DNA adduct levels in rat colon in part by increasing apoptosis during tumor initiation. Cancer Epidemiol Biomarkers Prev. 2000;9(8):819-26.
- 88. Ghorbanihaghjo A, Safa J, Alizadeh S, et al. Protective effect of fish oil supplementation on DNA damage induced by cigarette smoking. J Health Popul Nutr. 2013;31(3):343-9.
- 89. Stephenson JA, Al-Taan O, Arshad A, et al. The multifaceted effects of omega-3 polyunsaturated Fatty acids on the hallmarks of cancer. J Lipids. 2013;2013:261247.
- 90. Alzoubi K, Khabour O, Hussain N, et al. Evaluation of vitamin B12 effects on DNA damage induced by pioglitazone. Mutat Res. 2012;748(1-2):48-51.
- 91. Sweetman SF, Strain JJ, McKelvey-Martin VJ. Effect of antioxidant vitamin supplementation on DNA damage and repair in human lymphoblastoid cells. Nutr Cancer. 1997;27(2):122-30.
- 92. Cooke MS, Evans MD, Podmore ID, et al. Novel repair action of vitamin C upon in vivo oxidative DNA damage. FEBS Lett. 1998;439(3):363-7.
- 93. Tomasetti M, Alleva R, Borghi B, et al. In vivo supplementation with coenzyme Q10 enhances the recovery of human lymphocytes from oxidative DNA damage. Faseb j. 2001;15(8):1425-7.
- 94. Song Y, Leonard SW, Traber MG, et al. Zinc deficiency affects DNA damage, oxidative stress, antioxidant defenses, and DNA repair in rats. J Nutr. 2009;139(9):1626-31.
- 95. Hartwig A. Role of magnesium in genomic stability. Mutat Res. 2001;475(1-2):113-21.
- Mahabir S, Wei Q, Barrera SL, et al. Dietary magnesium and DNA repair capacity as risk factors for lung cancer. Carcinogenesis. 2008;29(5):949-56.

- 97. de Rosa V, Erkekoglu P, Forestier A, et al. Low doses of selenium specifically stimulate the repair of oxidative DNA damage in LNCaP prostate cancer cells. Free Radic Res. 2012;46(2):105-16.
- 98. Nichols JA, Katiyar SK. Skin photoprotection by natural polyphenols: anti-inflammatory, antioxidant and DNA repair mechanisms. Arch Dermatol Res. 2010;302(2):71-83.
- 99. Zattra E, Coleman C, Arad S, et al. Polypodium leucotomos extract decreases UV-induced Cox-2 expression and inflammation, enhances DNA repair, and decreases mutagenesis in hairless mice. Am J Pathol. 2009;175(5):1952-61.
- 100. Tan X, Zhao C, Pan J, et al. In vivo non-enzymatic repair of DNA oxidative damage by polyphenols. Cell Biol Int. 2009;33(6):690-6.
- 101. Katiyar SK, van Steeg H, Sharma SD. Abstract 1875: Dietary grape seed proanthocyanidins induce rapid repair of DNA damage via nucleotide excision repair genes in preventing UV-induced immunosuppression. Cancer Research. 2010;70(8 Supplement):1875-.
- 102. Mansouri E, Khorsandi L, Abedi HA. Antioxidant effects of proanthocyanidin from grape seed on hepatic tissue injury in diabetic rats. Iran J Basic Med Sci. 2014;17(6):460-4.
- 103. Roy M, Sinha D, Mukherjee S, et al. Curcumin prevents DNA damage and enhances the repair potential in a chronically arsenic-exposed human population in West Bengal, India. Eur J Cancer Prev. 2011;20(2):123-31.
- 104. Mukherjee S, Roy M, Dev S, et al. A Mechanistic Approach for Modulation of Arsenic Toxicity in Human Lymphocytes by Curcumin, an Active Constituent of Medicinal Herb Curcuma longa Linn. J Clin Biochem Nutr. 2007;41(1):32-42.
- 105. Astley SB, Elliott RM, Archer DB, et al. Increased cellular carotenoid levels reduce the persistence of DNA singlestrand breaks after oxidative challenge. Nutr Cancer. 2002;43(2):202-13.
- 106. Lorenzo Y, Azgueta A, Luna L, et al. The carotenoid beta-cryptoxanthin stimulates the repair of DNA oxidation damage in addition to acting as an antioxidant in human cells. Carcinogenesis. 2009;30(2):308-14.
- 107. Astley SB, Elliott RM, Archer DB, et al. Evidence that dietary supplementation with carotenoids and carotenoid-rich foods modulates the DNA damage: repair balance in human lymphocytes. Br J Nutr. 2004;91(1):63-72.
- 108. Chou YC, Chu CH, Wu MH, et al. Dietary intake of vitamin B(6) and risk of breast cancer in Taiwanese women. J Epidemiol. 2011;21(5):329-36.
- 109. Le Marchand L, White KK, Nomura AM, et al. Plasma levels of B vitamins and colorectal cancer risk: the multiethnic cohort study. Cancer Epidemiol Biomarkers Prev. 2009;18(8):2195-201.

- 110. Available at: http://www.lifeextension. com/Magazine/2014/6/The-2013-SENS-Foundation-Conference/Page-01. Accessed June 16, 2017.
- 111. Available at: http://www.lifeextension. com/Magazine/2016/7/Age-Reversal-Research-at-Harvard-Medical-School/ Page-01. Accessed June 16, 2017.
- 112. Thomas P, Wang YJ, Zhong JH, et al. Grape seed polyphenols and curcumin reduce genomic instability events in a transgenic mouse model for Alzheimer's disease. Mutat Res. 2009;661(1-2):25-34.
- 113. Shu L, Khor TO, Lee JH, et al. Epigenetic CpG demethylation of the promoter and reactivation of the expression of Neurog1 by curcumin in prostate LNCaP cells. Aaps j. 2011;13(4):606-14.
- 114. Bistulfi G, Vandette E, Matsui S, et al. Mild folate deficiency induces genetic and epigenetic instability and phenotype changes in prostate cancer cells. BMC Biol. 2010:8:6.
- 115. Duthie SJ, Narayanan S, Sharp L, et al. Folate, DNA stability and colo-rectal neoplasia. Proc Nutr Soc. 2004;63(4):571-
- 116. James SJ, Pogribny IP, Pogribna M, et al. Mechanisms of DNA damage, DNA hypomethylation, and tumor progression in the folate/methyl-deficient rat model of hepatocarcinogenesis. J Nutr. 2003;133(11 Suppl 1):3740s-7s.
- 117. Palmer HG, Sanchez-Carbayo M, Ordonez-Moran P, et al. Genetic signatures of differentiation induced by 1alpha,25-dihydroxyvitamin D3 in human colon cancer cells. Cancer Res. 2003;63(22):7799-806.
- 118. Fernandez-Garcia NI, Palmer HG, Garcia M, et al. 1alpha,25-Dihydroxyvitamin D3 regulates the expression of Id1 and Id2 genes and the angiogenic phenotype of human colon carcinoma cells. Oncogene. 2005;24(43):6533-44.
- 119. Stefanska B. Salame P. Bednarek A. et al. Comparative effects of retinoic acid, vitamin D and resveratrol alone and in combination with adenosine analogues on methylation and expression of phosphatase and tensin homologue tumour suppressor gene in breast cancer cells. Br J Nutr. 2012;107(6):781-90.
- 120. Kelley MR, Logsdon D, Fishel ML. Targeting DNA repair pathways for cancer treatment: what's new? Future Oncol. 2014;10(7):1215-37.
- 121. Park HJ. Chemotherapy induced peripheral neuropathic pain. Korean J Anesthesiol. 2014;67(1):4-7.
- 122. Piccolo J, Kolesar JM. Prevention and treatment of chemotherapy-induced peripheral neuropathy. Am J Health Syst Pharm. 2014;71(1):19-25.
- 123. Visovsky C, Collins M, Abbott L, et al. Putting evidence into practice: evidencebased interventions for chemotherapyinduced peripheral neuropathy. Clin J Oncol Nurs. 2007;11(6):901-13.

- 124. Stubblefield MD, McNeely ML, Alfano CM, et al. A prospective surveillance model for physical rehabilitation of women with breast cancer: chemotherapy-induced peripheral neuropathy. Cancer. 2012;118(8 Suppl):2250-60.
- 125. Feedback from chemotherapy patients using nicotinamide riboside dietary supplements. Life Extension Wellness Specialist Report. June 2017.
- 126. Hamity MV, White SR, Walder RY, et al. Nicotinamide riboside, a form of vitamin B3 and NAD+ precursor, relieves the nociceptive and aversive dimensions of paclitaxel-induced peripheral neuropathy in female rats. Pain. 2017;158(5): 962-72.
- 127. Available at: http://www.health.harvard. edu/heart-health/cancer-treatments-mayharm-the-heart. Accessed June 16, 2017.
- 128. Available at: https://www.texasoncology. com/cancer-treatment/side-effects-ofcancer-treatment/long-term-side-effects/ cardiac-toxicity. Accessed June 16, 2017.
- 129. Svoboda M, Poprach A, Dobes S, et al. Cardiac toxicity of targeted therapies used in the treatment for solid tumours: a review. Cardiovasc Toxicol. 2012:12(3):191-207.
- 130. Marinko T, Dolenc J, Bilban-Jakopin C. Cardiotoxicity of concomitant radiotherapy and trastuzumab for early breast cancer. Radiol Oncol. 2014;48(2):105-12.
- 131. Darby SC, Ewertz M, McGale P, et al. Risk of ischemic heart disease in women after radiotherapy for breast cancer. N Engl J Med. 2013;368(11):987-98.
- 132. Yusuf SW, Sami S, Daher IN. Radiationinduced heart disease: a clinical update. Cardiol Res Pract. 2011;2011:317659.
- 133. Cortes EP, Gupta M, Chou C, et al. Adriamycin cardiotoxicity: early detection by systolic time interval and possible prevention by coenzyme Q10. Cancer Treat Rep. 1978;62(6):887-91.
- 134. Iarussi D. Auricchio U. Agretto A. et al. Protective effect of coenzyme Q10 on anthracyclines cardiotoxicity: control study in children with acute lymphoblastic leukemia and non-Hodgkin lymphoma. Mol Aspects Med. 1994;15 Suppl:s207-12.
- 135. Albini A, Pennesi G, Donatelli F, et al. Cardiotoxicity of anticancer drugs: the need for cardio-oncology and cardiooncological prevention. J Natl Cancer Inst. 2010;102(1):14-25.
- 136. Rusciani L, Proietti I, Paradisi A, et al. Recombinant interferon alpha-2b and coenzyme Q10 as a postsurgical adjuvant therapy for melanoma: a 3-year trial with recombinant interferon-alpha and 5-year follow-up. Melanoma Res. 2007;17(3):177-83.
- 137. Folkers K. Brown R. Judy WV. et al. Survival of cancer patients on therapy with coenzyme Q10. Biochem Biophys Res Commun. 1993;192(1):241-5.

- 138. Lockwood K, Moesgaard S, Folkers K. Partial and complete regression of breast cancer in patients in relation to dosage of coenzyme Q10. Biochem Biophys Res Commun. 1994;199(3):1504-8.
- 139. Lockwood K, Moesgaard S, Hanioka T, et al. Apparent partial remission of breast cancer in 'high risk' patients supplemented with nutritional antioxidants, essential fatty acids and coenzyme Q10. Mol Aspects Med. 1994;15 Suppl:s231-40.
- 140. Smyth JF, Bowman A, Perren T, et al. Glutathione reduces the toxicity and improves quality of life of women diagnosed with ovarian cancer treated with cisplatin: results of a double-blind, randomised trial. Ann Oncol. 1997;8(6):569-
- 141. Lissoni P, Chilelli M, Villa S, et al. Five years survival in metastatic non-small cell lung cancer patients treated with chemotherapy alone or chemotherapy and melatonin: a randomized trial. JPineal Res. 2003;35(1):12-5.
- 142. Israel L, Hajji O, Grefft-Alami A, et al. Vitamin A augmentation of the effects of chemotherapy in metastatic breast cancers after menopause. Randomized trial in 100 patients. Ann Med Interne (Paris). 1985;136(7):551-4.
- 143. Zou YH, Liu XM. Effect of astragalus injection combined with chemotherapy on quality of life in patients with advanced non-small cell lung cancer. Zhongguo Zhong Xi Yi Jie He Za Zhi. 2003;23(10):733-5.
- 144. Beer TM, Ryan CW, Venner PM, et al. Double-blinded randomized study of high-dose calcitriol plus docetaxel compared with placebo plus docetaxel in androgen-independent prostate cancer: a report from the ASCENT Investigators. J Clin Oncol. 2007;25(6):669-74.
- 145. Pace A. Giannarelli D. Galie E. et al. Vitamin E neuroprotection for cisplatin neuropathy: a randomized, placebo-controlled trial. Neurology. 2010;74(9):762-6.
- 146. Rouse K, Nwokedi E, Woodliff JE, et al. Glutamine enhances selectivity of chemotherapy through changes in glutathione metabolism. Ann Surg. 1995;221(4):420-6.
- 147. Kim SR, Jo SK, Kim SH. Modification of radiation response in mice by ginsenosides, active components of Panax ginseng. In Vivo. 2003;17(1):77-81.
- 148. Xie FY, Zeng ZF, Huang HY. Clinical observation on nasopharyngeal carcinoma treated with combined therapy of radiotherapy and ginseng polysaccharide injection. Zhongguo Zhong Xi Yi Jie He Za Zhi. 2001;21(5):332-4.
- 149. Kiremidjian-Schumacher L, Roy M, Glickman R. et al. Selenium and immunocompetence in patients with head and neck cancer. Biol Trace Elem Res. 2000;73(2):97-111.

- 150. Malmberg KJ, Lenkei R, Petersson M, et al. A short-term dietary supplementation of high doses of vitamin E increases T helper 1 cytokine production in patients with advanced colorectal cancer. Clin Cancer Res. 2002;8(6):1772-8.
- 151. Kumar B, Jha MN, Cole WC, et al. Dalpha-tocopheryl succinate (vitamin E) enhances radiation-induced chromosomal damage levels in human cancer cells, but reduces it in normal cells. J Am Coll Nutr. 2002;21(4):339-43.
- 152. Kennedy RS, Konok GP, Bounous G, et al. The use of a whey protein concentrate in the treatment of patients with metastatic carcinoma: a phase I-II clinical study. Anticancer Res. 1995;15(6b):2643-9.
- 153. Todorova VK, Harms SA, Kaufmann Y, et al. Effect of dietary glutamine on tumor glutathione levels and apoptosisrelated proteins in DMBA-induced breast cancer of rats. Breast Cancer Res Treat. 2004;88(3):247-56.
- 154. Dorai T, Aggarwal BB. Role of chemopreventive agents in cancer therapy. Cancer Lett. 2004;215(2):129-40.
- 155. Hillman GG, Wang Y, Kucuk O, et al. Genistein potentiates inhibition of tumor growth by radiation in a prostate cancer orthotopic model. Mol Cancer Ther. 2004;3(10):1271-9.
- 156. Yashar CM, Spanos WJ, Taylor DD, et al. Potentiation of the radiation effect with genistein in cervical cancer cells. Gynecol Oncol. 2005;99(1):199-205.
- 157. Kotowski U, Heiduschka G, Brunner M, et al. Radiosensitization of head and neck cancer cells by the phytochemical agent sulforaphane. Strahlenther Onkol. 2011;187(9):575-80.
- 158. He H, Zhou X, Wang Q, et al. Does the couse of astragalus-containing chinese herbal prescriptions and radiotherapy benefit to non-small-cell lung cancer treatment: a meta-analysis of randomized trials. Evid Based Complement Alternat Med. 2013;2013:426207.
- 159. Kwan ML, Greenlee H, Lee VS, et al. Multivitamin use and breast cancer outcomes in women with early-stage breast cancer: the Life After Cancer Epidemiology study. Breast Cancer Res Treat. 2011;130(1):195-205.

The incredible health nutrient

CURCUMIN

in a super <u>absorbable</u> formula.

Super Bio-Curcumin® delivers a patented turmeric extract that is up to seven times more absorbable and lasts longer in your bloodstream.

NON-GMO

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.



For full product description and to order Super Bio-Curcumin®, call 1-800-544-4440 or visit www.LifeExtension.com

Item #00407 • 400 mg • 60 veg. caps Retail Your

Price \$38.00 1-bottle

Price

4-bottle

\$28.50 \$26.25 ea

Suggested dose is one capsule daily.



For full product description and to order **Super K with Advanced K2 Complex** or **Once-Daily Health Booster**, call **1-800-544-4440** or visit **www.LifeExtension.com**

Vitamin K1, vitamin K2 (MK-4), and vitamin K2 (MK-7) can also be found in Life Extension® Once-Daily Health Booster.

If you take Once-Daily Health Booster, you do not need additional Super K with Advanced K2 formula.

Warning to Coumadin® (warfarin) Drug Users: Patients prescribed vitamin K-antagonist anticoagulant prescription drugs like warfarin should consult their physician before taking vitamin K supplements like Super K and Super Booster. There is evidence, however, that users of drugs like warfarin could benefit from a consistent low dose of supplemental K. Ask your doctor if you can take a low dose (45 mcg a day) of vitamin K2 in the long-acting MK-7 form for the purpose of stabilizing your INR levels and also protecting your body against long-term vitamin K deficit. Do not initiate any form of vitamin K supplementation without full cooperation of your treating doctor, as your doctor may need to increase your dose of warfarin to compensate for your vitamin K supplement. Life Extension® provides several forms of low-dose vitamin K for physician consideration.



Higher Potency
NICOTINAMIDE
RIBOSIDE
Lower
Cost-Per-Milligram

When you ingest **nicotinamide riboside**, it converts to **NAD**+ in your body.

NAD+ is essential for cell **energy** production and **DNA function**. As we age, NAD+ levels markedly <u>decline</u>.

Save more than 30% with <u>new</u> higher-potency NAD+ Cell Regenerator™.



For full product description and to order the <u>new</u> NAD+ Cell Regenerator™, call 1-800-544-4440 or visit www.LifeExtension.com

Non-GMO	Price	Price	
1 bottle 4 bottles	\$42	\$31.50 \$28 each	

Item #02144 • 30 vegetarian capsules

NIAGEN® is a registered trademark of ChromaDex, Inc. Patents see: www.ChromaDexPatents.com.

POWER UP!

TURN ON YOUR LONGEVITY GENES

Optimized NAD+

Cell Regenerator™ with Resveratrol

NAD+ is essential for energy production and DNA function.

RESVERATROL promotes cell proteins called **sirtuins** that activate longevity **genes**.

Sirtuins are **dependent** on **NAD**+ for optimal function.

This <u>new</u> formula combines 250 mg of NAD+ precursor nicotinamide riboside with 100 mg of trans-resveratrol and other plant extracts in one daily capsule.



Retail Your Price

1 bottle \$50 \$37.50 \$4 bottles \$34 each

1 tem #02145 • 30 vegetarian capsules

For full product description and to order

Optimized NAD+ Cell Regenerator™

with Resveratrol, call 1-800-544-4440 or

visit www.LifeExtension.com

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



COQUINOL 10

BETTER ABSORPTION
WITH ADDED MITOCHONDRIAL SUPPORT

Non-GMO

LifeExtension
Super Ubiquinol
COO10
with followed Mitschondrial Superior
100 mg
Crihanced Delivery System
for Maximum Absorption
Superior

Item # 01426

For full product description and to order **Super Ubiquinol CoQ10**with Enhanced Mitochondrial Support,

call **1-800-544-4440** or visit **www.LifeExtension.com**

60 softgels • Retail Price is \$62 • Your price is \$46.50 • 4 bottles are only \$39 each

Q+®, Kaneka Ubiquinol™, and the quality seal™ are registered or pending trademarks of Kaneka Corp.

Boosting NAD+ Levels Slows Aging

A 2017 review of the literature has found that supplementation with NAD+ precursors **nicotinamide riboside** or **nicotinamide mononucleotide** increases lifespan in mice. It also improved their mitochondrial, brain, muscle, and melanocyte stem-cell function.*

In one of the papers discussed, researchers identified a protein that aids in DNA repair in young mice. The research shows NAD+ levels can be boosted through NAD+ precursor supplementation, reducing DNA damage and bringing cellular activity back to youthful levels.

Researchers believe the medical implications for <u>humans</u> indicate that supplementation with **nicotinamide riboside** at doses of **100-250 mg** or more can increase NAD+ levels systemically.

Editor's Note: The study's authors note that the exact mechanism of declining NAD+ levels and their basic importance to the aging process are still under investigation.

* Rejuvenation Res. 2017 May 24.

Attentive Diabetes Management Extends Life

Strict management of type II diabetes can make a significant difference in quality and length of life.*

A 20-year study divided 160 people—all of whom were at risk of type II diabetes—into two groups. One group stayed with their usual treatment, while the other changed to a more multitargeted, aggressive regimen.

Results showed the intensive-treatment group lived, on average, 7.9 years longer than the "normal" treatment group. Also, in the aggressive treatment group, the risk for a number of diseases (including kidney disease, heart disease, and blindness) was reduced.

When the study began, the average subject age was 55, and all were borderline obese.

According to senior study author Dr. Oluf Pedersen, the intensive treatment was aimed at reducing a comprehensive selection of adverse factors such as blood-clot risk, high glucose, high blood pressure, triglycerides and cholesterol. The regimen included behavior modification (exercise, healthy diet, no smoking) and medications when deemed necessary.

Editor's Note: Dr. Joel Zonszein, director of New York's Clinical Diabetes Center at Montefiore Medical Center, stated, "These results are impressive, and the message is important. Physicians are not being aggressive enough...If you look at all the factors they (the researchers) treated, about 80% of the U.S. population isn't treated correctly, according to national surveys."

*Diabetologia. 2016 Nov;59(11):2298-307.

Aspirin Fights Cancer

A recent study suggests **aspirin** could slow the growth of some types of cancer.*

The research was designed to determine how inhibition of platelet activation through the use of aspirin might affect the proliferation of colon and pancreatic cancer cells.

Platelets, when activated, cause blood to clot. They can also promote the growth of cancer cells through releasing growth factors and enhancing the response of *oncoproteins*, which regulate the development of tumor cells. Aspirin is an antiplatelet drug, and low doses have been known to reduce the risk of some gastrointestinal cancers by mechanisms still under investigation.

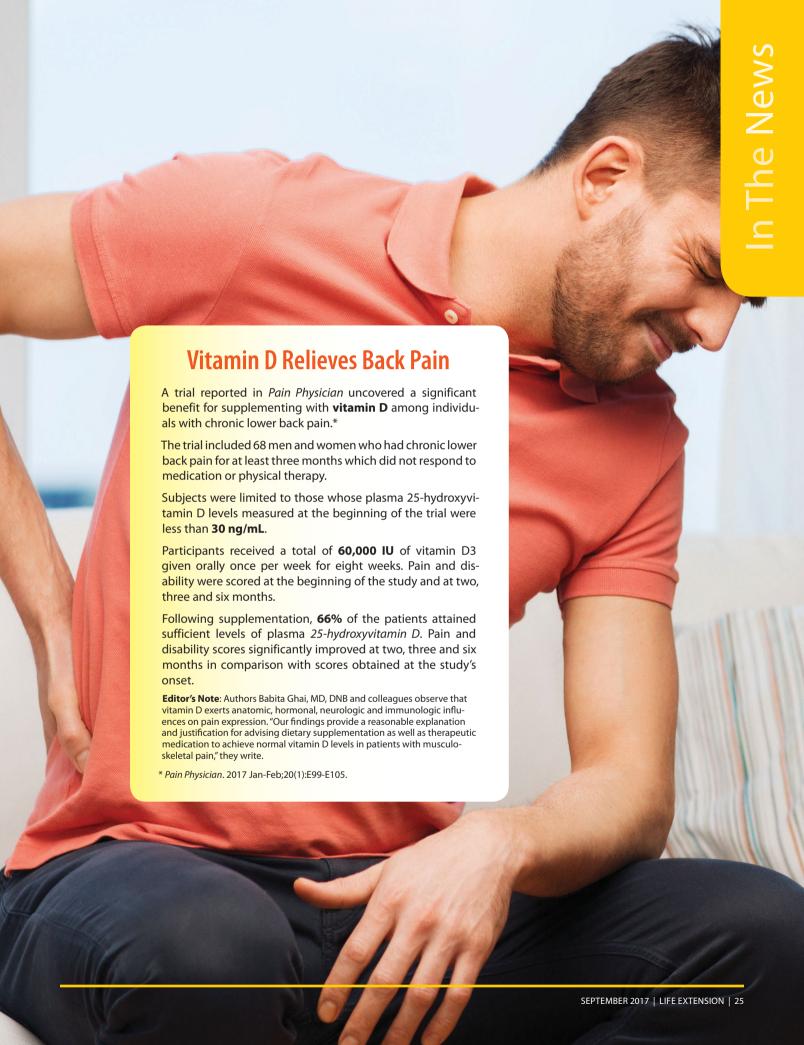
Researchers combined platelets with metastatic (cancer that has spread to other parts of the body) colon cancer cells, nonmetastatic (cancer that has not spread) colon cancer cells, and nonmetastatic pancreatic cancer cells. Aspirin was then added to all three groups.

Results showed that a low aspirin dose stopped platelets from prompting growth and replication of nonmetastatic pancreatic and colon cancer cells. The growth of metastatic pancreatic cancer cells could also be stopped with aspirin, but only at doses too large for humans to ingest. Metastatic colon cancer cells were unaffected at any dose.

These data are corroborated by human studies showing lower risk of many cancers in those taking low-dose aspirin daily.

Editor's Note: The researchers found these results promising. "Our study," they wrote, "reveals important differences and specificities in the mechanism of action of high- and low-dose aspirin in metastatic and nonmetastatic cancer cells with different tumor origins and suggests that the ability of aspirin to prevent platelet-induced c-MYC (an oncoprotein) expression might be selective for a nonmetastatic phenotype."

* Am J Physiol Cell Physiol. 2017 Feb 1;312(2):C176-C189.





"D"fend Your Health

VITAMIN D3

Systemic Support for Youthful Cell Function



Non-GMO Price Price

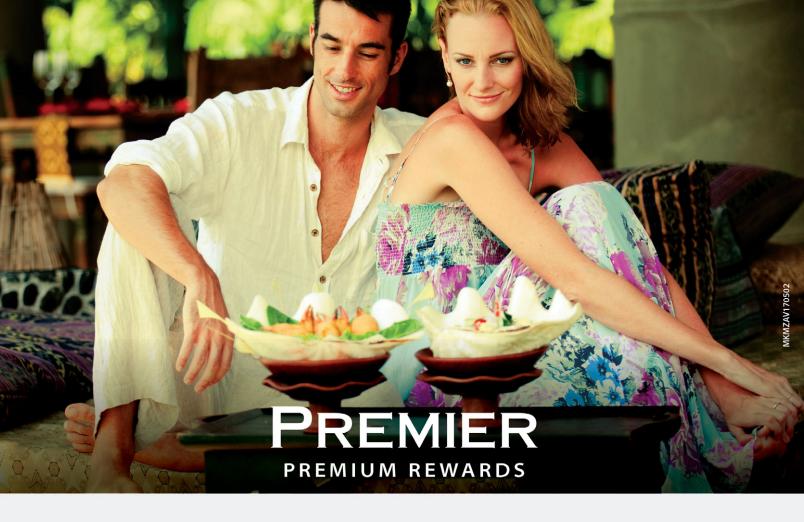
1 bottle \$10 \$7.50
4 bottles \$6.50 each

1 tem # 01713 • 5,000 IU • 60 softgels

For full product description and to order Vitamin D3, 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 II per day unless recommended by your doctor. Vitamin D supplementation is not recommended for individuals with high blood calcium levels.

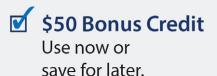








ALL YEAR LONG





Join Premier Today! Only \$49.95 per year

For details, visit www.LifeExtension.com/YourPremier Call 1-866-542-9357 toll-free

Mention code YRX703E

LifeExtension°

Premier service is good for a full 12 months from the date of purchase or renewal, and can only be renewed after 6 months from the date of your last Premier purchase or renewal. Redeem LE Dollars to purchase virtually anything we sell, including products, blood tests, sale items, and even shipping fees! At the rate of 1 LE Dollar equal to \$1 U.S. Dollar at checkout. 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. Enjoy all the rewards of Premier.

Ben Stiller Wants Men to Test for Prostate Cancer

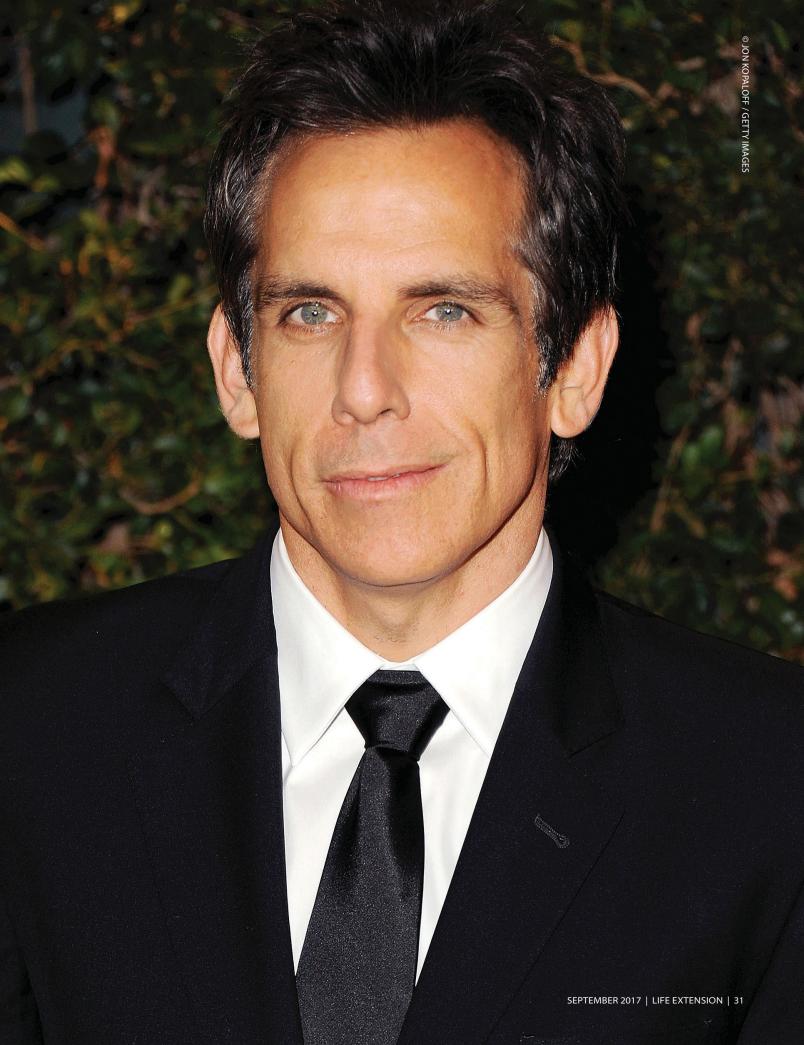
Actor Ben Stiller was as surprised as anyone when he heard these words: "So yeah, it's cancer."

After all, he was only **48** and had no real reason to suspect that he had cancer, especially prostate cancer, which many people think of as an older man's disease.

"I have no history of prostate cancer in my family and I'm not in the high-risk group," he wrote in a public posting detailing his experience. "I had no symptoms."

So how did the star of movies including *There's Something About Mary, Meet the Parents*, and *Zoolander* end up getting diagnosed in the first place? And what does his case have to say about the way we diagnose and treat prostate cancer in the United States?

Stiller's story began two years <u>before</u> the day in June 2014 when he was diagnosed with prostate cancer. This is when his doctor, a "thoughtful internist", gave him a simple and inexpensive **PSA** screening test. This was the first of many PSA tests over the next few years.



A one-time modest elevation of **PSA** blood levels can be explained by several factors that are often correctable. So the best course of action is to have follow-up PSA tests to monitor what direction the PSA is moving in.

As follow-up PSA tests were performed, Ben Stiller's doctor noted a gradual rise in Stiller's PSA over his earlier baseline. These rising levels triggered a referral to a urologist, who did further testing, including a digital rectal exam, an MRI, and finally a biopsy that confirmed the diagnosis.

Three months after his diagnosis, Stiller had undergone treatment—in his case a robotic-assisted laparoscopic radical prostatectomy, or removal of his prostate gland during a minimally invasive surgery—and was cancer free. That could have been the end of it, but after doing his research into prostate cancer screening and diagnosis, Stiller realized he couldn't be silent about his experience. He's been spreading the same message ever since: "Taking the PSA test saved my life."

This might not seem like a controversial statement—after all, it might seem hard to argue against a simple blood test that can identify prostate cancer early enough to treat it before it spreads and without major side effects. But in fact, due to recent chaos in the official recommendations for **PSA blood testing**, tens of thousands of American men are skipping the very test that possibly saved Stiller's life on the advice of their doctors and with potentially devastating consequences.

History of Screening Recommendations

The PSA test is used to measure **prostate-specific** antigen, a protein that is produced by the prostate gland.

PSA levels rise in aging men and can be the first signal of underlying prostate cancer. So the PSA blood

test is used to identify men who may have prostate malignancy and need further evaluation.

This simple blood test was approved by the FDA in 1994, allowing men to begin monitoring their PSA levels and identify possible tumors long before they become dangerous.1

Since PSA testing was introduced, the risk of dving from prostate cancer among men who were regularly screened declined by as much as 42%.2,3

Despite this drop, widespread PSA screening remained controversial in the medical community.

Prostate cancer is typically a slow-growing cancer, and the current biopsy and treatment methods. including the kind of less-invasive surgical removal that Stiller underwent, carry risks such as pain, incontinence and impotence. Some doctors worried that the PSA test, which can detect very slight increases in PSA levels, might be causing men with low-risk cancers to undergo biopsies and possibly unnecessary treatment.

Based on these concerns, in 2012, the US Preventive Service Task Force (USPSTF) issued a stunning update to prostate screening recommendations. Drawing its conclusions from the results of a \$400 million federal study, the USPSTF advised against PSA screening for healthy men, saving that PSA screening has "no net benefit."4-6 The American Cancer Society soon revised its recommendations, steering healthy, average-risk men away from PSA screening until age 50, with revised recommendations for men with a family history of prostate cancer.7

These guidelines caused immediate uproar in the medical community, including rebuttals from Life Extension[®] urging men over age 40 to continue having annual PSA blood tests. By 2016, the USPSTF announced it was reconsidering its prior recommendations against PSA screening.



In 2017, a new draft recommendation was released for public input. This time, the USPSTF slightly backtracked, saving that the risks and benefits of PSA screening are "closely balanced" in men between the ages of 55 and 69 and they should seek their doctor's advice on PSA screening. Men aged 54 and under and those over the age of 70 would still be counseled to avoid PSA screening. These new, slightly softer guidelines were still not finalized as of May 2017, and the agency was soliciting public input.8

In Stiller's case, following even the updated guidelines might have meant disaster—he was still too young to be screened according to the **USPSTF** (United States Preventive Service Task Force).

"If he [Ben Stiller's doctor] had waited, as the American Cancer Society recommends, until I was 50, I would not have known I had a growing tumor until two years after I got treated," he wrote. "If he [Ben Stiller's doctor] had followed the US Preventive Service Task Force guidelines, I would never have gotten tested at all, and not have known I had cancer until it was way too late to treat successfully."

The USPSTF's original recommendations against screening were partly based on the **Prostate**, **Lung**, Colorectal, and Ovarian Cancer Screening Trial. This huge trial assigned 76,685 men aged 55 – 74 years to one of two study arms. The first group (38,340 men) underwent annual PSA testing for 6 years and an annual digital rectal exam for 4 years. The control group (38,345 men) underwent normal care, with occasional "opportunistic screening" but no regular PSA monitoring. At the end of the 13-year follow-up period, researchers announced there was "no evidence of a mortality benefit" for annual PSA screening.9 The USPSTF recommendation against PSA screening soon followed.

Life Extension, which has long supported PSA screening, issued a detailed rebuttal challenging the findings of this study. In fact, the study was deeply flawed thanks to widespread "contamination" of the control arm.

While **Life Extension** was early in identifying the obvious flaws with this study, it wasn't long until astute research groups began to catch up. In early 2016, a group of urologists from the New York Presbyterian Hospital and Weill Cornell Medical College in New York published a letter in the New England Journal of *Medicine* confirming what **Life Extension** suspected.¹⁰

The shocking truth was that more than 80% of the men in the **control** group—which was supposed to only receive "occasional" PSA screening-reported at least one PSA test during the trial. In fact, by some measures, the men in the **control** group received **more** PSA screening than men in the **PSA screening** arm!¹⁰

Their conclusion? "We're going to have to reconsider this issue."11



Further support for this position was published in another large study, this one called the **European** Randomized Study of Screening for Prostate Cancer. This study randomized 182,000 men aged 50 to 74 to a "usual care" control group or a group with PSA screening every two to seven years. Spread across seven research centers in Europe, the group tracked prostate cancer mortality in both study arms. At the median follow-up of nine years, researchers reported that **PSA screening** resulted in a **20**% reduction in prostate cancer mortality!12

A study from the Göteborg center, one of the seven participating centers in this study, found that men aged 50 to 64 years of age who had a PSA screening every other year had a 44% reduced mortality risk from prostate cancer. The center used a PSA cutoff of 2.5 ng/mL to 3.0 ng/mL. Men with these cutoff PSA levels and higher were referred for additional testing, including a digital rectal exam, transrectal ultrasound, and prostate biopsy.13

Although it's too late to help the tens of thousands of men who likely skipped PSA screening, we are grateful the USPSTF is slowly grappling with the welldocumented issues in its original guidelines by issuing the new draft recommendations.14



The issue was further complicated by results from a study published in the New England Journal of Medicine in **2016**. This trial followed 1,643 men for a decade, each with prostate cancer that was first detected by PSA screening, to see which of the most popular treatment techniques was most effective, including "active waiting" and monitoring the disease, surgery to remove the prostate gland, or external radiation beam therapy to treat the cancer. While the prostate-cancer-specific survival rate was high (>98%) in all three groups, researchers found that men in the "active waiting" group were more likely to progress to metastatic disease, and about half of them needed surgery or radiation therapy within the 10-year study period. 15

These results suggest that men benefit from early detection and early treatment of prostate cancer.

Please note that Life Extension does not recommend "watchful or active waiting" in the presence of **high PSA** and/or **low-grade** prostate cancer. We instead advise men to follow an aggressive "active surveillance" program that involves an anticancer diet along with specific drugs and nutrients that may enable early-stage disease to be contained.

For information about Life Extension's "active surveillance" guidelines, refer to the article titled How to Reverse Markers of Prostate Cancer in the **June 2016** issue of this magazine. (Available free of charge at LifeExtension.com.)

Rise in Metastatic Cancer Rates

While various agencies continue to issue contradictory and confusing advice, men across the country have paid the price. In late **2016**, a research group from Northwestern Medicine released a stunning and

tragic finding: diagnoses of metastatic prostate cancer, the worst type, climbed an unbelievable 72% between 2004 and 2013.16

To reach these findings, the group studied a database of more than three-quarters of a million men in the National Cancer Data Base. What they found should alarm any man who skips his PSA screening.

"The fact that men in 2013 who presented with metastatic disease had much higher PSAs than similar men in 2004 hints that more aggressive disease is on the rise,"17 said study author Dr. Edward Schaeffer, chair of urology at Northwestern University Feinberg School of Medicine and Northwestern Medicine.

"One hypothesis is the disease has become more aggressive, regardless of the change in screening," said Dr. Schaeffer. "The other idea is since screening guidelines have become more lax, when men do get diagnosed, it's at a more advanced stage of disease. Probably both are true. We don't know for sure but this is the focus of our current work."17

This makes treatment more difficult, and it's exactly the situation Ben Stiller would have faced if his forward-thinking doctor hadn't established a PSA baseline early on and tracked it, allowing him to discover Stiller's troubling increase in PSA levels over time and recommend the movie star for further evaluation and surgery.

It's important to note that the increase in metastatic, aggressive prostate cancer almost perfectly aligns with the trend away from PSA screening that culminated with the USPSTF 2012 recommendation against any PSA screening.

Stiller's Happy Ending

The main concern with PSA screening is the potential for overdiagnosis and unnecessary treatment. These are real concerns—PSA screening frequently returns "false positives," which are stressful for the patients involved and result in unnecessary biopsies and additional tests.18

This is why **Life Extension** recommends regular, inexpensive PSA screening to establish a baseline and follow PSA numbers over time. If your PSA level rises above **1.0 ng/mL**, there are natural and safe measures you can take to reduce it. Further evaluation may be necessary if your PSA continues to rise over time.

In fact, this is exactly the course Stiller followed, and today he's alive and grateful for it.

"The bottom line for me: I was lucky enough to have a doctor who gave me what they call a 'baseline'

© TONY BARSON / GETTY IMAGE:

PSA test when I was about 46," he wrote in Medium, a popular blogging platform. "My doctor watched my PSA tests rise for over a year and a half, testing me every six months...I think men over the age of 40 should have the opportunity to discuss the test with their doctor and learn about it, so they can have the chance to be screened."19

More recently, two years after his diagnosis and treatment, Stiller went public with his experience with an interview with Matt Lauer on the *Today* show, alongside Dr. Schaeffer. While reporting that he wasn't experiencing any of the major complications of prostate surgery, Stiller gave a simple reason for going public. He wanted to educate as many men as possible about their options when it came to PSA screening.

"It's a whole new world," Stiller said. "You need to educate yourself."

We at Life Extension commend Bernard M. Kruger. **M.D.** for having the foresight to test Ben Stiller's **PSA** blood levels despite conventional "authorities" advising against PSA screening.

The good news for all men is that newer imaging techniques, as described in the June 2016 issue of Life Extension magazine, are reducing biopsy side effect risk and enabling men to eradicate prostate cancer without major surgery and radiation.

Prostate Cancer Survivors Due to Early Detection Name **Year Successfully Treated** Robert De Niro 2003 at age 60 John Kerry 2003 at age 60 Rudy Giuliani 2000 at age 56 **Robert Goulet** 1993 at age 60 Colin Powell 2003 at age 66 Michael Milken 1993 at age 46

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: https://www.cancer.gov/types/prostate/psa-fact-sheet. Accessed May 2, 2017.
- Arnsrud Godtman R, Holmberg E, Lilja H, et al. Opportunistic testing versus organized prostate-specific antigen screening: outcome after 18 years in the Goteborg randomized population-based prostate cancer screening trial. Eur Urol. 2015;68(3):354-60.



- Roobol MJ, Kranse R, Bangma CH, et al. Screening for prostate cancer: results of the Rotterdam section of the European randomized study of screening for prostate cancer. Eur Urol. 2013;64(4):530-9.
- 4. Available at: https://www.uspreventiveservicestaskforce.org/Page/ Document/RecommendationStatementFinal/prostate-cancerscreening. Accessed May 2, 2017.
- Moyer VA, on behalf of the USPSTF. Screening for prostate cancer: U.s. preventive services task force recommendation statement. Annals of Internal Medicine. 2012;157(2):120-34.
- 6. Available at: https://www.uspreventiveservicestaskforce.org/Page/ Name/grade-definitions. Accessed May 9, 2017.
- 7. Available at: https://www.cancer.org/cancer/prostate-cancer/earlydetection/acs-recommendations.html. Accessed May 2, 2017.
- Available at: https://www.uspreventiveservicestaskforce.org/Page/ Document/RecommendationStatementDraft/prostate-cancerscreening1. Accessed May 2, 2017.
- 9. Andriole GL, Crawford ED, Grubb RL, 3rd, et al. Prostate cancer screening in the randomized Prostate, Lung, Colorectal, and Ovarian Cancer Screening Trial: mortality results after 13 years of follow-up. J Natl Cancer Inst. 2012;104(2):125-32.
- 10. Shoag JE, Mittal S, Hu JC. Reevaluating PSA Testing Rates in the PLCO Trial. N Engl J Med. 2016;374(18):1795-6.
- 11. Available at: https://news.weill.cornell.edu/news/2016/05/new-england-journal-of-medicine-letter-to-the-editor-calls-prostate-cancerscreening-guidelines-into. Accessed May 2, 2017.
- 12. Schroder FH, Hugosson J, Roobol MJ, et al. Screening and prostate-cancer mortality in a randomized European study. N Engl J Med. 2009;360(13):1320-8.
- 13. Hugosson J, Carlsson S, Aus G, et al. Mortality results from the Goteborg randomised population-based prostate-cancer screening trial. Lancet Oncol. 2010;11(8):725-32.
- 14. Available at: https://www.uspreventiveservicestaskforce.org/Page/ Document/final-research-plan/prostate-cancer-screening1. Accessed May 2, 2017.
- 15. Hamdy FC, Donovan JL, Lane JA, et al. 10-Year Outcomes after Monitoring, Surgery, or Radiotherapy for Localized Prostate Cancer. N Engl J Med. 2016;375(15):1415-24.
- 16. Weiner AB, Matulewicz RS, Eggener SE, et al. Increasing incidence of metastatic prostate cancer in the United States (2004-2013). Prostate Cancer Prostatic Dis. 2016;19(4):395-7.
- 17. Available at: https://news.northwestern.edu/stories/2016/07/metastatic-prostate-cancer-cases-skyrocket/. Accessed May 2, 2017.
- 18. Lin K, Lipsitz R, Miller T, et al. Benefits and harms of prostatespecific antigen screening for prostate cancer: an evidence update for the U.S. Preventive Services Task Force. Ann Intern Med. 2008;149(3):192-9.
- 19. Available at: https://medium.com/cancer-moonshot/the-prostatecancer-test-that-saved-my-life-613feb3f7c00. Accessed May 2, 2017.



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.

ArthroMax® Advanced with UC-II® & AprèsFlex® is a multinutrient formula that supports healthy joint function.

<u>Two</u> clinically validated ingredients in this formula: AprèsFlex® and UC-II® maintain cartilage and ease inflammation and thus may promote comfortable joint structure and function.

For full product description and to order ArthroMax® Advanced with UC-II®

and AprèsFlex® call 1-800-544-4440 or visit www.LifeExtension.com

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.



Order by **August 31, 2017**, and receive a **44**% discount off the regular retail price!

TAKE ADVANTAGE OF THIS EXCELLENT OFFER TODAY.

To order a PSA blood test (Item Code LC010322) for only \$23.25,

call Life Extension® at 1-800-208-3444 or order online at LifeExtension.com/PSA





Homocysteine Resist

Item #02121 • 60 vegetarian capsules

Non-GMO	Retail Price	Your Price
1 bottle	\$26	\$19.50
4 bottles		\$17.50 each

Homocysteine Resist supports healthy levels of homocysteine, an unfavorable amino acid that can increase with normal aging.

Just one daily capsule of **Homocysteine Resist** provides:

5-MTHF (activated folate)	5,000 mcg
Methylcobalamin (activated vitamin B12)	1,000 mcg
Pyridoxal-5-phosphtate (activated vitamin B6)	100 mg
Riboflavin (vitamin B2)	25 mg

For full product description and to order **Homocysteine Resist**, call **1-800-544-4440** or visit **www.LifeExtension.com**



Tap the <u>Power</u> of N-Acetyl-L-Cysteine To Boost Glutathione Levels



N-acetyl-L-cysteine supports healthy levels of *glutathione*, a molecule utilized by all cells for protection against free radical damage and attacks from foreign compounds.

N-Acetyl-L-Cysteine

Item #01534 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$14	\$10.50
4 bottles	•••••	\$9.25 each



Non-GMO

Caution: Those who supplement with NAC should drink six to eight glasses of water daily in order to prevent cysteine renal stones. Cysteine renal stones are rare but do occur.

For full product description and to order **N-Acetyl-L-Cysteine**, call 1-800-544-4440 or visit **www.LifeExtension.com**



An Innovative Approach to Stopping COLDS and FLU

The Centers for Disease Control and Prevention estimate that 56,000 people a year die from flu-related illnesses.¹ During the 2015-2016 flu season 310,000 people were hospitalized due to flu-related illnesses.²

Colds and the flu are often treated with a variety of medications designed to *reduce* symptoms but that have no ability to *activate* the body's own immune response to fend off invading bacteria or viruses.

Scientists wanted to find a way to <u>stop</u> the common cold and flu before they take hold.

Our bodies have a built-in security system called secretory **IgA**, which is present in mucosal membranes that line the nose and upper respiratory tract. IgA can prevent cold and flu viruses from entering.

With age, our ability to generate IgA secretions declines and this first line of defense is weakened.

Researchers started their investigations knowing that a critical aspect of the immune system is the *microbiome* in the gut, where a complex microenvironment of beneficial bacteria exist and interact.

Scientists discovered that a targeted **probiotic cocktail** of bacteria can boost the body's immune defense system, in particular mucosal secretory **IgA**.

How IgA Protects Against Cold and Flu

Infections of the nose and upper respiratory tract pose a risk to older adults, who have a reduced immune response.3

This weakening of the immune system arises in part from reduced production of secretory antibodies that protect the nasal mucosa and respiratory tract mucosal surfaces from viral infection.

The decline of protective IgA helps explain why aging adults can be susceptible to infection by cold and flu viruses, as well as the Streptococcus pneumoniae bacterium, a cause of bacterial pneumonia in aging adults.4

To counteract this problem, researchers tested a unique **oral probiotic** blend designed to reduce the risk of respiratory infections by enhancing secretory immunity.

The term "secretory immunity" refers to production of specialized antibodies like **IgA** in the mucous membranes lining the nose, and portions of the windpipe and lungs.5

The importance of IgA antibodies is that they target both viral and bacterial invaders in the upper respiratory tract, deactivating them and presenting them for destruction by the immune system.⁶ This prevents both cold and flu viruses from gaining a foothold in the body. These pathogens are stopped before wreaking havoc in the body's respiratory tract.

Blocking viral attachment to mucous membranes. in turn, prevents viruses from injecting their genetic material into human cells, and hence from replicating to produce more viruses.6

IgA is the acronym for immunoglobulin A. Increasing **IgA secretion** and breaking a viral replication cycle can prevent development of colds, influenza, and other respiratory infections.

Probiotics Slash Respiratory Infections

Researchers have identified several unique strains of targeted probiotics that have potent preventive effects on human respiratory infections. Their weapon against microbes, especially viruses that target the respiratory tract, appears to be the **stimulation of IgA**.

To see how this unique probiotic blend works to prevent infection by cold and flu, let's look at a human clinical study. The trial was performed over the course of 90 days during cold and flu season with 250 subjects.⁷

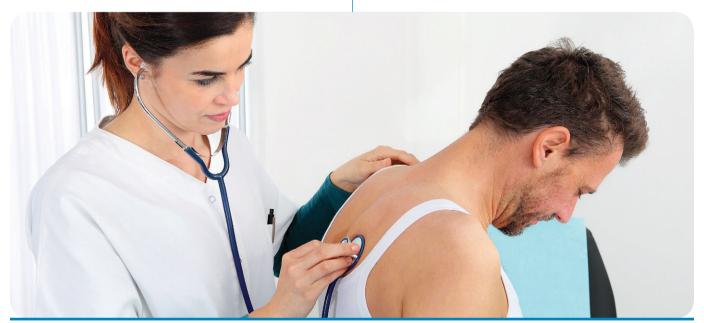
The subjects were randomly assigned to receive either a placebo or a mixture of five unique probiotic strains specifically formulated to stimulate protective IgA:

- L. plantarum (LP 01-LMG P-21021)
- *L. plantarum* (LP 02-LMG P-21020)
- L. rhamnosus (LR 04-DSM 16605)
- L. rhamnosus (LR 05-DSM 19739)
- B. lactis (BS 01-LMG P-21384)

During the course of the three-month study, subjects were asked to report all daily respiratory symptoms (runny nose, cough, fever, bronchitis, or pneumonia), along with the length and severity of symptoms.

All diseases accompanied by fever were classified as "flu-like syndromes," while a separate category of "influenza-like illnesses" was also used. Other categories were "bronchitis-like" diseases, upper respiratory tract infections, common cold, and cough without other symptoms.

The study showed reduction in symptoms and reduction in the duration of symptoms.





- 16 episodes of "influenza-like illnesses" in the placebo group compared to 3 such episodes in the **probiotic cocktail** group (a significant 81% reduction).
- 31 episodes of colds among placebo recipients compared to 20 reported cases in the probiotic group. This 35% reduction did not quite achieve statistical significance.
- Cold duration fell from 6 days in placebo recipients to **4.7** days in the **probiotic** supplemented patients, a 22% reduction.
- Cough duration fell in the patients given the **probiotic** cocktail from 7.3 to 4.5 days, a **39**% reduction.
- Total acute upper respiratory infections fell from **6.1** to **4.6** days in the **probiotic** group, a 25% reduction.

A similar study showed a **48**% reduction in flu episodes on subjects using the **probiotic cocktail**. The number of days with flu symptoms decreased significantly by 55%.8

- IgA antibodies, secreted from mucous membranes in the mouth, nose, and lungs, bind to and block respiratory viruses from invading human cells and producing symptoms of colds and flu. But as their production fades, so does our immunity to these microbes.
- Recent studies have revealed the surprising fact that selected strains of oral probiotic bacteria are capable of stimulating the intestinal immune system, resulting in body-wide increases in secretory immunity.
- In the respiratory tree, this increase in IgA production leads to enhanced protection against cold and flu viruses.
- Studies have now demonstrated convincingly that supplementing with these strains of probiotic bacteria results in significant reductions in the incidence and duration of year-round respiratory infections.
- IgA is the acronym for immunoglobulin A.

Adding Bacillus Subtilis CU1 to the Probiotic Cocktail

In another study, scientists identified a sixth probiotic that also provided immune-stimulating features among aging adults at risk for respiratory infections.9

This bacteria. Bacillus subtilis CU1. creates a natural protective shield that resists the acid in the stomach. promoting the probiotics' survival into the digestive tract.¹⁰ Previous studies have shown that this strain of probiotic can stimulate IgA in humans, a mechanism of great interest in preventing respiratory infections.9,11

A human clinical trial was conducted among healthy older adults (ages 60-74) who were randomly assigned to receive this probiotic or a placebo. Subjects took one capsule daily, containing two billion microorganisms per capsule.

Supplemented subjects experienced a significant **45%** drop in the frequency of respiratory infections. Notably, a concomitant significant 45% increase in concentrations of **IgA** was demonstrated in their saliva, strongly suggesting that increased IgA was at least in part responsible for the observed impact. No significant side effects were noted in either group.

Unleashing the Nuclear Bomb to Stop Colds and Flu

The importance of taking aggressive actions upon the first signs and symptoms of viral respiratory infections is critical. This protocol or "nuclear bomb" should be initiated within 24-48 hours of the manifestation of serious cold and flu symptoms.

- 1. **800 mg** of **cimetidine** (and higher). This drug is sold over the counter in pharmacies to combat heartburn, but its beneficial side effect is to boost immune function by reducing T-suppressor cells, thereby keeping the immune system active.¹⁵ Cimetidine can interact with prescription medications, so consult with a pharmacist and your physician before using. For most people, cimetidine provides immune system stimulation that is particularly effective against certain viruses.
- 2. 9,000 mg of high-allicin garlic once or twice daily. This potent form of garlic will cause painful stomach-esophageal burning if you don't eat food right afterward. Ingesting 9,000 mg of this kind of garlic will cause you to reek of a strong sulfur odor, but saturating the body with it is the objective. Garlic has shown direct virus-killing effects in a number of published studies.^{16,17}
- 3. 200 mg of DHEA early in the day. This is a high dose, 18 but DHEA has shown some unique benefits in boosting one's ability to mount a stronger immune response and also protecting against dangerous inflammatory cytokine responses that sometimes occur in response to viral infections.
- 4. 1,200 mg a day of lactoferrin. This natural constituent of mother's milk boosts natural killer-cell activity and can kill certain viruses.19

- 5. Two **18.5-mg zinc acetate lozenges** every two waking hours. Please be aware that this is a very high dose of zinc and is considered toxic if taken over the long term.^{20,21} You should only do this for a few days. Zinc has shown a direct effect of inhibiting cold viruses from latching onto your cells.¹⁹
- 6. 10-50 mg of melatonin at bedtime (ordinarily, melatonin is taken at levels of just 1-3 mg per evening). Melatonin induces a powerful immune response and this high dose can facilitate the deep sleep one often needs to fend off an infection. This dose of melatonin will make you extremely tired, so please only take this before bedtime and do not operate any machinery or vehicles after ingestion.²²
- 7. **3,600 mg** a day of **aged garlic extract**. There are unique immune-boosting compounds in aged garlic that work differently than those found in high-allicin garlic.23
- 8. As discussed on the sidebar on page 45, the prescription drug **Tamiflu**® in the dose of **75 mg** twice a day should be started within 24 hours of flu symptoms manifesting to block entry of certain viruses into cells where they multiply.

Do not delay in implementing the above regimen. Once a flu virus infects too many cells, it replicates out of control and strategies like zinc lozenges will not be effective. Treatment should be initiated as soon as possible after symptoms manifest!

The Importance of Dendritic Cells

How can ingestion of a **probiotic** lead to increased secretion of **IgA** in the nose and throat?

IgA production can be stimulated or reduced throughout the body, depending on the environment sensed by specialized cells called *dendritic cells*. Dendritic cells can detect molecular patterns on the surfaces of the billions of bacteria and viruses we swallow every day, whether they enter our bodies through the mouth or the nose.7

Once dendritic cells have encountered these microbial identifiers, they "teach" other immune system cells about the nature of the threat, prompting them to pump out **IgA**, among other defensive molecules.⁷

IgA is secreted throughout the body, including mucous membranes of the mouth and nose.¹² Increased IgA in those areas results in increased protection against invaders attempting to enter the body through those membranes.7

Studies have demonstrated that orally-ingested probiotics stimulate IgA in the mucous membranes of the bronchi (larger air tubes in the lungs).^{7,12} Similarly, probiotics have been shown to reduce the incidence and severity of respiratory infections in children.^{7,13}



When Influenza Turns Deadly: Life-threatening Influenza **Requiring Hospitalization**

Those over the age of 65 as well as aging individuals with chronic diseases that may weaken the immune system, cardiovascular system, and/or respiratory tract (e.g. diabetes, cardiovascular, and chronic lung disease) are at higher risk of developing potentially life-threatening infections including the dreaded influenza pneumonia that is linked to a high mortality rate.

In contrast to younger and generally healthy patients, these high-risk patients may not necessarily manifest a high fever initially (a body temperature in excess of 102.5 degrees Fahrenheit) since impaired thermoregulation is observed with aging.

Headache, dramatic fatigue, muscle aches, a nonproductive cough (initially), and nasal congestion are important signs and symptoms of influenza infection in older patients. Additional signs and symptoms in older patients may include cognitive dysfunction and confusion, difficulty walking, and falls.

Worrisome signs and symptoms for patients with influenza infection suggesting pneumonia include:

- Respiratory rate above 25 breaths per minute, reflecting difficulty with oxygenation
- Hypotension (blood pressure below 90/60 mm Hg),
- Bloody sputum

Labs tests that also suggest severe disease include:

- Elevated lactate dehydrogenase (LDH)
- Elevated creatine phosphokinase (CPK)
- Hypoxemia (the inability to oxygenate the blood) increases rapidly to the point of respiratory failure in many of these patients requiring mechanical ventilation, often after only 24 to 48 hours.

The most important pharmacologic intervention to reduce the risk of death in these patients, in addition to cardiovascular and respiratory supportive measures, is rapid identification of viral strain, and rapid antiviral treatment with neuraminidase inhibitors, e.g., oseltamivir (Tamiflu®) and zanamivir (Relenza®), initiated as soon as possible, ideally within 24 hours.

Thus, orally ingested probiotics appear capable of sealing the age-induced gap in *secretory immunity* that puts so many older adults at grave risk for respiratory infections every year.

The discovery of the unique **IgA**-stimulating properties of oral **probiotics** opens a new world in modulation and strengthening of our aging immune systems, and provides a new weapon in our battle against respiratory illness.

Summary

A cold or flu can pose serious risks for aging adults. Such infections can lead to potentially fatal bacterial infections, particularly pneumonia, which kills more than 50,000 Americans annually.14

One main source of the age-related increase of viral respiratory infections is the loss of *secretory immunity*, which is controlled by a class of antibodies known as IgA.

As aging adults lose IgA protection, they lose their ability to defend against viruses attacking the mucous membranes in the nose, lungs, and bronchi of the respiratory tract.

Specific strains of probiotic bacteria, ingested orally, have the ability to *stimulate* IgA production in the respiratory mucous membranes, thereby preventing attachment by, and infection with, common viruses.

Human studies demonstrate that supplementation with the proper blend of **probiotic** strains reduces the incidence of colds and flu-like illnesses, an effect largely attributable to increases in levels of IgA.



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. Available at: https://www.cdc.gov/flu/about/disease/us_flu-related_ deaths.htm. Accessed June 12, 2007.
- Available at: https://www.cdc.gov/flu/about/qa/hospital.htm. Accessed June 12, 2017.
- Fujihashi K, Kiyono H. Mucosal immunosenescence: new developments and vaccines to control infectious diseases. Trends Immunol. 2009:30(7):334-43
- 4. Fujihashi K, Sato S, Kiyono H. Mucosal adjuvants for vaccines to control upper respiratory infections in the elderly. Exp Gerontol. 2014;54:21-6.
- 5. Fagarasan S, Honjo T. Regulation of IgA synthesis at mucosal surfaces. Curr Opin Immunol. 2004;16(3):277-83.
- Snoeck V, Peters IR, Cox E. The IgA system: a comparison of structure and function in different species. Vet Res. 2006;37(3):455-67.
- Pregliasco F, Anselmi G, Fonte L, et al. A new chance of preventing winter diseases by the administration of synbiotic formulations. JClin Gastroenterol. 2008;42 Suppl 3 Pt 2:S224-33.
- 8. Belcaro G. Cesarone MR. Cornelli U. et al. Prevention of flu episodes with colostrum and Bifivir compared with vaccination: an epidemiological, registry study. Panminerva Med. 2010;52(4): 269-75.
- Lefevre M, Racedo SM, Ripert G, et al. Probiotic strain Bacillus subtilis CU1 stimulates immune system of elderly during common infectious disease period: a randomized, double-blind placebocontrolled study. Immun Ageing. 2015;12(1):24.
- 10. Cutting SM. Bacillus probiotics. Food Microbiol. 2011;28(2):214-20.
- 11. Corthesy B. Multi-faceted functions of secretory IgA at mucosal surfaces. Front Immunol. 2013;4:185.
- 12. Perdigon G, Alvarez S, Medina M, et al. Influence of the oral administration of lactic acid bacteria on iga producing cells associated to bronchus. Int J Immunopathol Pharmacol. 1999;12(2): 97-102
- 13. Hatakka K, Savilahti E, Ponka A, et al. Effect of long term consumption of probiotic milk on infections in children attending day care centres: double blind, randomised trial. BMJ. 2001:322(7298):1327.
- 14. Available at: https://www.cdc.gov/nchs/fastats/pneumonia.htm. Accessed May 25, 2017.
- 15. Kumar A. Cimetidine: an immunomodulator. Dicp. 1990;24(3):
- 16. Harris JC, Cottrell SL, Plummer S, et al. Antimicrobial properties of Allium sativum (garlic). Appl Microbiol Biotechnol. 2001:57(3):282-6.
- 17. Guo NL, Lu DP, Woods GL, et al. Demonstration of the anti-viral activity of garlic extract against human cytomegalovirus in vitro. Chin Med J (Engl). 1993;106(2):93-6.
- 18. van Vollenhoven RF, Morabito LM, Engleman EG, et al. Treatment of systemic lupus erythematosus with dehydroepiandrosterone: 50 patients treated up to 12 months. J Rheumatol. 1998;25(2):285-9.
- 19. Roxas M, Jurenka J. Colds and influenza: a review of diagnosis and conventional, botanical, and nutritional considerations. Altern Med Rev. 2007;12(1):25-48.
- 20. Solomons NW. Mild human zinc deficiency produces an imbalance between cell-mediated and humoral immunity. Nutr Rev. 1998;56(1 Pt 1):27-8
- 21. Lewis MR, Kokan L. Zinc gluconate: acute ingestion. J Toxicol Clin Toxicol. 1998;36(1-2):99-101.
- 22. Melatonin. Monograph. Altern Med Rev. 2005;10(4):326-36.
- 23. Kyo E, Uda N, Kasuga S, et al. Immunomodulatory effects of aged garlic extract. J Nutr. 2001;131(3s):1075s-9s.



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.

AGE LESS WITH GEROPROTECTORS AND ARTIFICIAL INTELLIGENCE

Longevity scientists have designed

Geroprotect™ Ageless Cell™

to combat aging at the cellular level.



Just <u>one</u> **Geroprotect™ Ageless Cell™** softgel daily provides:

Myricetin 50 mg
N-Acetyl-L-Cysteine 450 mg
Epigallocatechin Gallate [EGCG] 100 mg
Gamma tocotrienol 25 mg

Geroprotect™ Ageless Cell™ Item #02119 • 30 softgels

Retail Your Price

1 bottle \$40 \$30

4 bottles \$27 each

For full product description and to order **Geroprotect™ Ageless Cell™**, call **1-800-544-4440** or visit **www.LifeExtension.com**

Consult your healthcare provider before use if you have a bleeding disorder, are taking anticoagulant or antiplatelet medications or beta-blockers such as Nadolol. Keep out of reach of children. Do not exceed recommended dose.

FlorAssist® Immune Health

Probiotic Blend for Optimal Immune Support



ITEM # 02124 · 30 vegetarian capsules · Retail price \$26 · Your price \$19.50 · 4 bottles \$18 each

For full product description and to order FlorAssist® Immune Health, 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.



Mega Green Tea Extract provides powerful antioxidant effects throughout the body.

Each **725 mg** capsule of **Mega Green Tea Extract** is <u>standardized</u> to **98% polyphenols** that provide **326 mg** of **EGCG** to:

- Protect against DNA damage and oxidative stress¹
- Support healthy blood sugar levels²
- Enhance heart health³
- Boost brain function⁴
- Support strong bones⁵
- Maintain healthy cholesterol levels already within normal range⁶

Each cost-effective bottle lasts over three months!

References

- 1. Mutagenesis. 2015;30(1):129-37.
- 2. Curr Opin Clin Nutr Metab Care. 2013:16(6):688-97.
- 3. Circ J. 2010;74(3):578-88.
- 4. Nutrition. 2014;30(3):337-42.
- 5. BMC Musculoskelet Disord. 2009:10:110.
- 6. J Transl Med. 2015;13:79.

Note: **EGCG** is the acronym for **epigallocatechin gallate**, which is the polyphenol in green tea that has demonstrated the most robust health benefits.

Mega Green Tea Extract Decaffeinated

Item #00954 • 100 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$30	\$22.50
4 bottles		\$18 each
Non-GMO		



LifeExtension

Mega Green Tea

Extract

Per-Day Concentrated Polyphenol Extract

Mega Green Tea Extract Lightly Caffeinated

Item #00953 • 100 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$30	\$22.50
4 bottles		\$18 each
Non-GMO		



For full product description and to order Mega Green Tea Extract, call 1-800-544-4440 or visit www.LifeExtension.com

SUPPORT YOUR AGING IMMUNE **SYSTEM**

Life Extension® researchers have developed an innovative formula with three natural extracts—Cistanche, Pu-erh Tea, and Reishi Mushroom that supports more youthful immune function.

Cistanche

- Supports longer lifespan in animals.¹
- Optimizes the ratio of CD4 to CD8 cells, indicative of a more youthful immune system.¹

Pu-erh tea

 Supports decreases interleukin-6 (IL-6) while boosting natural killer cells and naïve T cells.2

Reishi

 Helps reduce biomarkers of immune senescence.3

For full product description and to order Immune Senescence Protection Formula™. call 1-800-544-4440 or visit www.LifeExtension.com

- 1. Anti-Aging Med. 2011;8(2):7-14.
- 2. Food Chem. 2012 Dec 15;135(4):2222-8.
- 3. Am J Chin Med. 2011;39(1):15-27.







Relieve Urinary-Tract Symptoms by Boosting Prostate Health

More than **50%** of men in their 60s and as many as **90%** of men in their 80s suffer **lower urinary-tract** symptoms related to **prostate** enlargement.¹

Common problems include urinary frequency, urgency, and weak stream. Over time, these urinary challenges can cause bladder infection, kidney stones, kidney damage, and erectile dysfunction.²⁻⁴

Frequent nighttime urination also results in chronic sleep deprivation—increasing the risk of heart attack,^{5,6} cancer,^{7,8} depression,⁹ inflammatory syndromes,^{10,11} and death.¹²

In lieu of drugs, **European** physicians have for decades prescribed natural **plant extracts** to alleviate urinary discomforts associated with benign prostate enlargement (BPH). Efficacy of these **plant extracts** has been demonstrated in a number of published medical studies.

Scientists have found that combining different **plant extracts** works better to alleviate prostate discomfort.

Conditions Causing Enlarged Prostate

A young healthy prostate gland weighs less than an ounce,13 but enlarges considerably as most men age.

Because of its location, even a small amount of prostate swelling can block the urethra, resulting in decreased urine flow and the potential for incomplete bladder emptying.

The majority of men who suffer from nonmalignant prostate enlargement have either **benign prostatic** hyperplasia or prostatitis. 14,15

Benign prostatic hyperplasia, or BPH, is a common cause of prostate enlargement that occurs when prostate cells increase in number. Men have a 50% chance of developing BPH over the course of their lives.^{2,16} It leads to progressive urinary symptoms^{2,17,18} and erectile dysfunction. 19-21 In severe cases it can cause kidney and bladder infections,²² bladder stones,²³ and other serious long-term problems.

Prostatitis is characterized by prostate inflammation or infection, and treating it is medically challenging.²⁴⁻²⁷ This is partly because prostatitis is an umbrella term for many conditions, including acute and chronic bacterial prostatitis, chronic nonbacterial prostatitis, inflammatory and noninflammatory prostatodynia (prostate pain), and asymptomatic inflammatory prostatitis.²⁸⁻³⁰

Scientists have found that men with persistent urinary problems can benefit by supplementing with prostate-specific plant extracts.

Saw Palmetto

The **saw palmetto berry** (*Serenoa repens*) was long ago shown to improve symptoms associated with an enlarged prostate.

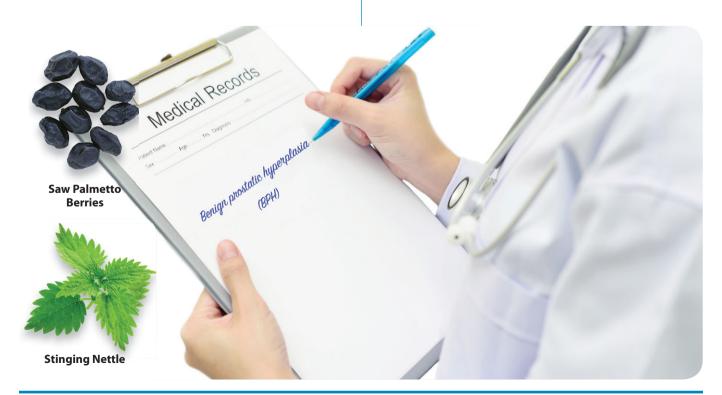
Some studies suggest it helps relieve BPH symptoms as well as finasteride and tamsulosin, 31,32 minus the sexual dysfunction associated with these drugs.³¹

One study found that 320 mg of saw palmetto extract daily improved the International Prostate Symptom Score of elderly men by 52% and their sexual dysfunction scores by 40%.32 Another review demonstrated saw palmetto's efficacy, especially for men with mild-to-moderate BPH symptoms.33

Two large meta-analyses showed that saw palmetto extract improved the **International Prostate Symptom** Score, reduced nighttime urination frequency, and improved peak urine-flow rates.^{34,35}

An underlying reason for these effects is that saw **palmetto** is rich in bioactive compounds—including beta-sitosterol—that support the aging prostate. One mechanism is to impede the adverse effect of dihydrotestosterone (DHT) on prostate cells. DHT is a hormone that increases prostate growth.³⁶ Blocking DHT receptors on prostate cells reduce DHT's potential negative impact.³⁷

Not all studies on **saw palmetto** by itself demonstrate clinical benefits.³⁸ For this reason, plant-based prostate formulas today include additional botanical extracts, and many are prescribed in Europe as "drugs" to alleviate urinary symptoms.



Nettle Root

Urtica dioica, or **stinging nettle root**, has been shown to shrink the prostate and relieve BPH symptoms 39-41

One study showed that nettle root extract improved lower urinary tract symptoms significantly better than placebo, with marked **improvements** in the International Prostate Symptom Score, increases in **peak urinary flow rates**, and reductions in **urine** volume remaining in the bladder.⁴¹

Other compelling research found that the combination of nettle root and saw palmetto extracts results in improvements that are similar to those of prescription BPH medications, but with substantially fewer adverse events.42-44

Scientists demonstrated that stinging nettle combined with saw palmetto reduced nighttime urination by **one episode** nightly, a significant difference.⁴³ And, in elderly BPH patients, this combination reduced the International Prostate Symptom Score by 53%. improved urinary flow by 19%, and reduced residual urine volume 44% compared to placebo. 45

While producing improvements similar to BPH drugs, this combination (nettle root and saw palmetto) results in far fewer adverse events. 42,44

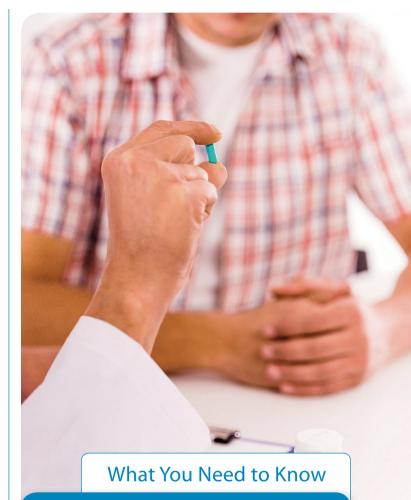
Pygeum Africanum

Pygeum africanum is a plum tree in tropical Africa, and an extract derived from this tree has been widely used in Central and Eastern Europe for decades. Numerous human studies have demonstrated the clinical efficacy of pygeum extract in the management of mild to moderate BPH.46,47

Pygeum extract has been shown to improve International Prostate Symptom Scores by 38%-46%,⁴⁷ reduce the frequency of nighttime urination by 32%,46 and raise peak urinary flow rates by 16%-19%, 47—all at typical doses of **100 mg** per day.

In two different studies, scientists demonstrated that pygeum extract improves quality of life—a critical factor in this condition—by about 30%.46,47

In a meta-analysis of 18 randomized clinical trials that involved a total of 1,562 men, researchers found that **pygeum extract** reduced nocturnal urination by 19% and increased urine flow by 23%. Pygeum use also resulted in a critical reduction in the volume of urine remaining in the bladder after urination, which decreases the risk of urinary tract infections. The extract was also found to more than **double** the odds of a man reporting an overall improvement in urinary tract symptoms compared to men using a placebo.48



Naturally Treat Urinary-Tract Symptoms

- **BPH causes lower urinary tract symptoms** in aging men and can result in serious complications, including bladder infections, kidney damage, and erectile dysfunction.
- It involves frequent nighttime urination that results in chronic sleep deprivation, in turn increasing the risk of heart attack, cancer, depression, inflammatory syndromes, and death.
- Several natural extracts have been found to safely shrink an enlarged prostate—as well as improve many of the unpleasant symptoms associated with BPH.



Rye Flower Pollen Extract

Rye flower pollen extract reduces lower urinary-tract symptoms by selectively inhibiting the growth of prostate cells and effectively treating BPH.⁴⁹⁻⁵³

Confirming early findings,⁵⁰⁻⁵³ scientists treated 79 BPH patients (ages 62 to 89) with **126 mg** of pollen extract three times daily for at least 12 weeks. Maximum urine-flow rates increased about **18**%, average urine-flow rates increased nearly **18**%, and residual urine volume (urine left in the bladder after voiding) plummeted **45**%. Those taking the extract beyond one year experienced an average decrease in prostate volume of about **20**%, without adverse reaction.⁵⁴

A subsequent review revealed that BPH patients who were treated with pollen extract were **2.4 times** more likely to experience improvement and over **two times** as likely to reduce nighttime urination as the placebo group.⁵⁵

Pollen extracts have also demonstrated an ability to suppress prostatitis and prostatodynia (chronic pelvic pain). In early human studies, pollen extracts eliminated these conditions from many patients.^{56,57} This is exciting, given how difficult these conditions are to treat.

One study divided 90 chronic **prostatitis** patients into two groups, one <u>without</u> related complicating factors and one <u>with</u> complications such as prostate stones or bladder-neck narrowing. All took pollen extract three times daily for six months.⁵⁷ In the group without complicating factors, **78**% had a favorable response in their symptoms and **36**% experienced *complete elimination* of symptoms. Patients in the group with

complicating factors did not respond as well,⁵⁷ suggesting that pollen extract is most useful in patients who lack complications.

Landmark research showed that pollen extract may help patients for *whom no standardized treatment exists*. Patients with inflammatory prostatitis/chronic pelvic pain syndrome received pollen extract for 12 weeks. Over **70**% of supplemented patients showed at least a **25**% improvement in their **NIH Chronic Prostatitis Symptom Index** score.⁵⁸

Flower pollen extract was shown to improve the quality of life of patients with chronic prostatitis or chronic pelvic pain syndrome better than ibuprofen,⁵⁹ without severe side effects.^{59,60}

Pumpkin Seed

Pumpkin seeds have been used in traditional medicine for generations to treat prostate issues. Now, modern medicine is confirming their ability to reduce prostate size as well as improve symptoms associated with enlarged prostate.

A **2016** study on hyperplastic (proliferated) cells from prostate tissue showed that pumpkin seed extract safely inhibited cell growth by a remarkable **40%-50%**.⁶¹

The study author concluded that this "corroborates the (traditional medicine) use of pumpkin seeds for treatment of benign prostate hyperplasia."⁶¹

Clinical studies have shown that pumpkin seed and pumpkin-seed oil significantly decreased **International Prostate Symptom Scores** within 3-12 months.^{62,63}

In **2016**, researchers conducted a review of studies related to lower urinary tract symptoms and BPH and found that all six clinical studies demonstrated that pumpkin seeds led to improvement in **International Prostate Symptom Score** and in volume and speed of urinary flow.64

Flaxseed

BPH patients who took flaxseed lignan extract experienced improvements in International Prostate Symptom Score and life-quality scores. 65 In a 2017 study of animals with induced BPH, a flaxseed diet was shown to reduce the thickness of the prostate epithelium, the outer layer of the prostate surface.⁶⁶

Summary

The life-disrupting, lower urinary-tract symptoms of BPH affect aging men and can lead to complications such as bladder infections, kidney damage, and sexual dysfunction.

The frequent nighttime urination associated with BPH also results in chronic sleep deprivation, which has been shown to boost the odds of heart attack, 5,6 cancer, 7,8 depression, 9 inflammatory syndromes, 10,11 and death.12

Abundant research demonstrates that a number of prostate-specific **plant extracts** can improve many of the symptoms associated with an enlarged prostate and may also reduce prostate gland volume without significant side effects.



Natural Prevention of Prostate Cancer

Prostate cancer is the second most common malignancy among men,67 with more than 160,000 American men expected to be diagnosed in 2017, according to the American Cancer Society.⁶⁸ It is also one of the most preventable cancers—because it tends to be very slow-growing and nutritional approaches can be highly effective.⁶⁹ The following is a list of nutrients that have been found to protect against the development of prostate cancer.

Lycopene: High consumption of lycopene—the carotenoid pigment abundant in tomatoes and other red fruits and vegetables—delivers potent effects against prostate cancer and is associated with a 59% lower risk of dying from more aggressive prostate cancers. 70-73 One 2016 clinical trial on 79 prostate cancer patients demonstrated that lycopene-rich tomato products significantly decreased PSA compared to controls.⁷⁴ By suppressing critical "master regulatory molecules," such as nuclear factor-kappa B (NF-kB), lycopene inhibits inflammatory processes that promote prostate—and many other—cancers.75

Pygeum: Research demonstrated that administering the bark extract of Pygeum africanum to mice specifically bred to develop prostate cancers significantly lowered their risk of developing this malignancy. Pygeum applied directly to prostate cancer cells in culture inhibited cell proliferation, induced apoptosis, and bound to androgen

receptors used by the tumor to sustain growth.⁶⁹ When serum taken from a man who was supplementing with pygeum extract was applied to prostate cells in culture, it decreased proliferation of prostate cells and upregulated genes involved in tumor suppression.76

Boswellia extract: When an extract of Boswellia serrata is applied to cultured prostate-cancer cells, it induces apoptosis.77-80 Research showed that boswellia components may prevent tumor growth by blocking receptors for androgen, the male hormone,81 and by inhibiting the formation of new blood vessels (angiogenesis) which further deprives tumors of nutrients.54

Flaxseed: Scientists demonstrated that flaxseed lowers PSA levels and significantly reduces proliferation of both normal and cancerous prostate cells.82,83 In a clinical study, flaxseed reduced tumor proliferation rates in prostate cancer patients in as few as 30 days.82

Boron: Men with the highest dietary boron intakes have a 54% lower risk of prostate cancer compared to those with the lowest intake.84 Boron blocks growth factors necessary for tumor development and inhibits the enzymatic action of PSA.85 Human prostate cancers implanted in mice were smaller by 38% following lowdose boron supplementation while serum PSA levels fell 89%.85

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. Sarma AV, Wei JT. Clinical practice. Benign prostatic hyperplasia and lower urinary tract symptoms. N Engl J Med. 2012;367(3):248-
- 2. Available at: https://www.niddk.nih.gov/health-information/ urologic-diseases/prostate-problems/prostate-enlargement-benignprostatic-hyperplasia. Accessed May 1, 2017.
- Roehrborn CG. Male lower urinary tract symptoms (LUTS) and benign prostatic hyperplasia (BPH). Med Clin North Am. 2011;95(1):87-100.
- Speakman MJ, Cheng X. Management of the complications of BPH/BOO. Indian J Urol. 2014;30(2):208-13.
- 5. Chandola T, Ferrie JE, Perski A, et al. The effect of short sleep duration on coronary heart disease risk is greatest among those with sleep disturbance: a prospective study from the Whitehall II cohort. Sleep. 2010;33(6):739-44.
- Nagai M, Hoshide S, Kario K. Sleep duration as a risk factor for cardiovascular disease- a review of the recent literature. Curr Cardiol Rev. 2010;6(1):54-61.
- 7. Kakizaki M, Inoue K, Kuriyama S, et al. Sleep duration and the risk of prostate cancer: the Ohsaki Cohort Study. Br J Cancer. 2008;99(1):176-8.
- Graci G. Pathogenesis and management of cancer-related insomnia. J Support Oncol. 2005;3(5):349-59.
- Germain A, Kupfer DJ. Circadian rhythm disturbances in depression. Hum Psychopharmacol. 2008;23(7):571-85.
- 10. Power JD, Perruccio AV, Badley EM. Pain as a mediator of sleep problems in arthritis and other chronic conditions. Arthritis Rheum. 2005;53(6):911-9.
- 11. Vgontzas AN, Papanicolaou DA, Bixler EO, et al. Circadian interleukin-6 secretion and quantity and depth of sleep. J Clin Endocrinol Metab. 1999;84(8):2603-7.
- 12. Available at: http://www.webmd.com/urinary-incontinence-oab/ news/20100601/nighttime-urination-linked-to-higher-death-risk. Accessed May 1, 2017.
- 13. Available at: https://www.ncbi.nlm.nih.gov/books/NBK301/. Accessed May 1, 2017.
- 14. Schiller DS, Parikh A. Identification, pharmacologic considerations, and management of prostatitis. Am J Geriatr Pharmacother. 2011:9(1):37-48.
- 15. Hamilton W, Sharp D. Symptomatic diagnosis of prostate cancer in primary care: a structured review. Br J Gen Pract. 2004;54(505):617-21.
- 16. Vahlensieck W, Jr. With alpha blockers, finasteride and nettle root against benign prostatic hyperplasia. Which patients are helped by conservative therapy? MMW Fortschr Med. 2002;144(16):33-6.
- 17. Berges R. Epidemiology of benign prostatic syndrome. Associated risks and management data in German men over age 50. Urologe A. 2008:47(2):141-8.
- 18. Kok ET, Schouten BW, Bohnen AM, et al. Risk factors for lower urinary tract symptoms suggestive of benign prostatic hyperplasia in a community based population of healthy aging men: the Krimpen Study. J Urol. 2009;181(2):710-6.
- 19. Glina S, Glina FP. Pathogenic mechanisms linking benign prostatic hyperplasia, lower urinary tract symptoms and erectile dysfunction. Ther Adv Urol. 2013;5(4):211-8.
- 20. Singh DV, Mete UK, Mandal AK, et al. A comparative randomized prospective study to evaluate efficacy and safety of combination of tamsulosin and tadalafil vs. tamsulosin or tadalafil alone in patients with lower urinary tract symptoms due to benign prostatic hyperplasia. J Sex Med. 2014;11(1):187-96.

- 21. Yan H, Zong H, Cui Y, et al. The efficacy of PDE5 inhibitors alone or in combination with alpha-blockers for the treatment of erectile dysfunction and lower urinary tract symptoms due to benign prostatic hyperplasia: a systematic review and meta-analysis. J Sex Med. 2014;11(6):1539-45.
- 22. Available at: https://my.clevelandclinic.org/health/articles/kidneyinfection-pyelonephritis. Accessed May 3, 2017.
- 23. Available at: http://www.mayoclinic.org/diseases-conditions/bladder-stones/symptoms-causes/dxc-20233507. Accessed May 3, 2017.
- 24. Wagenlehner FM, Bschleipfer T, Pilatz A, et al. Pollen extract for chronic prostatitis-chronic pelvic pain syndrome. Urol Clin North Am. 2011;38(3):285-92.
- 25. Monden K, Tsugawa M, Ninomiya Y, et al. [A Japanese version of the National Institutes of Health Chronic Prostatitis Symptom Index (NIH-CPSI, Okayama version) and the clinical evaluation of cernitin pollen extract for chronic non-bacterial prostatitis]. Nihon Hinyokika Gakkai Zasshi. 2002;93(4):539-47.
- 26. Nickel JC. Treatment of chronic prostatitis/chronic pelvic pain syndrome. Int J Antimicrob Agents. 2008;31 Suppl 1:S112-6.
- 27. Potts JM. Therapeutic options for chronic prostatitis/chronic pelvic pain syndrome. Curr Urol Rep. 2005;6(4):313-7.
- 28. Available at: http://emedicine.medscape.com/article/2002872-overview. Accessed May 3, 2017.
- 29. Gurunadha Rao Tunuguntla HS, Evans CP. Management of prostatitis. Prostate Cancer Prostatic Dis. 2002;5(3):172-9.
- 30. Nickel JC. Perplexing problem of persistently painful prostatitis. Rev Urol. 1999;1(3):160-9.
- 31. Fong YK, Milani S, Djavan B. Role of phytotherapy in men with lower urinary tract symptoms. Curr Opin Urol. 2005;15(1):45-8.
- 32. Suter A, Saller R, Riedi E, et al. Improving BPH symptoms and sexual dysfunctions with a saw palmetto preparation? Results from a pilot trial. Phytother Res. 2013;27(2):218-26.
- 33. Allkanjari O, Vitalone A. What do we know about phytotherapy of benign prostatic hyperplasia? Life Sci. 2015;126:42-56.
- 34. Boyle P, Robertson C, Lowe F, et al. Updated meta-analysis of clinical trials of Serenoa repens extract in the treatment of symptomatic benign prostatic hyperplasia. BJU Int. 2004;93(6):751-6.
- 35. Wilt T, Ishani A, Mac Donald R. Serenoa repens for benign prostatic hyperplasia. Cochrane Database Syst Rev. 2002(3):Cd001423.
- 36. Comhaire F, Mahmoud A. Preventing diseases of the prostate in the elderly using hormones and nutriceuticals. Aging Male. 2004;7(2):155-69.
- 37. Gerber GS. Saw palmetto for the treatment of men with lower urinary tract symptoms. J Urol. 2000;163(5):1408-12.
- 38. Tacklind J, MacDonald R, Rutks I, et al. Serenoa repens for benign prostatic hyperplasia. Cochrane Database Syst Rev. 2009(2):Cd001423.
- 39. Chrubasik JE, Roufogalis BD, Wagner H, et al. A comprehensive review on the stinging nettle effect and efficacy profiles. Part II: urticae radix. Phytomedicine. 2007;14(7-8):568-79.
- 40. Moradi HR, Erfani Majd N, Esmaeilzadeh S, et al. The histological and histometrical effects of Urtica dioica extract on rat's prostate hyperplasia. Vet Res Forum. 2015;6(1):23-9.
- 41. Safarinejad MR. Urtica dioica for treatment of benign prostatic hyperplasia: a prospective, randomized, double-blind, placebocontrolled, crossover study. J Herb Pharmacother. 2005;5(4):1-11.
- 42. Engelmann U, Walther C, Bondarenko B, et al. Efficacy and safety of a combination of sabal and urtica extract in lower urinary tract symptoms. A randomized, double-blind study versus tamsulosin. Arzneimittelforschung. 2006;56(3):222-9.
- 43. Oelke M, Berges R, Schlafke S, et al. Fixed-dose combination PRO 160/120 of sabal and urtica extracts improves nocturia in men with LUTS suggestive of BPH: re-evaluation of four controlled clinical studies. World J Urol. 2014;32(5):1149-54.
- 44. Sokeland J. Combined sabal and urtica extract compared with finasteride in men with benign prostatic hyperplasia: analysis of prostate volume and therapeutic outcome. BJU Int. 2000;86(4):439-
- 45. Lopatkin N, Sivkov A, Schlafke S, et al. Efficacy and safety of a combination of Sabal and Urtica extract in lower urinary tract symptoms--long-term follow-up of a placebo-controlled, doubleblind, multicenter trial. Int Urol Nephrol. 2007;39(4):1137-46.

- 46. Breza J, Dzurny O, Borowka A, et al. Efficacy and acceptability of tadenan (Pygeum africanum extract) in the treatment of benign prostatic hyperplasia (BPH): a multicentre trial in central Europe. Curr Med Res Opin. 1998;14(3):127-39.
- 47. Chatelain C, Autet W, Brackman F. Comparison of once and twice daily dosage forms of Pygeum africanum extract in patients with benign prostatic hyperplasia: a randomized, double-blind study. with long-term open label extension. Urology. 1999;54(3):473-8.
- 48. Wilt T, Ishani A, Mac Donald R, et al. Pygeum africanum for benign prostatic hyperplasia. Cochrane Database Syst Rev. 2002(1):Cd001044.
- 49. Habib FK, Ross M, Buck AC, et al. In vitro evaluation of the pollen extract, cernitin T-60, in the regulation of prostate cell growth. Br J Urol. 1990:66(4):393-7
- 50. Buck AC, Cox R, Rees RW, et al. Treatment of outflow tract obstruction due to benign prostatic hyperplasia with the pollen extract, cernilton. A double-blind, placebo-controlled study. Br J Urol. 1990;66(4):398-404.
- 51. Horii A, Iwai S, Maekawa M, et al. [Clinical evaluation of cernilton in the treatment of the benign prostatic hypertrophy]. Hinyokika Kivo. 1985:31(4):739-46.
- 52. Hayashi J, Mitsui H, Yamakawa G, et al. [Clinical evaluation of Cernilton in benign prostatic hypertrophy]. Hinyokika Kiyo. 1986;32(1):135-41.
- 53. Ueda K. Jinno H. Tsujimura S. [Clinical evaluation of Cernilton on benign prostatic hyperplasia]. Hinyokika Kiyo. 1985;31(1):187-91.
- Yasumoto R, Kawanishi H, Tsujino T, et al. Clinical evaluation of long-term treatment using cernitin pollen extract in patients with benign prostatic hyperplasia. Clin Ther. 1995;17(1):82-7.
- 55. MacDonald R, Ishani A, Rutks I, et al. A systematic review of Cernilton for the treatment of benign prostatic hyperplasia. BJU Int. 2000;85(7):836-41.
- 56. Buck AC, Rees RW, Ebeling L. Treatment of chronic prostatitis and prostatodynia with pollen extract. Br J Urol. 1989;64(5):496-9.
- 57. Rugendorff EW, Weidner W, Ebeling L, et al. Results of treatment with pollen extract (Cernilton N) in chronic prostatitis and prostatodynia. Br J Urol. 1993;71(4):433-8.
- 58. Wagenlehner FM, Schneider H, Ludwig M, et al. A pollen extract (Cernilton) in patients with inflammatory chronic prostatitischronic pelvic pain syndrome: a multicentre, randomised, prospective, double-blind, placebo-controlled phase 3 study. Eur Urol. 2009;56(3):544-51.
- 59. Cai T. Wagenlehner FM. Luciani LG. et al. Pollen extract in association with vitamins provides early pain relief in patients affected by chronic prostatitis/chronic pelvic pain syndrome. Exp Ther Med. 2014;8(4):1032-8.
- 60. Iwamura H, Koie T, Soma O, et al. Eviprostat has an identical effect compared to pollen extract (Cernilton) in patients with chronic prostatitis/chronic pelvic pain syndrome: a randomized, prospective study. BMC Urol. 2015;15:120.
- 61. Medjakovic S, Hobiger S, Ardjomand-Woelkart K, et al. Pumpkin seed extract: Cell growth inhibition of hyperplastic and cancer cells, independent of steroid hormone receptors. Fitoterapia. 2016;110:150-6.
- 62. Hong H, Kim CS, Maeng S. Effects of pumpkin seed oil and saw palmetto oil in Korean men with symptomatic benign prostatic hyperplasia. Nutr Res Pract. 2009;3(4):323-7.
- 63. Vahlensieck W, Theurer C, Pfitzer E, et al. Effects of pumpkin seed in men with lower urinary tract symptoms due to benign prostatic hyperplasia in the one-year, randomized, placebo-controlled GRANU study. Urol Int. 2015;94(3):286-95.
- 64. Damiano R, Cai T, Fornara P, et al. The role of Cucurbita pepo in the management of patients affected by lower urinary tract symptoms due to benign prostatic hyperplasia: A narrative review. Arch Ital Urol Androl. 2016;88(2):136-43.
- 65. Zhang W, Wang X, Liu Y, et al. Effects of dietary flaxseed lignan extract on symptoms of benign prostatic hyperplasia. J Med Food. 2008;11(2):207-14.
- 66. de Amorim Ribeiro IC, da Costa CA, da Silva VA, et al. Flaxseed reduces epithelial proliferation but does not affect basal cells in induced benign prostatic hyperplasia in rats. Eur J Nutr. 2017;56(3):1201-10.

- 67. Mariani S, Lionetto L, Cavallari M, et al. Low prostate concentration of lycopene is associated with development of prostate cancer in patients with high-grade prostatic intraepithelial neoplasia. Int J Mol Sci. 2014;15(1):1433-40.
- 68. Available at: https://www.cancer.org/cancer/prostate-cancer/about/ key-statistics.html. Accessed May 3, 2017.
- 69. Shenouda NS, Sakla MS, Newton LG, et al. Phytosterol Pygeum africanum regulates prostate cancer in vitro and in vivo. Endocrine. 2007;31(1):72-81.
- 70. Borel P, Desmarchelier C, Nowicki M, et al. Lycopene bioavailability is associated with a combination of genetic variants. Free Radic Biol Med. 2015;83:238-44.
- 71. Grainger EM, Hadley CW, Moran NE, et al. A comparison of plasma and prostate lycopene in response to typical servings of tomato soup, sauce or juice in men before prostatectomy. Br J Nutr. 2015:114(4):596-607.
- 72. Holzapfel NP, Holzapfel BM, Champ S, et al. The potential role of lycopene for the prevention and therapy of prostate cancer: from molecular mechanisms to clinical evidence. Int J Mol Sci. 2013;14(7):14620-46.
- 73. Wang Y, Jacobs EJ, Newton CC, et al. Lycopene, tomato products and prostate cancer-specific mortality among men diagnosed with nonmetastatic prostate cancer in the Cancer Prevention Study-II Nutrition Cohort. Int J Cancer. 2016.
- 74. Paur I, Lilleby W, Bohn SK, et al. Tomato-based randomized controlled trial in prostate cancer patients: Effect on PSA. Clin Nutr. 2017;36(3):672-9.
- 75. Assar EA, Vidalle MC, Chopra M, et al. Lycopene acts through inhibition of IkappaB kinase to suppress NF-kappaB signaling in human prostate and breast cancer cells. Tumour Biol. 2016:37(7):9375-85
- 76. Larre S, Camparo P, Comperat E, et al. Biological effect of human serum collected before and after oral intake of Pygeum africanum on various benign prostate cell cultures. Asian J Androl. 2012;14(3):499-504.
- 77. Buchele B, Zugmaier W, Estrada A, et al. Characterization of 3alpha-acetyl-11-keto-alpha-boswellic acid, a pentacyclic triterpenoid inducing apoptosis in vitro and in vivo. Planta Med. 2006;72(14):1285-9.
- 78. El Gaafary M, Buchele B, Syrovets T, et al. An alpha-acetoxytirucallic acid isomer inhibits Akt/mTOR signaling and induces oxidative stress in prostate cancer cells. J Pharmacol Exp Ther. 2015:352(1):33-42
- 79. Lu M, Xia L, Hua H, et al. Acetyl-keto-beta-boswellic acid induces apoptosis through a death receptor 5-mediated pathway in prostate cancer cells. Cancer Res. 2008;68(4):1180-6.
- 80. Pang X, Yi Z, Zhang X, et al. Acetyl-11-keto-beta-boswellic acid inhibits prostate tumor growth by suppressing vascular endothelial growth factor receptor 2-mediated angiogenesis. Cancer Res. 2009:69(14):5893-900.
- 81. Yuan HQ, Kong F, Wang XL, et al. Inhibitory effect of acetyl-11-keto-beta-boswellic acid on androgen receptor by interference of Sp1 binding activity in prostate cancer cells. Biochem Pharmacol. 2008;75(11):2112-21.
- 82. Demark-Wahnefried W, Polascik TJ, George SL, et al. Flaxseed supplementation (not dietary fat restriction) reduces prostate cancer proliferation rates in men presurgery. Cancer Epidemiol Biomarkers Prev. 2008;17(12):3577-87.
- 83. Demark-Wahnefried W, Robertson CN, Walther PJ, et al. Pilot study to explore effects of low-fat, flaxseed-supplemented diet on proliferation of benign prostatic epithelium and prostate-specific antigen. Urology. 2004;63(5):900-4.
- 84. Cui Y, Winton MI, Zhang ZF, et al. Dietary boron intake and prostate cancer risk. Oncol Rep. 2004;11(4):887-92.
- 85. Gallardo-Williams MT, Chapin RE, King PE, et al. Boron supplementation inhibits the growth and local expression of IGF-1 in human prostate adenocarcinoma (LNCaP) tumors in nude mice. Toxicol Pathol. 2004;32(1):73-8.



Super Miraforte with Standardized Lignans Item #01940 • 120 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$62	\$46.50
4 bottles		\$42 each



* J Clin Endocrinol Metab. 2002 Feb;87(2):589-98.



Low Testosterone Levels May Lead to:

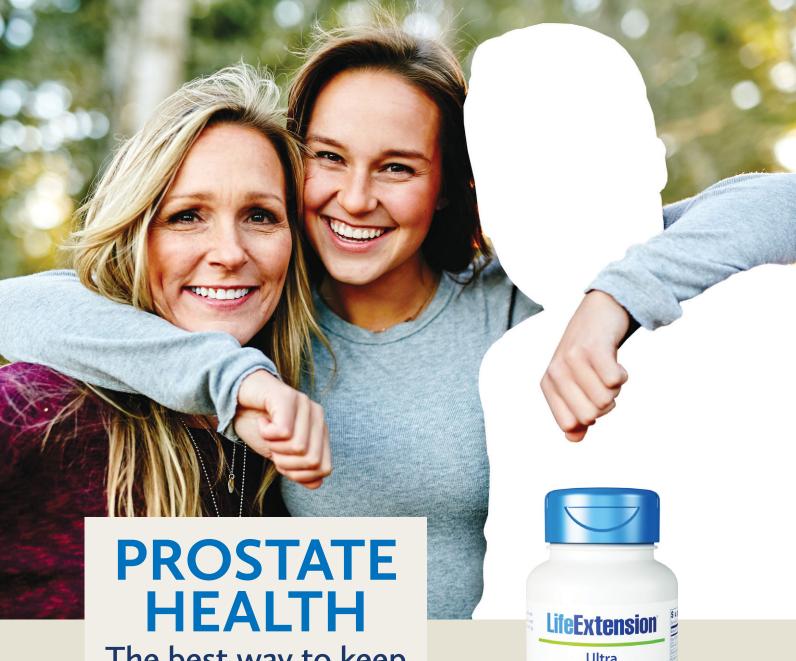
- Reduced Sex Drive
- Less Energy
- Cloudy Thinking
- Weight Gain
- Cardiovascular Issues

For full product description and to order **Super Miraforte with Standardized Lignans**, call **1-800-544-4440** or visit **www.LifeExtension.com**

Caution: If you are taking any medication, use only under physician supervision. Men with existing prostate cancer may not be able to use this product.

Elevations in free testosterone can unmask an occult (hidden) prostate cancer. Anyone with this concern should have a baseline PSA prior to using this product and a follow-up PSA test 60 days later. If a significant elevation of PSA is found, discontinue this product and advise physician. Do not take more than 15 mg per day of Bioperine®.

Bioperine[®] is a registered trademark of Sabinsa Corp. HMRlignan[®] is a trademark used under sublicense from Linnea S.A.



The best way to keep You in the picture.

As a man ages, maintaining a healthy prostate is key. We created Ultra Natural Prostate to help you maintain prostate health, so you can focus on what's important. With over a dozen natural ingredients, this supplement promotes healthy prostate function, supports easier urination, inhibits inflammatory factors, and encourages natural division of prostate cells. Ultra Natural Prostate. The most comprehensive prostate health supplement.



Item #01928 • 60 softgels

6	Retail Price	YOUR PRICE
1-bottle	\$38.00	\$28.50
4-bottle	_	\$26.25 ea
Non-GMO		

For full product description and to order Ultra Natural Prostate, call 1-800-544-4440 or visit www.LifeExtension.com

ApresFlex® 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.



Now Delivering into California!

- Full Compounding Lab
- Full Retail Pharmacy
- Bio-Identical Hormone Replacement Therapy
- Free Standard Delivery/Shipping
- Durable Medical Equipment
- Trilingual (English, Spanish, French)
- Licensed to Ship into 42 States

Renew Rx's online to

RECEIVE \$1 OFF EACH Rx

(Cash Rx's only)

Did you know we carry a full line of PET products? Mention this ad to

RECEIVE 10% OFF your first order

Quality, Reliability, Integrity

Lowest Prices: Ask for Customer Discount Pricing

Providing Trusted Prescription Compounding for Over 45 Years!

Proud Members of



(877) 877-9700

Phone: (954) 989-6524 | Fax: (866) 892-3432

4401 Sheridan St. Hollywood, FL 33021 www.PostHastePharmacy.com



active forms.

BioActive Complete B-Complex provides *enzymati*cally active forms of meaningful potencies of each B vitamin. This includes the *pyridoxal 5'-phosphate* form of vitamin B6 (the metabolically active form, shown to protect lipids and proteins against glycation reactions) and the most biologically active form of **folate** called 5-methyltetrahydrofolate (5-MTHF), which is up to 7 times more bioavailable than folic acid.1

For full product description and to order **BioActive Complete B-Complex**, call 1-800-544-4440 or visit www.LifeExtension.com

BioActive Complete B-Complex

Item #01945 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$12	\$9
4 bottles		\$8 each

Non-GMO

1. Br J Pharmacol. 2004 Mar: 141(5):825-30.





REVERSING a Root Cause of GLAUCOMA

Glaucoma is one of the leading causes of blindness, affecting approximately **2.7 million** Americans.¹

In the past year, studies have found new links between **glaucoma** and a host of diverse risk factors, such as tooth loss,² obstructive sleep apnea,³ genetic predisposition⁴ and potentially diabetes.⁵ Drugs such as corticosteroids are also implicated in glaucoma risk.⁶

The most common risk factor for glaucoma is increased pressure in the eye, which can cause irreversible damage to the optic nerve.⁷

There is usually no pain associated with increased eye pressure,⁸ which means you could be at risk for glaucoma—and not even know it.

But while glaucoma itself is irreversible, studies have shown that it is possible to prevent—or even *reverse*—the major underlying <u>cause</u>.

In a human study, a proprietary extract of **French** maritime pine bark combined with bilberry extract reduced eye pressure by as much as **24%**—with reductions of **40%** when combined with standard therapy.⁹

A Vision-Robbing Disease— Without Warning Signs

People who are developing **glaucoma** generally have no symptoms. They feel no pain.^{8,10}

One day their vision is normal, the next they begin to realize that they are missing some objects that would normally fall within their peripheral vision. At this first sign, the progression that may eventually lead to blindness is well underway.¹⁰

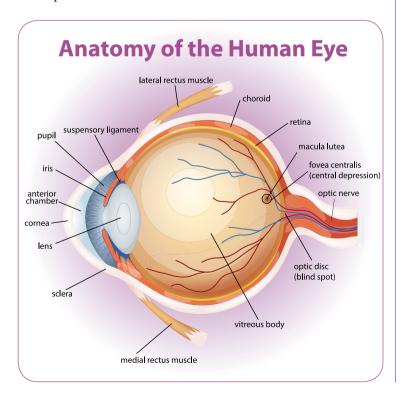
In most cases, glaucoma occurs when fluid builds up, creating abnormal **intraocular pressure** within the eye. Over time, this pressure damages the retina and the **optic nerve**, resulting in reduced visual acuity and possibly leading to blindness.¹¹

What Causes Increased Eye Pressure?

The transparent fluid that fills the anterior part of the eye between the lens and the cornea is called *aqueous humor*. ¹² This fluid has numerous jobs, including providing nutrition to the anterior part of the eye and transporting the metabolic debris produced there to the bloodstream so that we can see clearly.

The appropriate production, circulation, and drainage of this fluid are essential for eye health.

Open angle glaucoma, which is diagnosed in at least **90%** of glaucoma patients, is the most common form of the disease. ¹³ Over time, the drainage channels become blocked, fluid builds up, and intraocular pressure rises. ¹⁰



Additionally, endothelial dysfunction and vascular structural changes can substantially alter blood flow within the tissues and elevate intraocular pressure, leading eventually to open angle glaucoma.¹⁴

Whether or not you develop glaucoma as a result of increased intraocular pressure depends on the level of pressure your optic nerve can tolerate without being damaged. ¹⁰ However, once the optic nerve is damaged, it can't be repaired—*even if the raised intraocular pressure is corrected*. ^{15,16}

This makes it extremely urgent to find a way to reverse high intraocular pressure *before* it causes the irreversible damage of glaucoma.

The Search for a Solution

Scientists turned to past research to find the best possible natural candidates for treating this condition. Their search led them to **French maritime pine bark** and standardized **bilberry extract**.

Previous studies had shown that these extracts could successfully counteract **retinopathy**, which is persistent or acute damage to the retina. ^{9,17} This led investigators to examine the usefulness of these plant extracts in countering the drivers behind increased intraocular pressure.

What they found was that French maritime pine bark could improve the function of the **endothelium**, the delicate layer of cells lining the blood vessels. Disorders of *endothelial function* are contributing factors to the development or progression of glaucoma.^{9,14}

Other studies showed that bilberry extract could counteract hyperpermeability of the ciliary capillaries. The beneficial effect is significantly increased ocular blood flow, resulting in <u>reduced</u> intraocular pressure.¹⁷

It became clear that these **two extracts** may work together to:

- Decrease inflow of aqueous humor;
- Improve microvascular tone and integrity;
- Decrease resistance across the region of the eye responsible for fluid drainage, and possibly;
- Contribute to better fluid outflow.

The ability of both bilberry and French maritime pine bark to target critical aspects of increased eye pressure led scientists to formulate a compound that combined these two. The next step was to conduct human studies that tested the dual-extract formulation.

Remarkable Drop in Eye Pressure

In an initial controlled study of this dual compound, scientists measured blood flow in the eyes of 38 volunteers who had high **intraocular pressure** but who had not yet shown evidence of glaucoma. One group took the **pine bark-bilberry compound** orally for six months and the second group did not.¹⁷

At three months, the group taking the pine bark-bilberry compound showed a statistically significant 13% reduction in intraocular pressure. Compared to untreated participants, the treated group also had improved ocular blood flow in three different blood vessels.17

A follow-up study showed that taking the same pine bark-bilberry compound for longer led to even greater improvements.

In this study, 79 individuals with intraocular pressure who had not yet shown signs of open angle glaucoma were divided into three groups:

- 1. The first group received the pine bark-bilberry extract.
- 2. The second group received standard medical therapy with latanoprost (Xalatan[®]) eye drops,
- 3. The third group received both the pine barkbilberry compound and the latanoprost drops.9

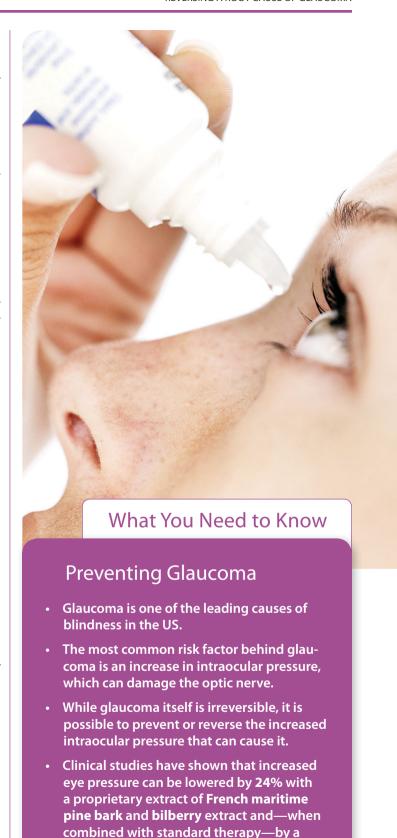
All three treatment groups demonstrated a reduction in intraocular pressure. Subjects using the prescription eye drops lowered their eye pressure by an average of 28%, beginning from the fourth treatment week. Those participants taking the **pine bark-bilberry** formulation reduced their eye pressure significantly beginning in the sixth treatment week and throughout the study, leading to a **24**% reduction in the sixteenth week—comparable to the drug, but with a better safety profile.9

But by far, the most compelling results were seen in the group that used the combination of pine bark**bilberry** formulation and the latanoprost drops. A significant, average reduction in intraocular pressure of 28% began at four weeks—but when the study ended at 24 weeks, the decrease in eye pressure had reached an approximate 40%!9

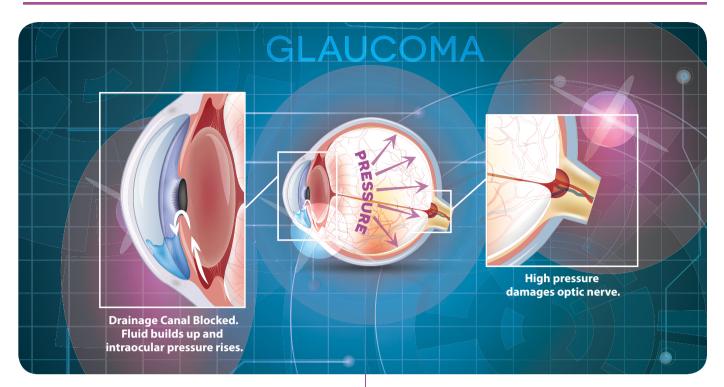
The **pine bark-bilberry** compound appeared to have an additive effect with the latanoprost drops to **amplify** the reduction of intraocular pressure better than either agent alone.9

Critically, the subjects experienced a significant increase of ocular blood flow.9

The study author noted that, "No serious side effects occurred during the study, apart from standard side effects in patients related to latanoprost."9



compelling 40%!



French Maritime Pine-Bark Extract

Numerous studies have given us insight into why these two extracts have such beneficial effects on eye pressure.

Each extract has its own range of actions that appear particularly suited to aiding the complex balance at the level of the eye chambers.⁹

French maritime pine bark is rich in plant-based *proanthocyanidins* and was found to have numerous biologic effects, including:

- The scavenging and neutralization of harmful free radicals,
- Regulation of the cell's antioxidant network and associated genes,
- Anti-inflammatory effects (through the dampening of gene expression related to the nuclear factor-kappaB-dependent pathways inside cells),^{18,19}
- Improved vascular endothelial function,²⁰ and
- Improved microcirculation from antiplatelet effects and clot-formation prevention.^{21,22}

One **2015** study appeared to show beneficial effects on intraocular pressure when volunteers were given **French maritime pine bark** combined with extracts of blueberry and green tea.²³

French maritime pine bark's powerful antioxidative capacity—which can protect the eye's drainage system—is mirrored in the strong, free radical-quenching effects of bilberry extract.

Standardized Bilberry Extract

Bilberry (*Vaccinium myrtillus*) and other related berries are known for superior free radical-scavenging activity as well as genetic signaling ability.²⁴ Bilberry has been shown to bolster the body's defense systems against dangerous oxidative stress,²⁵ and it has also been shown to be beneficial in atherosclerosis.²⁶

Specifically using tissue from the pigmented layer of the retina, scientists found that bilberry positively influenced beneficial pathways involved in the antioxidant response effort.²⁵

Bilberry has also been shown to provide protective effects in other models of inflammatory disease such as uveitis in a dose-dependent manner.²⁷

A Significant Step Toward Prevention

Even with standard medical or surgical therapies, some glaucoma patients still progress to vision loss. ⁸ Unfortunately, this loss is permanent. Until a cure is found, research is urgently needed to identify ways to *prevent* this devastating disease.

There currently is no accepted preventive strategy for glaucoma. The best defense to date involves rigorous and regular eye examinations by a trained professional.

However, delicate eve tissues—under assault by environmental toxins and cellular byproducts associated with aging—can greatly benefit from nutritional and other therapeutic support against glaucoma and other sight-robbing diseases.28

The human studies described in this article are promising and mark an important initial step toward finding ways to prevent glaucoma. The dual-extract formulation of pine bark and bilberry has been shown to lower intraocular pressure by almost 40% in conjunction with prescribed eve drops that do involve some risk.9,29

While this may bring hope to those with elevated eve pressure, it is important to note that high intraocular pressure may not always be the defining characteristic for diagnosing glaucoma or predicting whether the disease will worsen.³⁰ Statistics show that **15%** of patients with characteristic glaucomatous nerve damage have intraocular pressure measurements that fall within the normal range.31 Such cases may be partly due to poor blood flow to the optic nerve.³²

Until a cure is identified, greater research into prevention strategies is needed.

Summary

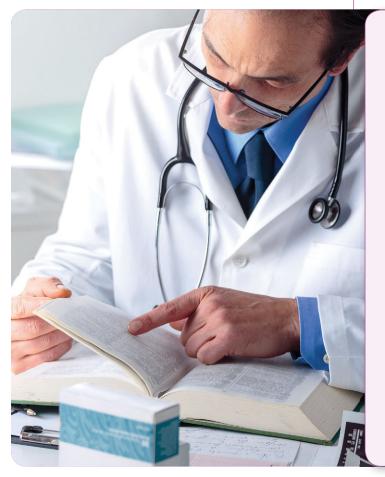
Increased pressure in the eye is the most common underlying cause of glaucoma, and it usually occurs without pain or other warning signs.

Human studies demonstrate that increased intraocular pressure can be significantly reversed with a proprietary extract of **French maritime pine bark** and **bilberry** extract.

In a human study, this formulation reduced eye pressure by 24% and—when combined with standard therapy—by up to **40**%!

While there is no cure for glaucoma, the **pine** bark-bilberry formulation we have described appears to represent a substantial breakthrough along the road to even greater preventive or curative discoveries.

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



Understanding Glaucoma

The term "glaucoma" refers to a common group of similar conditions that damage the retina and optic nerve, leading to visual impairment. There are many risk factors for glaucoma that range from genetics and age to lifestyle factors.

Increased intraocular pressure is by far the most significant factor, and the one most associated with glaucoma. In general, those aged 60 and older are at a higher risk of developing glaucoma regardless of increased intraocular pressure. Amongst ethnic groups, African-Americans have the highest risk for glaucoma in the US. In addition, individuals with history of high blood pressure or diabetes are also at an increased risk. Certain medications, like corticosteroids also increase the risk of alaucoma.33

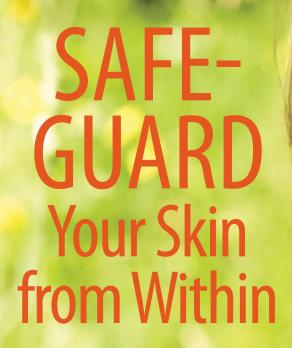
It is important to note that **normal tension** glaucoma can develop in the absence of increased intraocular pressure and cause optic nerve damage. People with family history and those of Japanese ancestry are at a higher risk for this type of glaucoma.34

References

- 1. Available at: https://www.cdc.gov/visionhealth/research/projects/ ongoing/glaucoma.htm#ref1. Accessed May 30, 2017.
- 2. Pasquale LR, Hyman L, Wiggs JL, et al. Prospective Study of Oral Health and Risk of Primary Open-Angle Glaucoma in Men: Data from the Health Professionals Follow-up Study. Ophthalmology. 2016;123(11):2318-27.
- Chaitanya A, Pai VH, Mohapatra AK, et al. Glaucoma and its association with obstructive sleep apnea: A narrative review. Oman J Ophthalmol. 2016;9(3):125-34.
- 4. Miller MA, Fingert JH, Bettis DI. Genetics and genetic testing for glaucoma. Curr Opin Ophthalmol. 2017;28(2):133-8.
- Song BJ, Aiello LP, Pasquale LR. Presence and Risk Factors for Glaucoma in Patients with Diabetes. Curr Diab Rep. 2016:16(12):124.
- Schlote T. Impact of Drugs on Glaucoma and Intraocular Pressure. Klin Monbl Augenheilkd. 2017;234(2):179-84.
- Available at: http://www.glaucoma.org/glaucoma/does-blood-pressure-affect-glaucoma.php. Accessed May 31, 2017.
- Available at: http://www.glaucoma.org/glaucoma/glaucoma-factsand-stats.php. Accessed May 31, 2017.
- Steigerwalt RD, Jr., Belcaro G, Morazzoni P, et al. Mirtogenol potentiates latanoprost in lowering intraocular pressure and improves ocular blood flow in asymptomatic subjects. Clin Ophthalmol. 2010;4:471-6.
- 10. Available at: https://nei.nih.gov/health/glaucoma/glaucoma_facts. Accessed May 31, 2017.
- 11. Aung T, Lim MC, Chan YH, et al. Configuration of the drainage angle, intraocular pressure, and optic disc cupping in subjects with chronic angle-closure glaucoma. Ophthalmology. 2005;112(1):28-32.
- 12. Available at: https://www.mercy.net/healthinfo/hw121946. Accessed May 31, 2017.
- 13. Available at: http://www.glaucoma.org/glaucoma/types-of-glaucoma. php. Accessed May 31, 2017.
- 14. Resch H, Garhofer G, Fuchsjager-Mayrl G, et al. Endothelial dysfunction in glaucoma. Acta Ophthalmol. 2009;87(1):4-12.
- Moore DL, Goldberg JL. Four steps to optic nerve regeneration. J Neuroophthalmol. 2010;30(4):347-60.
- 16. Shum JW, Liu K, So KF. The progress in optic nerve regeneration, where are we? Neural Regen Res. 2016;11(1):32-6.
- 17. Steigerwalt RD, Gianni B, Paolo M, et al. Effects of Mirtogenol on ocular blood flow and intraocular hypertension in asymptomatic subjects. Mol Vis. 2008;14:1288-92.
- 18. Rohdewald P. A review of the French maritime pine bark extract (Pycnogenol), a herbal medication with a diverse clinical pharmacology. Int J Clin Pharmacol Ther. 2002;40(4):158-68.

- 19. Peng Q, Wei Z, Lau BH. Pycnogenol inhibits tumor necrosis factoralpha-induced nuclear factor kappa B activation and adhesion molecule expression in human vascular endothelial cells. Cell Mol Life Sci. 2000;57(5):834-41.
- 20. Nishioka K, Hidaka T, Nakamura S, et al. Pycnogenol, French maritime pine bark extract, augments endothelium-dependent vasodilation in humans. Hypertens Res. 2007;30(9):775-80.
- 21. Cesarone MR, Belcaro G, Rohdewald P, et al. Prevention of edema in long flights with Pycnogenol. Clin Appl Thromb Hemost. 2005;11(3):289-94.
- 22. Belcaro G, Cesarone MR, Rohdewald P, et al. Prevention of venous thrombosis and thrombophlebitis in long-haul flights with pycnogenol. Clin Appl Thromb Hemost. 2004;10(4):373-7.
- 23. Karhanova M, Eliasova M, Kubena T, et al. [ProVens(R) in the Therapy of Glaucoma and Ocular Hypertension]. Cesk Slov Oftalmol.71(6):288-92.
- 24. Zafra-Stone S, Yasmin T, Bagchi M, et al. Berry anthocyanins as novel antioxidants in human health and disease prevention. Mol Nutr Food Res. 2007;51(6):675-83.
- 25. Milbury PE, Graf B, Curran-Celentano JM, et al. Bilberry (Vaccinium myrtillus) anthocyanins modulate heme oxygenase-1 and glutathione S-transferase-pi expression in ARPE-19 cells. Invest Ophthalmol Vis Sci. 2007;48(5):2343-9.
- 26. Mauray A, Felgines C, Morand C, et al. Bilberry anthocyanin-rich extract alters expression of genes related to atherosclerosis development in aorta of apo E-deficient mice. Nutr Metab Cardiovasc Dis. 2012;22(1):72-80.
- 27. Yao N, Lan F, He RR, et al. Protective effects of bilberry (Vaccinium myrtillus L.) extract against endotoxin-induced uveitis in mice. J Agric Food Chem. 2010;58(8):4731-6.
- 28. Baltmr A, Duggan J, Nizari S, et al. Neuroprotection in glaucoma -Is there a future role? Exp Eye Res. 2010;91(5):554-66.
- 29. Available at: http://www.glaucoma.org/gleams/glaucoma-medications-and-their-side-effects.php. Accessed June 2, 2017.
- 30. Varma R, Peeples P, Walt JG, et al. Disease progression and the need for neuroprotection in glaucoma management. Am J Manag Care. 2008;14(1 Suppl):S15-9.
- 31. Distelhorst JS, Hughes GM. Open-angle glaucoma. Am Fam Physician. 2003;67(9):1937-44.
- 32. Available at: http://www.umm.edu/health/medical/reports/articles/ glaucoma. Accessed June 2, 2017.
- 33. Available at: http://www.mayoclinic.org/diseases-conditions/glaucoma/basics/risk-factors/con-20024042. Accessed June 12, 2017.
- 34. Available at: http://www.glaucoma.org/glaucoma/normal-tensionglaucoma.php. Accessed June 12, 2017.





Unique ORAL formula provides
Polypodium leucotomos fern extract
along with nicotinamide and
red orange extract.



Non-GMO Retail Your Price

1 bottle \$44 \$33
4 bottles \$30 each

Item # 01938 • 120 vegetarian capsules

For full product description and to order **Shade Factor**™, call **1-800-544-4440** or visit www.LifeExtension.com

This product is not a substitute for topical sunscreens.

COMPREHENSIVE EYE HEALTH FORMULA



trans-zeaxanthin, and meso-zeaxanthin to help maintain structural integrity of the macula and retina.1-5

Alpha-carotene is included based on new evidence that it helps support the macular pigment.1

People supplementing with saffron showed an improvement in **vision** as measured by them seeing an average of **two additional lines** on the eye chart commonly used by doctors to test vision.1

This formula provides the optimal dose of saffron along with cyanidin-3-glucoside to support healthy vision.6-8

Item #01992 • 60 softgels • Non-GMO

	Retail Price	Your Price
1 bottle	\$25	\$18.75
4 bottles		\$17.50 each

Each bottle lasts for two months.

- **References**1. *JAMA Ophthalmol.* 2015;133(12):1415-24.
- 2. Nutrients. 2013 April;5(4):1169-85.
- 2. Nutrition. 2011 Sep;27(9):960-6. 4. Free Radic Biol Med. 2012;53(6):1298-307. 5. J Ophthalmol. 2015;2015:523027.
- 6. Evid Based Complement Alternat Med. 2012;2012:429124.
- 7. Invest Ophthalmol Vis Sci. 2010;51(12):6118-24
- 8. J Agric Food Chem. 2003 Jun 4;51(12):3560-3



For full product description and to order MacuGuard® Ocular Support, call 1-800-544-4440 or visit www.LifeExtension.com

LuteinPlus® and Mz® are registered trademarks of NutriProducts Ltd., UK, licensed under U.S. Patent 8,623,428.



Clinically Studied in Europe

PRESSURE SUPPORT WITH Mirtogenol®

Eye Pressure Support with Mirtogenol® formula may support healthy blood flow within the tiny vessels (*microvasculature*) of your eyes, helping to promote healthy fluid pressure.

Eye Pressure Support with Mirtogenol®

Item #01514 • 30 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$38	\$28.50
4 bottles		\$25.50 each



For full product description and to order

Eye Pressure Support with Mirtogenol®

call 1-800-544-4440

or visit www.LifeExtension.com

Non-GMO

Mirtogenol® is a registered trademark of Horphag Research, Ltd.
Mirtoselect® is a registered trademark of Indena, S.p.A., Milan, Italy.
Pycnogenol® is a registered trademark of Horphag Research, Ltd.
Pycnogenol® is protected by U.S. patents #5,720,956 and #6,372,266
and other international patents.

CHAILENGE

TODAY'S MEDICAL LIMITS AND TAKE CHARGE OF YOUR LIFESPAN



What if you are living as healthfully as possible but your body finally fails and today's doctors give up on you? Give medicine of the future a chance to repair and rejuvenate you. Let us help you get there.

Call for the facts on cryopreservation.

Cryopreservation is the science of using ultra-cold temperature to preserve human life with the intent of restoring good health when technology becomes available to do so. Call Alcor or visit our website today for your free information package.

Consider opting for the most advanced protection science currently offers to help you preserve your life!



480-905-1906 ext. 101 alcor.org

BY ALMA ROSS

Pomegranate Improves Markers of Aging

Pomegranate's heart benefits have led researchers to investigate in what ways this red fruit can keep us healthy.

In a recent finding, Swiss researchers have identified a new molecule that results from digesting two compounds found in pomegranates: punicalagins and ellagitannins. This unique molecule, known as **urolithin A**, helps rejuvenate **mitochondria**, our cellular powerhouses.¹

Urolithin A opens the door to potential new therapeutic treatments against age-related disorders, including **frailty**, which is a risk factor for disability, hospitalizations, and mortality.²

Research Update



What is Urolithin A?

Urolithin A is produced by the body after ingesting compounds found in pomegranate such as punicalagins and ellagitannins and can help recycle defective mitochondria.

What A New Study Showed

Researchers first studied **Urolithin A** on a common worm called *C. elegans*. This worm is often used in anti-aging studies because after just 8-10 days it's considered elderly. Its short lifespan allows scientists to observe and measure the effects of aging in a little over a week.

The researchers administered **urolithin A** to a group of these worms and noted that **lifespan** in the **urolithin A** group increased by more than **45%** compared to the control group.¹

Next, the team performed several rodent studies and found that **urolithin A** improved muscle function and removed damaged mitochondria before they accumulate and cause cellular dysfunction throughout the body. Scientists know that with age, mitochondria lose their strength and die off thereby "clogging up" cells with debris that impedes their function.

In the first mouse study, urolithin A administration over the long-term was found to *increase* muscle function of aging mice. Compared to the control group, the supplemented group showed a **57**% *increase* in the level of spontaneous exercise measured by the running wheel and a **9**% *increase* in grip strength.¹

In a second mouse study, this time involving a shorter treatment regimen on aged mice, urolithin A was found to increase running endurance by an average of **42**%.¹

Following these findings on aging mice, the team performed another study, also evaluating the impact of urolithin A on <u>muscle function</u> in young rats. Muscle function was evaluated by measuring voluntary running in activity wheels. Once again, treatment with urolithin A proved to be effective, this time by *increasing* the running capacity by **65**% compared to controls.¹

The administration of urolithin A resulted in an enhanced exercise capacity in young and older rodents. Muscle strength *increased* and running endurance was robustly augmented. Together, these different studies highlight that the administration of urolithin A, both short- and long-term, improved muscle function throughout different stages of life by improving muscle quality.

Translating these Findings into Humans

Over time, the constant strain of energy production takes a toll on the mitochondria and energy output declines. At this point, these mitochondria function poorly and are basically useless. In young, healthy cells, the drop in performance of the mitochondria is identified by the body and the mitochondria are swiftly broken down, disassembled, and eliminated in a process called mitophagy. In this way, defective or less-than-optimal mitochondria are eliminated, giving room to new mitochondria and ensuring that optimal cellular function is maintained.

With age, our cells struggle to recycle defective mitochondria, leading to a progressive build-up of malfunctioning mitochondria that take up valuable space in the

Research Update

body's cellular system. This mitochondrial degradation affects the health of the cells, gradually weakening tissues. This process has been suspected of playing a role in many disorders of aging, such as Parkinson's disease.³

In humans, the inability to remove these useless mitochondria in skeletal muscle has been linked to reduced mobility in the elderly. The progressive decline of muscle function contributes to a progressive state of generalized frailty.

In addition, the **frailty** associated with old age is an important risk factor for disability, hospitalization and mortality.² Thus, the muscle weakness seen in the elderly might be due to an increase in the accumulation of useless mitochondria. Results from the rodent studies strongly suggest that improvement of muscle quality may be achieved by enhancing mitochondrial function with **urolithin A**.

The results from these recent Swiss studies suggest that supplementation with **pomegranate extract** to boost the body's content of urolithin A may be an innovative approach to maintaining healthy mitochondrial and muscle function.

Encouraged by their initial findings, the study authors are currently conducting clinical trials testing a special delivery method of finely calibrated doses of urolithin A in humans. These trials are currently taking place in a number of hospitals across Europe.⁴

Urolithin A Helps Fight Cancer

Despite aggressive surgical care and chemotherapy, nearly **50**% of people with colorectal cancers develop recurrent tumors.⁵ This

may be due in part to the survival of dangerous colon-cancer stem cells that resist conventional chemotherapy and act as "seeds" for subsequent cancers.⁶

In an interesting finding, researchers exposed colon-cancer stem cells from a patient with colorectal cancer to either a mixture containing **85%** urolithin A or **30%** urolithin A. The results were impressive. The higher urolithin A concentration mixture was most effective at inhibiting the number and size of colon-cancer stem cells and inhibiting the activity of *aldehyde dehydrogenase*, a marker of chemoresistance.⁷

This therapeutic approach is exciting because traditional therapies against cancer lack the ability to kill or stop the proliferation of cancerous *stem cells*. These new findings support the notion that a nutrient approach may prove valuable as an alternative treatment or preventive intervention for targeting these harmful cells.

Neuroprotective Effects

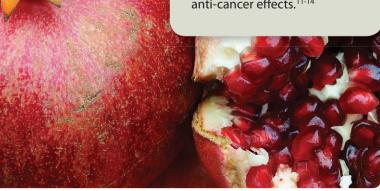
The connection between pomegranate and its neuroprotective effects against Alzheimer's disease has been well established in animal studies. However, the bioactive constituents for this action were unknown until now.

Alzheimer's disease is expected to affect over 115 million people worldwide by the year 2050.9 A group of researchers looked at a previous animal study that reported on the anti-Alzheimer's effects of pomegranate extract constituents.8

The team evaluated the ability of these components to cross the blood-brain barrier and found that a methylated form of urolithin A (mUA), derived from pomegranate, along with other urolithins were capable of doing so.

Urolithin A

Urolithin A is generated in the digestive tract as a natural metabolite of **punicalagins** and **ellagitannins**, polyphenol compounds found in pomegranates. Pomegranates have been identified as a potent weapon in the fight against aging due to their high content of ellagitannins. Ellagitannins are a natural substance with anti-inflammatory, anti-cancer effects. 11-14



Research Update

And, although more research is needed, the authors concluded that **urolithins** are the possible compounds responsible for the anti-Alzheimer's effects that include protection against neurotoxicity and β -amyloid fibrillation. These results are promising, and suggest the need for exploring other naturally-based dietary intervention strategies for preventing or slowing down the progression of Alzheimer's.

The results and data from these various studies further support the importance of polyphenol metabolite compounds like urolithin A from pomegranate and their role in the fight against colon cancer and neurodegenerative diseases.

Summary

The discovery of urolithin A, that results from the punical agins and ellagitannins compounds found in pomegranates, provides new opportunities to fight agerelated decline of mitochondrial function and the resulting frailty and loss of muscle.

By helping cells renew themselves and optimizing muscle performance, pomegranate extract and its newly identified metabolite, urolithin A—could prove successful.

Along with these findings, there is supportive evidence of the powerful effects that urolithin A has against Alzheimer's disease and cancer, offering yet another tool to fight against these devastating conditions that affect many aging individuals.

This nutritional approach opens up possibilities that traditional pharmaceutical approaches have never explored.

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

- Ryu D, Mouchiroud L, Andreux PA, et al. Urolithin A induces mitophagy and prolongs lifespan in C. elegans and increases muscle function in rodents. *Nat Med.* 2016;22(8):879-88.
- Fried LP, Tangen CM, Walston J, et al. Frailty in older adults: evidence for a phenotype. *J Gerontol A Biol Sci Med Sci*. 2001;56(3):M146-56.
- Perier C, Vila M. Mitochondrial biology and Parkinson's disease. *Cold Spring Harb Perspect Med.* 2012;2(2):a009332.
- 4. Available at: http://labiotech.eu/switzerland-unlocks-the-secret-to-pomegranates/. Accessed June 5, 2017.
- Patel BB, Majumdar AP. Synergistic role of curcumin with current therapeutics in colorectal cancer: minireview. *Nutr Cancer*. 2009;61(6):842-6.
- Subramaniam D, Ramalingam S, Houchen CW, et al. Cancer stem cells: a novel paradigm for cancer prevention and treatment. *Mini Rev Med Chem*. 2010;10(5):359-71.
- 7. Nunez-Sanchez MA, Karmokar A, Gonzalez-Sarrias A, et al. In vivo relevant mixed urolithins and ellagic acid inhibit phenotypic and molecular colon cancer

- stem cell features: A new potentiality for ellagitannin metabolites against cancer. *Food Chem Toxicol.* 2016; 92:8-16.
- 8. Yuan T, Ma H, Liu W, et al. Pomegranate's Neuroprotective Effects against Alzheimer's Disease Are Mediated by Urolithins, Its Ellagitannin-Gut Microbial Derived Metabolites. ACS Chem Neurosci. 2016;7(1):26-33.
- Available at: https://www.nia.nih.gov/ research/publication/longer-lives-anddisability/burden-dementia. Accessed June 5, 2017.
- Seeram NP, Henning SM, Zhang Y, et al. Pomegranate juice ellagitannin metabolites are present in human plasma and some persist in urine for up to 48 hours. J Nutr. 2006;136(10):2481-5.
- Mele L, Mena P, Piemontese A, et al. Antiatherogenic effects of ellagic acid and urolithins in vitro. *Arch Biochem Biophys*. 2016;599:42-50.
- Hollebeeck S, Winand J, Herent MF, et al. Anti-inflammatory effects of pomegranate (Punica granatum L.) husk ellagitannins in Caco-2 cells, an in vitro model of human intestine. Food Funct. 2012;3(8):875-85.
- 13. Ismail T, Calcabrini C, Diaz AR, et al. Ellagitannins in Cancer Chemoprevention and Therapy. *Toxins*. 2016;8(5):151.
- 14. Adams LS, Zhang Y, Seeram NP, et al. Pomegranate Ellagitannin-Derived Compounds Exhibit Anti-proliferative and Anti-aromatase Activity in Breast Cancer Cells In Vitro. Cancer prevention research (Philadelphia, Pa.). 2010;3(1):108-13.



TUMMY TROUBLES?

Get relief with **Digest RC**™, the European herbal formula for smoother digestion.



- Relieves fullness and bloating.
- Speeds digestion of fats and proteins.
- Prevents food stagnation in the digestive tract.

Digest RC™

Item #01358 • 30 tablets

	Retail Price	Your Price
1 box	\$19.95	\$14.96
4 boxes		\$12.75 each

For full product description and to order **Digest RC**™, call 1-800-544-4440 or visit www.Lifeextension.com



Polyphenols contained in berries are the focus of intense research into brain health.

Obtaining berries from food sources can be expensive, but standardized berry extracts are remarkably cost effective.

Here are two **berry extracts** used by **Life Extension**® consumers:



Blueberry Extract contains three different sources of blueberry in each 500 mg capsule, providing a broad-spectrum of active polyphenols.

The suggested supplemental dose for most individuals is <u>one</u> capsule daily.



Enhanced Berry Complete with Acai provides in each 700 mg capsule, extracts of 12 berries including blueberries and strawberries.

The suggested supplemental dose for most individuals is <u>one</u> capsule daily.



Blueberry Extract

Item #01214 • 60 vegetarian capsules

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

Enhanced Berry Complete with Acai

Item #01496 • 60 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$29	\$21.75
4 bottles		\$19.50 each

For full product description and to order Blueberry Extract or Enhanced Berry Complete with Acai, call 1-800-544-4440 or visit www.LifeExtension.com

Non-GMO Contains soybeans.

AuroraBlue® is a registered trademark of Denali BioTechnologies, Inc., AnthoComplete™ and VitaBerry® are trademarks of VDF Futureceuticals, Inc., used under license. POMELLA® Extract is covered under U.S. Patent 7,638,640 and POMELLA® is a registered trademark of Verdure Sciences, Inc.

Pomegranate Complete



THE NEXT-GENERATION POMEGRANATE FORMULA

Pomegranate Complete combines extracts from the whole fruit and flower along with pomegranate seed oil to support system-wide health. And it augments these polyphenols with newly discovered compounds from other parts of the pomegranate plant to help ease inflammation and to combat age-related metabolic changes.

Pomegranate Complete

Item #01953 • 30 softgels • Non-GMO

	Retail Price	Your Price
1 bottle	\$24	\$18
4 bottles		\$15.75 each



For full product description and to order **Pomegranate Complete**, call **1-800-544-4440** or visit **www.LifeExtension.com**

POMELLA® extract is covered under U.S. Patent 7,638,640 and POMELLA® is a registered trademark of Verdure Science, Inc.





Lactoferrin provides support for the body's immune system.1-2

In addition, new research shows lactoferrin increases tear production following cataract surgery by 95% and tear break-up time by 77%, which promotes eye protection.3

Contains milk.

Bioferrin[®] is a registered trademark of Glanbia.

References

1. Invest Ophthalmol Vis Sci. 2009 Apr;50(4):1636-43.

2. Curr Eye Res. 2013 Nov;38(11):1110-7.

3. J Clin Diagn Res. 2015 Oct;9(10):NC06-9.



Item #01681 • 60 capsules

	Retail Price	Your Price
1 bottle	\$44	\$33
4 bottles		\$30 each

For full product description and to order Lactoferrin, call 1-800-544-4440 or visit www.LifeExtension.com

BOOST **BRAIN** PERFORMANCE

Cognitex® with Pregnenolone & Brain Shield®

Cognitex® is designed to improve cerebral performance and supports brain and nervous system function.

Scientifically formulated **Cognitex**® contains validated ingredients shown at two weeks to improve:*

- Spatial short-term memory 42%
- Recall 15%
- Recognition 11%
- Attention 12%
- Visual learning 33%
- Activities of daily living by over 10%

Cognitex® with Pregnenolone & Brain Shield® (Gastrodin)

Item #01897 • 90 softgels

	Retail Price	Your Price
1 bottle	\$62	\$46.50
4 bottles		\$39.75 each

^{*} J Diet Suppl. 2011 Jun; 8(2):158-68

Non CMC

Caution: Do not take this product if you have breast cancer, prostate cancer, or other hormone-sensitive diseases. If you are taking anticoagulant or anti-platelet medications, or have a bleeding disorder, consult with your health care provider before taking this product.

Leucoselect® is a registered trademark of Indena S.p.A. Perluxan® is used with permission. Sensoril® is protected under US Patents Nos. 6,153,198 and 6,713,092 and is a registered trademark of Natreon, Inc. Sharp-PS® is a registered trademark of Enzymotec Ltd.

For full product description and to order

Cognitex® with Pregnenolone & Brain Shield®,

call 1-800-544-4440 or

visit www.LifeExtension.com







The best way to prevent cardiovascular disease (atherosclerosis, high blood pressure, heart attack, stroke, etc.) is with a healthy lifestyle—eat healthy foods, exercise, don't smoke, and don't gain weight.

For many people this lifestyle is too difficult or troublesome. Physicians attempting to encourage this lifestyle are often not successful and must resort to drugs.

Drugs are usually effective in reducing blood pressure. **Statin** drugs lower LDL and total cholesterol. One in four Americans age 45 and over takes a statin.¹

Blood plasma is **92%** water. Water and oil don't mix, so in order to transport fat in the bloodstream from the liver to body tissues, fats must be bound to **lipoproteins**. The two main forms of fat transported in the bloodstream by lipoproteins are **triglycerides** and **cholesterol**.

Two predominant lipoprotein particles are **LDL** (low density lipoprotein) and **HDL** (high density lipoprotein). Cholesterol attached to LDL (LDL cholesterol) is often called "bad cholesterol," because the LDL particle can deposit the cholesterol behind blood vessel walls, causing atherosclerosis.

HDL cholesterol is called "good cholesterol" because the HDL particle transports cholesterol back to the liver. This classification is misleading because both LDL and HDL can be either beneficial or harmful, depending on particle size, oxidation, and other factors.

Statin drugs reduce plasma levels of LDL cholesterol by inhibiting cholesterol formation in the liver.

PCSK9 is a naturally occurring human *enzyme* that causes LDL receptors to be degraded. If PCSK9 activity is blocked, more LDL receptors will be present on cells to remove LDL cholesterol from the blood. This results in <u>decreased</u> levels of blood LDL cholesterol. (PCSK9 stands for "Proprotein Convertase Subtilisin Kexin⁹.")

PCSK9 inhibitors are a new and expensive class of drugs that sharply reduce plasma LDL cholesterol.

Dr. Michael Ozner is a member of the **Life Extension Scientific Advisory Board**, a board certified cardiologist, and director of an annual symposium on preventing cardiovascular disease. This report concerns presentations made at his **February 2017** symposium.

Conference Overview

Michael Ozner, MD (medical director, Center for Prevention and Wellness, Baptist Health South Florida,



Miami, Florida) as director of this symposium gave an overview of topics related to cardiovascular disease prevention. Dr. Ozner emphasizes the importance of a healthy lifestyle, noting that even in patients with high genetic risk for coronary artery disease, a healthy lifestyle can reduce that risk by half.²

Dr. Ozner is a firm believer in the benefits of the diet eaten in the region of the Mediterranean sea (one of his books is titled *The Complete Mediterranean Diet*). He mentioned a study of over 100,000 health professionals which found that replacing **5**% of dietary saturated fat with either polyunsaturated fat, monounsaturated fat, or whole-grain carbohydrate reduced coronary heart disease risk by **25**%, **15**%, or **9**%, respectively.³ Aerobic exercise can reduce plasma triglycerides by up to **20**%.⁴

Concerning blood lipids, Dr. Ozner is very concerned about apolipoprotein B (**apoB**), which is the primary protein portion of all cholesterol particles other than HDL cholesterol.

High **apoB** is a better predictor of cardiovascular disease than high levels of LDL cholesterol.⁵ High apoB indicates numerous small, dense LDL cholesterol particles, the form of LDL which is most likely to be oxidized and cross blood vessel walls to cause atherosclerosis.⁶ High apoB in young adults predicts coronary artery calcification in midlife.⁷

Reduction of LDL cholesterol with statin drugs has been shown to reduce the incidence of heart attack and stroke.^{8,9} Using both a statin and anti-PCSK9 antibody is even more effective at lowering LDL cholesterol than statin alone.^{10,11}

Triglycerides and Cardiovascular Disease

Peter Libby, MD (cardiovascular specialist, Brigham and Women's Hospital, Boston, Massachusetts) spoke



about the increasing levels of plasma triglycerides in Americans. Roughly a quarter of American adults have excessively high levels of blood triglycerides. Triglyceride-laden lipoprotein is an even greater cause of coronary heart disease than LDL cholesterol. 13

Dr. Libby has noted that high HDL cholesterol tends to be associated

with low plasma triglycerides, and vice versa.¹⁴ But

when HDL is loaded with triglycerides, the HDL can become proinflammatory and cause atherosclerosis. ¹⁴ High levels of triglycerides on any of the lipoproteins cause inflammation. ¹⁵

Elevated triglycerides are associated with cardiovascular disease even in patients who have been successfully treated with statins.¹² When type II diabetics have high triglycerides, they show greater coronary artery calcification (atherosclerosis).¹⁶

Potential Benefit of HDL Cholesterol

Sergio Fazio, MD, PhD (director of the Center for Preventive Cardiology at Oregon Health & Science



University, Portland, Oregon) is concerned with the effects of HDL cholesterol on cardiovascular disease. Even among persons with low LDL cholesterol, those with the highest HDL cholesterol have less risk of cardiovascular disease than those with low HDL cholesterol. But while clinical trials have succeeded in raising HDL cholesterol in patients, this

did not reduce cardiovascular disease risk. 18,19

HDL cholesterol can become proinflammatory when LDL cholesterol is high.²⁰ Another reason why raising HDL cholesterol was not effective may have been that there are different forms of HDL cholesterol, with some forms being more protective than others.²¹ HDL subclasses appear to have different functions. For example, small HDL particles appear to have the capacity to remove cholesterol from atherosclerotic plaques.^{22,23}

A newly discovered compound known as **CSL112** is capable of making HDL cholesterol particles smaller, and thereby more efficient at removing cholesterol from atherosclerotic plaques.²⁴ CSL112 has not shown any harmful effects in clinical trials.



High Blood Pressure

William Cushman, MD (professor, Preventive Medicine, University of Tennessee, Memphis,



Tennessee) is concerned with high blood pressure as a cause of cardiovascular disease.

Nearly one third of American adults have **systolic blood pressure** (when the heart contracts) greater than **140 mmHg** and **diastolic pressure** (when the heart relaxes) greater than **90 mmHg**.²⁵

This high level (>140 mmHg systolic) of blood pressure raises the risk of coronary artery disease 44%, raises the risk of stroke 57%, raises the risk of heart failure 88%, and raises the risk of kidney failure 95%.²⁶ A clinical study investigated whether persons having a systolic pressure above 130 mmHg would benefit from therapy to reduce systolic pressure to less than 120 mmHg. Three years after systolic blood pressure was lowered, death rates dropped by about 25%.²⁷

Approximately a quarter of persons with high blood pressure (systolic pressure above **140 mmHg**) cannot reduce their blood pressure with three medications (**resistant hypertension**).²⁸ Most often, people with resistant hypertension are obese or elderly.

People with high blood pressure are more likely to have their blood pressure increased by salt consumption than people with normal blood pressure.²⁹ Eating foods higher in potassium, such as fruits and vegetables (rather than cereals and meats) can lower blood pressure.³⁰ The American Heart Association has estimated that increasing potassium consumption can decrease blood pressure and lengthen lifespan by several years.³⁰

Life Extension® has long advocated optimal blood pressure in most people to be **115 mmHg** systolic and **75 mmHg** diastolic. Newer studies corroborate the benefits of having lower normal blood pressure readings.

Stroke Risk

Ian del Conde Pozzi, MD (cardiologist, West Kendall Baptist Hospital, Miami, Florida) spoke of the risk of



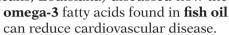
stroke. More than **60**% of patients with type II diabetes die of cardiovascular disease. But glucose control does not affect their risk of stroke.³¹ High blood pressure is the major risk factor for stroke.

A meta-analysis of 16 trials involving more than 70,000 patients showed that blood pressure-reducing

medications lowered the incidence of stroke by **22**%.³² Statin drugs were also shown beneficial. Clinical trials have shown that every **39 mg/dL** decrease in **LDL** cholesterol resulted in a greater than **21**% reduction of **stroke** risk.³³

Omega-3 Fatty Acids in Fish Oil

Carl Lavie, Jr., MD (cardiologist, Ochsner Medical Center, New Orleans, Louisiana) discussed how the





Fish oil supplements have been shown to reduce **inflammation** and blood vessel constriction,³⁴ while reducing irregular heartbeats (cardiac arrhythmias).³⁵

Approximately a third of Americans have excessively high blood triglycerides.³⁶ High blood tri-

glycerides are a strong predictor of residual risk of cardiovascular disease in patients receiving maximal doses of statins.³⁶

Omega-3 fatty acids reduce blood triglycerides significantly.³⁷ The minimal effective dose is more than **2,000 mg** of **EPA/DHA** from fish oil per day.³⁷

Eskimos show prolonged bleeding times with their dietary consumption of **20 grams** of omega-3 fatty acids per day, but Dr. Lavie has noted that doses of up to **7 grams** per day do not cause prolonged bleeding.³⁸ Consuming fish oil is safer than eating fish because toxic mercury attaches to fish meat, but is distilled out of quality omega-3 oil concentrates.³⁸ One study showed that **4,000 mg** of omega-3 fatty acids from fish oil reduce triglycerides by **45**% in patients with high triglycerides.³⁹

Lipoprotein(a)

Paul Ziajka, MD, PhD (clinical assistant professor, Florida University School of Medicine, Orlando,



Florida) spoke on the subject of lipoprotein(a) [**Lp(a)**], which is a highly atherosclerosis-causing particle attached to an LDL cholesterol particle. Patients with low LDL cholesterol nonetheless have a high cardiovascular disease risk if Lp(a) is high.⁴⁰

Lp(a) has less resistance to oxidation than plain LDL cholesterol.⁴¹

Niacin has been used to lower Lp(a).⁴² Combining niacin with statin was effective in lowering Lp(a) while at the same time increasing HDL cholesterol.⁴³

PCSK9 Action and Benefit

Peter Toth, MD, PhD (director, Preventive Cardiology, CGH Medical Center, Sterling, Illinois) discussed



PCSK9 inhibition. Experiments with mice have confirmed that PCSK9 inhibition lowers LDL cholesterol.⁴⁴ LDL cholesterol lowering with PCSK9 inhibition has also been shown in humans.⁴⁵

PCSK9 has been shown to lower not only plasma LDL cholesterol but also triglycerides⁴⁶ and Lp(a).⁴⁷

A clinical trial demonstrated that adding PCSK9 inhibition to statin therapy results in additional reduction of LDL cholesterol as well as additional reduction of cardiovascular death and disease.⁴⁸ PCSK9 inhibitors are much more expensive than statins, costing in excess of \$1,000 per month.⁴⁹

Coronary Artery Calcium

Khurram Nasir, MD (cardiologist, Baptist Health Medical Group, Miami Beach, Florida) is an advocate



of testing for **coronary artery calcium (CAC)**. This test uses high-speed radiological imaging to measure calcium in atherosclerotic plaques in the arteries of the heart. This helps determine the extent of coronary atherosclerosis.

Cardiovascular deaths and disease events are rare for persons who have no detectable CAC, mainly occurring

in diabetics and former smokers.⁵⁰ CAC is a more direct measurement of atherosclerosis than plasma LDL cholesterol or C-reactive protein, which are risk factors for atherosclerosis.⁵¹

More than one-quarter of American adults over age 40 take statin drugs.⁵² But Dr. Nasir has determined that people with no detectable CAC usually do not need to take a statin.⁵³

Bariatric Surgery

Anthony Gonzalez, MD (chief of surgery, Baptist Hospital of Miami, Miami, Florida) discussed the



benefits of gastric surgery for obese patients. Bariatric surgery reduces the size of the stomach, thereby reducing the amount of food a person can eat. Bariatric surgery substantially reduces cardiovascular death rates in obese patients⁵⁴ and substantially reduces symptoms of type II diabetes.⁵⁵

In 2010, **gastric bypass** was the most common form of bariatric surgery, but by 2013 **sleeve gastrectomy** had become the more common form.⁵⁶

Sleeve gastrectomy is less technically difficult to perform, but gastric bypass produces better results. According to one study, gastric bypass reduced body weight 23%, reduced triglycerides 40%, and reduced insulin use by 35%. For sleeve gastrectomy, the reductions were 19%, 29%, and 34%, respectively.⁵⁷ Earlier studies have shown similar results.⁵⁸ In all cases, bariatric surgery produced far better results than could be obtained by intensive treatment with medications.^{57,58}

Concluding Remarks

Cholesterol is not a toxic substance. On the contrary, cholesterol is an essential component of all cell membranes. Steroid hormones (testosterone, estrogen, etc.), cortisol, bile acids, and vitamin D are synthesized from cholesterol. Cholesterol is so essential for mental function that nearly one fourth of the body's cholesterol is in the brain, despite the fact that the brain accounts for only about **2**% of total body weight. Suicidal patients typically have lower plasma cholesterol than nonsuicidal patients.

LDL cholesterol becomes toxic when oxidized. Thus, smokers and other persons with high levels of oxidized LDL in their bloodstream are the main beneficiaries of cholesterol-lowering therapies. Considerable oxidation of LDL cholesterol occurs once it is behind blood vessels. Diabetes and chronic inflammation from other causes makes blood vessel walls more permeable to LDL and increases oxidative stress, thereby making LDL cholesterol more susceptible to oxidation. 63,64

LDL is beneficial when it is transported from the liver to body tissues, as opposed to being deposited into the endothelium. HDL is "good cholesterol" because it attaches to oxidized cholesterol in LDL for transport back to the liver to be detoxified.⁶⁵

Statins have cardiovascular benefits apart from LDL cholesterol lowering, including reduced inflammation.⁶⁶

One of the most common side effects associated with statins (**statin intolerance**), is muscle pain. Statin intolerance, real or imagined, is usually seen as subjective complaints rather than objective measurable quantities.⁶⁷ Especially in the elderly, muscle problems can be due to aging rather than statins. There are, nonetheless, objective problems seen with statin therapy, including an increased incidence of diabetes.⁶⁸

Statins reduce the body's synthesis of coenzyme Q10 and vitamin D. Statin intolerance could result from coenzyme Q10 depletion,⁶⁹ L-carnitine deficiency,⁷⁰ or vitamin D deficiency,⁷¹ all of which can be corrected with supplementation.



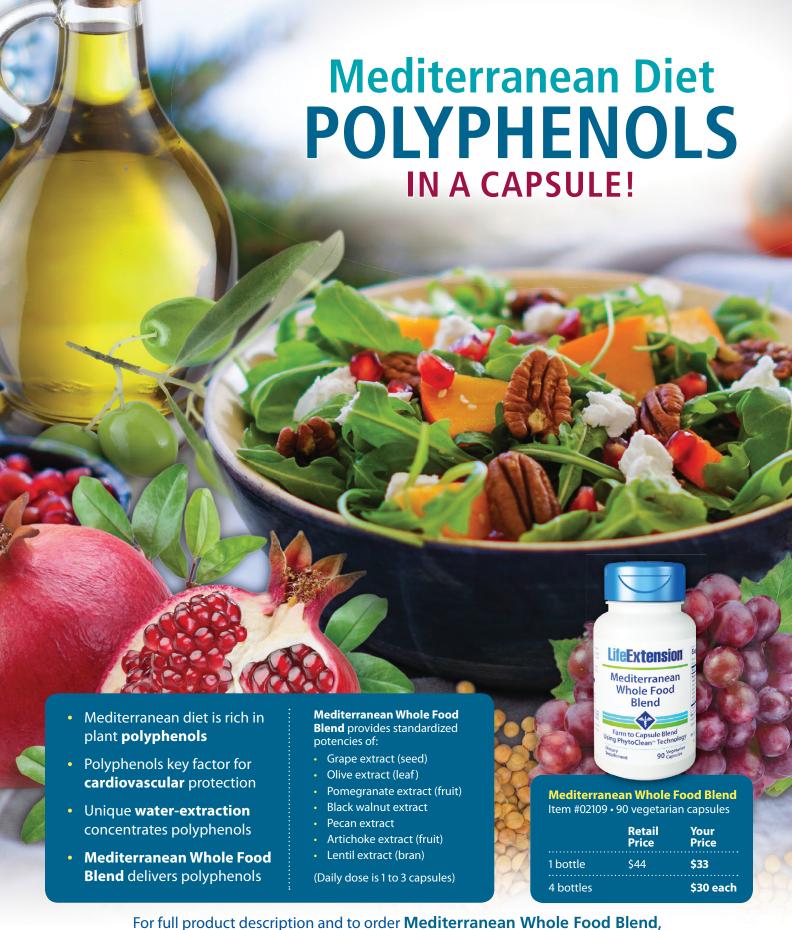
References

- 1. Available at: http://www.health.harvard.edu/blog/statin-use-isup-cholesterol-levels-are-down-are-americans-hearts-benefiting-201104151518. Accessed June 2, 2017.
- 2. Khera AV, Emdin CA, Drake I, et al. Genetic Risk, Adherence to a Healthy Lifestyle, and Coronary Disease. N Engl J Med. 2016;375(24):2349-58.
- Li Y, Hruby A, Bernstein AM, et al. Saturated Fats Compared With Unsaturated Fats and Sources of Carbohydrates in Relation to Risk of Coronary Heart Disease: A Prospective Cohort Study. J Am Coll Cardiol. 2015;66(14):1538-48.
- Watts GF, Ooi EM, Chan DC. Demystifying the management of hypertriglyceridaemia. Nat Rev Cardiol. 2013;10(11):648-61.
- Benn M, Nordestgaard BG, Jensen GB, et al. Improving prediction of ischemic cardiovascular disease in the general population using apolipoprotein B: the Copenhagen City Heart Study. Arterioscler Thromb Vasc Biol. 2007;27(3):661-70.
- Lamarche B, Tchernof A, Moorjani S, et al. Small, dense low-density lipoprotein particles as a predictor of the risk of ischemic heart disease in men. Prospective results from the Quebec Cardiovascular Study. Circulation. 1997;95(1):69-75.
- Wilkins JT, Li RC, Sniderman A, et al. Discordance Between Apolipoprotein B and LDL-Cholesterol in Young Adults Predicts Coronary Artery Calcification: The CARDIA Study. J Am Coll Cardiol. 2016;67(2):193-201.
- Cholesterol Treatment Trialists C, Baigent C, Blackwell L, et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26 randomised trials. Lancet. 2010;376(9753):1670-81.
- Toth PP, Thanassoulis G, Williams K, et al. The Risk-Benefit Paradigm vs the Causal Exposure Paradigm: LDL as a primary cause of vascular disease. J Clin Lipidol. 2014;8(6):594-605
- 10. Puri R, Nissen SE, Somaratne R, et al. Impact of PCSK9 inhibition on coronary atheroma progression: Rationale and design of Global Assessment of Plaque Regression with a PCSK9 Antibody as Measured by Intravascular Ultrasound (GLAGOV). Am Heart J. 2016:176:83-92
- 11. Gouni-Berthold I, Berthold HK. PCSK9 antibodies for the treatment of hypercholesterolemia. Nutrients. 2014;6(12):5517-33.
- 12. Toth PP. Triglyceride-rich lipoproteins as a causal factor for cardiovascular disease. Vasc Health Risk Manag. 2016;12:171-83.
- 13. Rosenson RS, Davidson MH, Hirsh BJ, et al. Genetics and causality of triglyceride-rich lipoproteins in atherosclerotic cardiovascular disease. J Am Coll Cardiol. 2014;64(23):2525-40.
- 14. Libby P. Triglycerides on the rise: should we swap seats on the seesaw? Eur Heart J. 2015;36(13):774-6.

- 15. Nordestgaard BG. Triglyceride-Rich Lipoproteins and Atherosclerotic Cardiovascular Disease: New Insights From Epidemiology. Genetics, and Biology. Circ Res. 2016;118(4):547-63.
- 16. Qamar A, Khetarpal SA, Khera AV, et al. Plasma apolipoprotein C-III levels, triglycerides, and coronary artery calcification in type 2 diabetics. Arterioscler Thromb Vasc Biol. 2015;35(8):1880-8.
- 17. Barter P, Gotto AM, LaRosa JC, et al. HDL cholesterol, very low levels of LDL cholesterol, and cardiovascular events. N Engl J Med. 2007:357(13):1301-10
- 18. Schwartz GG, Olsson AG, Abt M, et al. Effects of dalcetrapib in patients with a recent acute coronary syndrome. N Engl J Med. 2012;367(22):2089-99.
- 19. Khera AV, Rader DJ. Cholesterol efflux capacity: full steam ahead or a bump in the road? Arterioscler Thromb Vasc Biol. 2013;33(7):1449-51.
- 20. Aryan Z, Noshad S, Afarideh M, et al. Comment on Sharif et al. HDL Cholesterol as a Residual Risk Factor for Vascular Events and All-Cause Mortality in Patients With Type 2 Diabetes. Diabetes Care 2016;39:1424-1430. Diabetes Care. 2016;39(10):e189.
- 21. Williams PT, Feldman DE. Prospective study of coronary heart disease vs. HDL2, HDL3, and other lipoproteins in Gofman's Livermore Cohort. Atherosclerosis. 2011;214(1):196-202.
- 22. Anastasius M, Kockx M, Jessup W, et al. Cholesterol efflux capacity: An introduction for clinicians. Am Heart J. 2016;180:54-63.
- 23. Fazio S, Pamir N. HDL Particle Size and Functional Heterogeneity. Circ Res. 2016;119(6):704-7.
- 24. Didichenko SA, Navdaev AV, Cukier AM, et al. Enhanced HDL Functionality in Small HDL Species Produced Upon Remodeling of HDL by Reconstituted HDL, CSL112: Effects on Cholesterol Efflux, Anti-Inflammatory and Antioxidative Activity. Circ Res. 2016;119(6):751-63.
- 25. Cushman WC, Ford CE, Einhorn PT, et al. Blood pressure control by drug group in the Antihypertensive and Lipid-Lowering Treatment to Prevent Heart Attack Trial (ALLHAT). J Clin Hypertens (Greenwich). 2008;10(10):751-60.
- 26. Muntner P, Davis BR, Cushman WC, et al. Treatment-resistant hypertension and the incidence of cardiovascular disease and end-stage renal disease: results from the Antihypertensive and Lipid-Lowering Treatment to Prevent Heart Attack Trial (ALLHAT). Hypertension. 2014;64(5):1012-21.
- 27. Group SR, Wright JT, Jr., Williamson JD, et al. A Randomized Trial of Intensive versus Standard Blood-Pressure Control. N Engl J Med. 2015;373(22):2103-16.
- 28. Calhoun DA, Jones D, Textor S, et al. Resistant hypertension: diagnosis, evaluation, and treatment. A scientific statement from the American Heart Association Professional Education Committee of the Council for High Blood Pressure Research. Hypertension. 2008;51(6):1403-19.

- 29. McCallum L, Lip S, Padmanabhan S. The hidden hand of chloride in hypertension. Pflugers Arch. 2015;467(3):595-603.
- Stone MS, Martyn L, Weaver CM. Potassium Intake, Bioavailability, Hypertension, and Glucose Control. Nutrients. 2016;8(7).
- 31. Duckworth W, Abraira C, Moritz T, et al. Glucose control and vascular complications in veterans with type 2 diabetes. N Engl J Med. 2009;360(2):129-39.
- 32. Sipahi I, Swaminathan A, Natesan V, et al. Effect of antihypertensive therapy on incident stroke in cohorts with prehypertensive blood pressure levels: a meta-analysis of randomized controlled trials. Stroke. 2012:43(2):432-40.
- 33. Amarenco P, Labreuche J. Lipid management in the prevention of stroke: review and updated meta-analysis of statins for stroke prevention. Lancet Neurol. 2009;8(5):453-63.
- 34. Mozaffarian D, Bryson CL, Lemaitre RN, et al. Fish intake and risk of incident heart failure. J Am Coll Cardiol. 2005;45(12):2015-21.
- 35. Yokoyama M, Origasa H, Matsuzaki M, et al. Effects of eicosapentaenoic acid on major coronary events in hypercholesterolaemic patients (JELIS): a randomised open-label, blinded endpoint analysis. Lancet. 2007:369(9567):1090-8.
- 36. Benes LB, Bassi NS, Davidson MH. Omega-3 carboxylic acids monotherapy and combination with statins in the management of dyslipidemia. Vasc Health Risk Manag. 2016;12:481-90.
- 37. Shearer GC, Savinova OV, Harris WS. Fish oil -- how does it reduce plasma triglycerides? Biochim Biophys Acta. 2012;1821(5):843-51.
- 38. Lavie CJ, Milani RV, Mehra MR, et al. Omega-3 polyunsaturated fatty acids and cardiovascular diseases. J Am Coll Cardiol. 2009:54(7):585-94.
- 39. Jacobson TA. Role of n-3 fatty acids in the treatment of hypertriglyceridemia and cardiovascular disease. Am J Clin Nutr. 2008:87(6):1981s-90s.
- 40. Konishi H, Miyauchi K, Kasai T, et al. Impact of lipoprotein(a) as residual risk on long-term outcomes in patients after percutaneous coronary intervention. Am J Cardiol. 2015:115(2):157-60.
- 41. Cantin B, Gagnon F, Moorjani S, et al. Is lipoprotein(a) an independent risk factor for ischemic heart disease in men? The Quebec Cardiovascular Study. J Am Coll Cardiol. 1998;31(3):519-25.
- 42. Nordestgaard BG, Chapman MJ, Ray K, et al. Lipoprotein(a) as a cardiovascular risk factor: current status. Eur Heart J. 2010;31(23):2844-53.
- 43. Levy DR, Pearson TA. Combination niacin and statin therapy in primary and secondary prevention of cardiovascular disease. Clin Cardiol. 2005:28(7):317-20.
- 44. Abifadel M, Rabes JP, Devillers M, et al. Mutations and polymorphisms in the proprotein convertase subtilisin kexin 9 (PCSK9) gene in cholesterol metabolism and disease. Hum Mutat. 2009:30(4):520-9
- 45. Steinberg D, Witztum JL. Inhibition of PCSK9: a powerful weapon for achieving ideal LDL cholesterol levels. Proc Natl Acad Sci USA. 2009;106(24):9546-7.
- 46. Seidah NG. New developments in proprotein convertase subtilisinkexin 9's biology and clinical implications. Curr Opin Lipidol. 2016;27(3):274-81.
- 47. Banerjee Y, Santos RD, Al-Rasadi K, et al. Targeting PCSK9 for therapeutic gains: Have we addressed all the concerns? Atherosclerosis. 2016:248:62-75.
- 48. Sabatine MS, Giugliano RP, Keech AC, et al. Evolocumab and Clinical Outcomes in Patients with Cardiovascular Disease, N Engl J Med. 2017;376(18):1713-22.
- 49. Available at: http://www.medscape.com/viewarticle/867527. Accessed June 23, 2017.
- 50. Budoff MJ, McClelland RL, Nasir K, et al. Cardiovascular events with absent or minimal coronary calcification: the Multi-Ethnic Study of Atherosclerosis (MESA). Am Heart J. 2009;158(4):554-61.
- 51. Blaha MJ, Budoff MJ, DeFilippis AP, et al. Associations between C-reactive protein, coronary artery calcium, and cardiovascular events: implications for the JUPITER population from MESA, a population-based cohort study. Lancet. 2011;378(9792):684-92.
- 52. Salami JA, Warraich H, Valero-Elizondo J, et al. National Trends in Statin Use and Expenditures in the US Adult Population From 2002 to 2013: Insights From the Medical Expenditure Panel Survey. JAMA Cardiol. 2017;2(1):56-65.

- 53. Nasir K, Bittencourt MS, Blaha MJ, et al. Implications of Coronary Artery Calcium Testing Among Statin Candidates According to American College of Cardiology/American Heart Association Cholesterol Management Guidelines: MESA (Multi-Ethnic Study of Atherosclerosis). J Am Coll Cardiol. 2015;66(15):1657-68.
- 54. Sjostrom L, Peltonen M, Jacobson P, et al. Bariatric surgery and long-term cardiovascular events. JAMA. 2012;307(1):56-65.
- 55. Carlsson LM, Peltonen M, Ahlin S, et al. Bariatric surgery and prevention of type 2 diabetes in Swedish obese subjects. N Engl J Med. 2012;367(8):695-704.
- 56. Spaniolas K, Kasten KR, Brinkley J, et al. The Changing Bariatric Surgery Landscape in the USA. Obes Surg. 2015;25(8):1544-6.
- 57. Schauer PR, Bhatt DL, Kirwan JP, et al. Bariatric Surgery versus Intensive Medical Therapy for Diabetes - 5-Year Outcomes. N Engl J Med. 2017;376(7):641-51.
- 58. Schauer PR, Kashyap SR, Wolski K, et al. Bariatric surgery versus intensive medical therapy in obese patients with diabetes. N Engl J Med. 2012:366(17):1567-76.
- 59. Dietschy JM, Turley SD. Thematic review series: brain Lipids. Cholesterol metabolism in the central nervous system during early development and in the mature animal. J Lipid Res. 2004;45(8):1375-
- 60. Pfrieger FW. Role of cholesterol in synapse formation and function. Biochim Biophys Acta. 2003;1610(2):271-80.
- 61. Wu S, Ding Y, Wu F, et al. Serum lipid levels and suicidality: a meta-analysis of 65 epidemiological studies. J Psychiatry Neurosci. 2016:41(1):56-69.
- 62. Nishi K, Itabe H, Uno M, et al. Oxidized LDL in carotid plaques and plasma associates with plaque instability. Arterioscler Thromb Vasc Biol. 2002;22(10):1649-54.
- 63. Obradovic MM, Trpkovic A, Bajic V, et al. Interrelatedness between C-reactive protein and oxidized low-density lipoprotein. Clin Chem Lab Med. 2015;53(1):29-34.
- 64. Tangvarasittichai S. Oxidative stress, insulin resistance, dyslipidemia and type 2 diabetes mellitus. World J Diabetes. 2015;6(3):456-
- 65. Navab M, Ananthramaiah GM, Reddy ST, et al. The oxidation hypothesis of atherogenesis: the role of oxidized phospholipids and HDL. J Lipid Res. 2004;45(6):993-1007.
- 66. Wang JC, Bennett M. Aging and atherosclerosis: mechanisms, functional consequences, and potential therapeutics for cellular senescence. Circ Res. 2012;111(2):245-59.
- 67. Stulc T, Ceska R, Gotto AM, Jr. Statin Intolerance: the Clinician's Perspective. Curr Atheroscler Rep. 2015;17(12):69.
- 68. Ganda OP. Statin-induced diabetes: incidence, mechanisms, and implications. F1000Res. 2016;5.
- 69. Potgieter M, Pretorius E, Pepper MS. Primary and secondary coenzyme Q10 deficiency: the role of therapeutic supplementation. Nutr Rev. 2013:71(3):180-8.
- 70. Spence JD, Dresser GK. Overcoming Challenges With Statin Therapy. J Am Heart Assoc. 2016;5(1).
- 71. Saxon DR, Eckel RH. Statin Intolerance: A Literature Review and Management Strategies. Prog Cardiovasc Dis. 2016;59(2):153-64.



For full product description and to order **Mediterranean Whole Food Blend**, call **1-800-544-4440** or visit **www.LifeExtension.com**

The PhytoClean™ Method is a patented bioactive extraction method. **CAUTION:** If you have gallstones or a biliary tract obstruction, do not use this product. Do not take if pregnant or lactating. Contains tree nuts (walnut, pecan). Keep out of reach of children. Do not exceed recommended dose. **Non-GMO**



Taurine, a free amino acid, is "one of the most essential substances in the body." 1 But as we age, taurine levels decline.

Research has found that taurine can promote **new cell** formation in the area of the brain associated with learning and memory.

Taurine also enhances **neurites**, tiny projections that help brain cells communicate with each other.

Past research has also shown the ability of taurine to maintain and support: 2-8

- · Cardiovascular health
- Insulin sensitivity
- · Modulation of the immune system
- Regulation of the central nervous system
- Liver function
- Eye health
- Hearing function

Item #01827 • 90 vegetarian capsules

	Retail Price	Your Price
1 bottle	\$13	\$9.75
4 bottles		\$9 each

Non-GMO

For full product description and to order Taurine, call 1-800-544-4440 or visit www.LifeExtension.com

- eferences
 Mol Vis. 2012;18;2673-86.
 Exp Clin Cardiol. 2008 Summer;13(2):57-65.
 Exp Mol Med. 2012 Nov 30;44(11):665-73.
 Curr Opin Clin Nutr Metab Care. 2006 Nove;9(6):728-33.
 J Biomed Sci. 2010 Aug 24;17 Suppl1:51.
 Amino Acids. 2008 Aug;35(2):469-73.
 Amino Acids. 2012 Nov;43(5):1979-93.
 Neurosci Lett. 2006 May 15;399(1-2):23-6.

BY GARRY MESSICK

Celery is related to parsley and fennel, a member of the Umbelliferae family of plants. Mankind has cultivated it as a vegetable for thousands of years.

Let's consider a few of the numerous if often overlooked health benefits contained in this popular, fibrous vegetable.

Fiber

At about 1.6 grams per cup, celery is high in fiber, which makes it beneficial for helping to fight everything from diabetes, heart disease, and high cholesterol to colon cancer and constipation.1

Anti-inflammatory

Celery contains beneficial phytonutrients such as the flavonols quercetin and kaempferol, flavones such as luteolin, and phenolic acids. These antioxidants are known for their anti-inflammatory properties.² A study has shown that celery helps inhibit the activity of two proteins linked to inflammation—nuclear factor-kappa B (NF-kB) and tumor necrosis factor alpha (TNF-alpha).3

Minerals

Celery is rich with a number of important minerals. These include iron, zinc, copper, magnesium, calcium, and selenium, but chiefly potassium, which helps reduce the risk of heart disease and support cellular function in muscles.1

It is best to choose organic celery whenever possible as commercially grown celery has been exposed to a great deal of pesticides.4,5

- 1. Available at: http://woman.thenest.com/ health-benefits-celery-2024.html. Accessed June 6, 2017.
- 2. Available at: http://www.livescience. com/50640-celery-nutrition.html. Accessed June 6, 2017.
- 3. Mol Nutr Food Res. 2012;56(4):558-69.
- 4. Available at: https://www.ewg.org/foodnews/ summary.php. Accessed June 6, 2017.
- 5. Available at: http://www.cnn.com/2010/ HEALTH/06/01/dirty.dozen.produce.pesticide/ index.html. Accessed June 6, 2017.

Not Eating Enough Veggies? No Problem!

Get Protective Benefits Of

Cruciferous Vegetables

with Apigenin

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

Triple Action Cruciferous Vegetable Extract combines vital plant **extracts** that have been shown to protect cellular DNA.

The formula provides optimal potencies of cruciferous extracts like **I3C** (*Indole-3-carbinol*) and **DIM** (*di-indolyl-methane*) to favorably modulate estrogen metabolism,¹⁻⁵ along with **apigenin**.

For full product description and to order
Triple Action Cruciferous
Vegetable Extract, call 1-800-544-4440
or visit www.LifeExtension.com

- 1. Biochem Pharm. 2002, 64;393-404.
- 2. Toxicol Appl Pharm. 2001 Jul 15;174(2):146-52.
- 3. In Vivo. 2006 Mar-Apr;20(2):221-8.
- 4. Cancer Detect Prevent. 2004;28:72-9.
- 5. *Mol Carcinog*. 2012 Mar;51(3):244-56.



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





BY JON VANZILE

HUMBERTO FASANO

A Heart Transplant Success Story

In 2001, Humberto Fasano received a staggering diagnosis: he had severe cardiomyopathy and congestive heart failure (CHF).

With these conditions, the heart muscle becomes flabby and stretched out. As a result, its pumping ability is weakened, and not enough oxygenated blood is circulated to the body, causing symptoms like exhaustion and arrhythmia. The condition is considered degenerative and incurable, meaning that once someone is diagnosed with congestive heart failure, there is no way to restore their heart function back to its normal level.

Doctors measure CHF by tracking the size of the left ventricle and by measuring how much blood it expells with every contraction, a measurement known as the "ejection fraction." In general, doctors consider an ejection fraction above **55%** to be normal.

Fasano had an ejection fraction of just **14%**. And at 10.5 centimeters, his left ventricle was the largest ever recorded at the University of Miami Health System.



Humberto Fasano and his wife, Maria.

Wellness Profile

Based on this grim diagnosis, Fasano needed to be on the heart transplant list. Doctors told him that **50%** of CHF patients usually die within five years of being diag-

But Fasano, who says he always maintains an optimistic and hopeful attitude, wasn't buying it. Instead of giving in to his diagnosis, he decided he would fight back and do everything he could to survive. Today, more than 15 years later, Fasano is thriving—and he's always happy to tell the story of how he managed to outlast a chronic, seemingly hopeless diagnosis with the help of first-rate cardiac care and Life Extension®.

"I was never in fear or depressed," Fasano says. "'Give up' is not in my dictionary, period."

The Road Back

Initially, Fasano experienced symptoms such as shortness of breath and fatigue while performing everyday activities such as climbing stairs. In the short term, to help relieve his symptoms and support his weak heart, his doctors recommended immediate placement of a left-sided pacemaker/ defibrillator. This device regulated his heart rhythm and, if necessary, shocked it back into a normal rhythm if something went wrong. Two years later, the device was upgraded to a biventricular pacemaker/defibrillator, which provides more comprehensive support to keep a normal heart rhythm.

While the devices went to work and Fasano adapted to an aggressive program of prescription drugs to reduce his symptoms, he launched an all-out effort to find relief. In 2004, "by the grace of God," he came across an Internet forum where people with congestive heart failure were singing the praises of an orthomolecular therapist in Holland named Corrij Kooij. According to her patients, Kooij had developed a special program of supplements that was able to increase their ejection fraction by up to 15%.

"I immediately wrote Corrij and asked if she could help me," Fasano says. "She accepted and requested I fill out a questionnaire, in addition to sending her my last two blood lab reports. She performed this service at zero cost to me."

After reviewing his case, Kooii recommended a program of nine highly-targeted nutrients and supplements. These supplements (see sidebar) were designed to make his heart stronger by increasing intracardiac energy and to help his heart function more efficiently. The list included stalwarts like coenzyme Q10, magnesium, potassium, R-lipoic acid, and vitamins C and E. Each of the recommendations was backed up by extensive research showing a positive effect on cardiac function.

Naturally, Fasano wanted to make sure he was getting the best supplements possible. At the time, he was already taking a few supplements from **Life Extension**, so he wrote Kooij back and mentioned that he was taking Life Extension

products already and did she have a particular brand she recommended. In his letter, he remarked that he would go anywhere in the world to obtain the very best.

"If she said I needed a Japanese supplement, I was prepared to go to Japan to buy it. But she immediately wrote back and said, 'Humberto, here in Holland my husband and I take Life Extension," he recalls. "I found all nine products within Life Extension's line-up."

This recommendation would end up launching a long-standing relationship between Fasano and Life Extension—but first he had to make sure his health didn't deteriorate.

The Transplant

In 2014, 13 years after his diagnosis, Fasano had already outlived all expectations, but he was still looking for every advantage possible. That year, he signed up for an experimental stem-cell study at the University of Miami.

Unfortunately, the treatment didn't have the effect he'd hoped for. Later that year, with his condition worsening, he had to pull himself from the program. That autumn, he had two serious emergencies, one in October and one in Novemberthen three more in January 2015.



By this time, it had become obvious that Fasano would have to take the next step. He would need a heart transplant. He applied to the Mavo Clinic Transplant Program and was accepted. While he was waiting, he spent several months taking a class of drugs called inotropes and, in summer 2015, he received a left-ventricular assist device. or LVAD. This specialized device acts as a kind of mechanical heart. LVADs were originally designed as a "bridge to transplant," for shortterm use, but recent advances in technology have stretched the horizon of time people can survive with an LVAD device. This was excellent news for Fasano and gave him time while the search for a donor heart began.

In January 2015, Fasano and his wife moved to Jacksonville, Florida, where the Mayo Clinic transplant hospital was located, to wait for his donor heart. With his typical optimism, Fasano loved Jacksonville.

"I spent months within 15 minutes of the heart transplant center," Fasano says. "And I'll tell you, we had a great time. I didn't have any dietary restrictions anymore, so I told my wife, 'Let's go find all the good restaurants!' So we did, and there are some great restaurants in Jacksonville!"

But that doesn't mean it was easy. While they waited, Fasano's doctors continued to refine his medication program, and his symptoms posed a daily challenge. He was tired more often and dealt with the side effects of his medications.

"I was limited physically," he said. "I couldn't run or go upstairs. I couldn't do a lot of the things I used to do. But emotionally, I never had a problem."

Finally, that autumn, Fasano got the news—a suitable donor heart had been found. On November 18, he received a new heart. It was a huge step forward, and it began his journey back. Eighteen months after his transplant, Fasano felt much better and has been able to participate more fully in life.

"I have been able to go back to most of my normal activities, including work, walking every day, enjoying my family, and helping others as much as I can," he says. "I also participate in a foundation in Panama that raises funds for people who need a transplant."

Fasano's Lasting Connection with Life Extension

More than anything else, Fasano is deeply grateful for all the help he's received.

"I owe my new life to God and his team first, including my caregiver 24/7 for sixteen continuous months: my wife Maria, who never had one complaint and never got tired. I still cannot believe her strength and persistence. My brother Max was also 24/7 dedicated to taking care of me uninterrupted. I'm extremely thankful for my doctors, my nurses and the whole medical team at Mayo Clinic and University of Miami. as well as the prayers of my family and friends. I'm also grateful to Corrij, my angel, and to Life Extension for the supplements I took for more than 10 years to control my illness."

In fact, Fasano was so impressed with his results that shortly after he started his supplement regimen, he reached out to **Life Extension** to set up an appointment with the international team. Soon after, he became a general distributor for the company in Panama—a position he continues to hold.

"When I started this journey, I knew very little about vitamins

Humberto Fasano's Heart Failure Protocol

After Humberto Fasano reached out to Corrij Kooij, she sent him a supplement protocol designed to provide extra energy to a stressed heart muscle. The list of supplements Kooij recommended includes:

- MSM: 1,000 mg, 3x daily
- Super Ubiquinol CoQ10:
 100 mg, 2x daily with meals
- Chromium polynicotinate: 200 mcg, 2x twice daily with meals
- Optimized Carnitine with GlycoCarn®: 2 capsules twice daily, a total of 2,800 mg, half-hour before or two hours after meals
- · Magnesium: 2x daily
- R-Lipoic acid: 150 mg, 2x daily
- Potassium: **750 mg**, 1x daily
- Vitamin C: 1,000 mg, 3x daily
- Vitamin E: 400 IU, 1x daily

Now, after the transplant, Fasano has also added the following supplements to his regimen:

- Super Bio-Curcumin
- Bone Restore
- Arthromax

and supplements," he says. "I was a civil engineer who worked in construction and project management. Distributing **Life Extension** products was something I started on the side, but now it's grown, and I'm doing great."

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

EXTEND-RELEASE

MAGNESIUM

WHEN YOU NEED IT

Unique delivery system provides **immediate** <u>and</u> **extended release magnesium** for full-body coverage of this essential mineral.



Non-GMO

Retail Your Price Price

1 bottle \$13 **\$9.75** 4 bottles **\$8.75 each**

Item # 02107 • 60 vegetarian capsules



For full product description and to order **Extend-Release Magnesium**, call **1-800-544-4440** or visit **www.LifeExtension.com**

ZümXR° is a registered trademark and protected by patents. See www.ZümXR.com



Most people obtain **alpha-tocopherol** in their multivitamin, but miss out on critical **gamma-tocopherol**.

Gamma E Mixed Tocopherols provides a full spectrum of natural vitamin E.



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

Item# 02075 • 60 softgels

Non-GMO

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

For full product description and to order Gamma E Mixed Tocopherol with Sesame Lignans, call 1-800-544-4440 or visit www.LifeExtension.com

ENHANCE YOUR IMMUNE SYSTEM WITH THE HIGHEST POTENCY AHCC® PRODUCT



Item #24404 **Retail Price \$84.95 Your Price \$63.71**

Quality of Life's AHCC products stand as the most clinically-researched immune support supplements available.* Backed by over 20 human clinical studies conducted at some of the finest research institutions worldwide, including lyy League universities and major health centers, AHCC provides you with immune support in times of need, and in times of maintenance.*

Kinoko Platinum is the ultimate choice as it delivers 750 mg of AHCC per vegicap — the highest potency available anywhere — making it easier to achieve the 3-gram amount recommended for advanced immune system support.*

FOR FULL PRODUCT DESCRIPTION AND TO ORDER KINOKO® PLATINUM AHCC®,
PLEASE CALL 1-800-544-4440 OR VISIT WWW.LIFEEXTENSION.COM



QUALITY OF LIFE | 877-937-2422 | www.QualityOfLife.net

■ Facebook.com/QualityofLifeLabs ■ @QOLsupplements

Quality of Life is proud to have taken the Natural Products Foundation's "Truth in Advertising Pledge," a formal commitment to disserninating only truthful, non-misleading, and substantiated information.



THE PROBIOTIC FOR WOMEN

Clinically Documented Probiotic Strains That Promote Healthy Vaginal Microflora and Urinary Tract Health*

Jarro-Dophilus® Women contains the four predominant *Lactobacilli* strains of the healthy vaginal tract.*

All four strains were isolated from the vaginal tracts of healthy pregnant women and have been clinically tested for efficacy L. crispatus LbV 88

L. jensenii LbV 116

L. gasseri LbV 150N

L. rhamnosus LbV 96

in helping to maintain protective, healthy vaginal microflora and urinary tract health.*

When it comes to choosing effective probiotics, clinically documented strains matter.™

Choose science.

Choose Jarro-Dophilus® probiotics.

Jarro-Dophilus® Women 5 Billion Per Capsule 30 Veggie Caps

Item # **52142** Retail Price **\$27.95** Your Price **\$20.96** For full product description or to order Jarro-Dophilus® Women call 1-800-544-4440 or visit www.LifeExtension.com



Jarro-Dophilus® Women contains the clinically tested Astarte® strains which are protected by U.S. Patent 8,846,027 and European Patent 2,509,610. Astarte® is owned by HSO Health Care GmbH, Vienna, Austria, and licensed in the U.S. to Jarrow Formulas, Inc. Other international patents pending.

NEW FOOD-GRADE ENTERIC COATING



ROOM TEMPERATURE STABLE™

Jarro-Dophilus



PROMOTES HEALTHY VAGINAL MICROFLORA*
PROMOTES URINARY TRACT HEALTH*

5 BILLION PER CAPSULE

4 CLINICALLY DOCUMENTED VAGINAL STRAINS

30 ENTERIC COATED VEGGIE CAPS PROBIOTIC SUPPLEMENT

*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.



FIVE EASY STEPS FOR ORDERING BLOOD TESTS:

 Call 1-800-208-3444 to discuss and place your order with one of our knowledgeable Wellness Specialists.
 Online orders can also be placed at

www.LifeExtension.com/labtesting

- 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.
- Your blood test results will be mailed, emailed, or faxed directly to you by Life Extension.
- 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

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.

LifeExtension

Blood Testing The Ultimate Information

YOUR

\$35

\$199

\$198

\$198

\$375

\$299

\$149

PSA (PROSTATE SPECIFIC ANTIGEN) (LC010322)

Screening for Prostate Cancer

SALE \$23.25

CBC/CHEMISTRY PROFILE (LC381822) includes:

Blood Sugar: Glucose

Lipid Profile:

Total cholesterol • Triglycerides
HDL cholesterol • LDL cholesterol (calc.)
VLDL cholesterol (calc.)
Total cholesterol/HDL ratio
Estimated Coronary Heart Disease risk

Kidney Function:

Uric acid • BUN (blood urea nitrogen) Creatinine • BUN/creatinine ratio eGFR (estimated glomerular filtration rate)

Liver Function:

Alkaline phosphatase • LDH (lactate dehydrogenase) AST (aspartate aminotransferase) ALT (alanine transaminase) Total protein • Albumin • Globulin Albumin/globulin ratio • Bilirubin

DH (lactate Complete Blood Count:

Red blood cell count • Hemoglobin
Hematocrit • MCV (mean corpuscular volume)
MCH (mean corpuscular hemoglobin)
MCHC (mean corpuscular hemoglobin
concentration)
RDW (red blood cell distribution)
White blood cell count
Immune Cell Differentiation Count
Platelet count

Electrolytes and Minerals:

Sodium • Potassium • Chloride Calcium • Phosphorus • Iron

NEUROTRANSMITTER BASIC PANEL**(LC100058)

Serotonin, Dopamine, Epinephrine, Norepinephrine, GABA, Glutamate. Alternations in these six neurotransmitters play a significant role in contributing to symptoms such as cognitive disorders, depression, anxiety, diminished drive, fatigue and sleep difficulties, cravings, addictions, pain and more! Not available in NY.

→ FOOD SAFE ALLERGY TEST - BASIC** (LCM73001)

This test measures delayed (lgG) food allergies for 95 common foods.

FOOD SAFE ALLERGY TEST — EXTENDED** (LCM73002)

This test measures delayed (IgG) food allergies to an additional 95 foods.

FOOD SAFE ALLERGY TEST — COMBO** (LCM73003)

This test measures delayed (IgG) food allergies to all 190 foods found in our Basic and Extended panels.

NEW GENETIC TESTING

DNA GENETIC CANCER RISK PROFILE**(LC100057)

With only a saliva sample, you can identify your risk for 25 hereditary cancers by analyzing 98 genes from your DNA including the well-known BRCA1, BRCA2, TP53, and APC. Not available in FL, NY, and RI.

APOE GENETIC TEST FOR ALZHEIMER'S AND CARDIAC RISK **(LC100059)

Apolipoprotein E (ApoE) is an important regulator of cholesterol and triglycerides levels in your blood and supports lipid transport and injury repair in your brain. Genetically, E4 is the strongest risk factor for developing Late Onset Alzheimer's disease. According to the National Institute of Health, inheriting a single copy of ApoE4 increases the risk of Alzheimer's disease by about three-fold. Inheriting two copies increases the risk by about 12-fold. In fact, almost 40% of AD patients have inherited an E4 allele.

In the cardiovascular system ApoE is involved in the transportation of fat molecules into your cells. E4 is associated with increased levels of cholesterol and triglycerides, which leads to atherosclerosis, heart disease and stroke.



$\overline{}$			
()	MAIFIIFF	EXTENSION PANEL	(1(3))

CBC/Chemistry Profile • DHEA-S • PSA (prostate-specific antigen) Homocysteine • C-Reactive Protein (high-sensitivity) Free Testosterone • Total Testosterone • Estradiol • TSH for thyroid function • Vitamin D (25-hydroxyvitamin D) • Hemoglobin A1c

MALE HORMONE ADD-ON PANEL* (LCADDM)

Pregnenolone and Dihydrotestosterone (DHT)

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. This panel provides information about a testosterone metabolite that can affect the prostate; and the hormone pregnenolone that acts as a precursor to all other steroid hormones.

MALE ELITE PANEL (LC100016)*

CBC/Chemistry Profile • 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 • HbA1c Vitamin D 25-OH • hs-CRP, ferritin • Homocysteine • Hemoglobin A1c

MALE COMPREHENSIVE HORMONE PANEL (LC100010)*

CBC/Chemistry Profile • DHEA-S, Estradiol • DHT • PSA
Pregnenolone • Total and Free Testosterone • SHBG • TSH • Free T3
This panel now includes Free T4 and Cortisol
with no increase in price!

MALE BASIC HORMONE PANEL (LC100012)

DHEA-S • Estradiol • Total and Free Testosterone • PSA

THYROID ADD-ON PANEL (LCTHYROID)

Free T3 & Free T4.

○ INSULIN (LC004333)

Helpful to assess insulin resistance.

NMR LIPOPROFILE® (LC123810)

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.

○ ADVANCED OXIDIZED LDL PANEL*(LC100035)

This panel looks at vascular inflammatory biomarkers, beginning with lifestyle choices to the development of metabolic and cardiovascular disease as well as the formation of vulnerable plaque. The panel contains the following tests: F2-Isoprostanes, Myeloperoxidase and Oxidized LDL.

HOMOCYSTEINE (LC100061)

High homocysteine is associated with heart attack, stroke, and dementia. Find out your homocysteine level so you can take steps to lower it if necessary.

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.

YOUR	YUUK
YOUR PRICE	PRICE

\$269

\$125

\$575

\$299

\$75

\$275

\$249

\$129

\$269 FEMALE LIFE EXTENSION PANEL (LC322535) CBC/Chemistry Profile • DHEA-S • Estradiol • Homocysteine C-Reactive Protein (high-sensitivity) • Progesterone • Free Testosterone Total Testosterone • TSH for thyroid function

Vitamin D (25-hydroxyvitamin D) • Hemoglobin A1c \$120 FEMALE HORMONE ADD-ON PANEL* (LCADDF)

Pregnenolone and Total EstrogenTo provide an even more in-depth analysis of a woman's hormone status, Life Extension has created this panel as an addition to the Female Life Extension Panel. This panel provides information about

Female Life Extension Panel. This panel provides information about total estrogen status and the hormone pregnenolone that acts as a precursor to all other steroid hormones.

\$575 FEMALE ELITE PANEL (LC100017)*

\$75

\$29.90

\$99

\$54

CBC/Chemistry Profile • 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 • HbA1c Vitamin D 25-OH • hs-CRP • Ferritin • Homocysteine • Hemoglobin A1c

FEMALE COMPREHENSIVE HORMONE PANEL (LC100011)*

\$299 **CBC/Chemistry Profile •** DHEA-S, Estradiol • Total Estrogens Progesterone • Pregnenolone • Total and Free Testosterone • SHBG TSH • Free T3

This panel now includes Free T4 and Cortisol with no increase in price!

FEMALE BASIC HORMONE PANEL (LC100013)

DHEA-S • Estradiol • Total and Free Testosterone • Progesterone

\$55 WEIGHT LOSS PANEL-COMPREHENSIVE (LC100028)

CBC/Chemistry Profile • DHEA-S • Free andTotal Testosterone Estradiol • Progesterone • Cortisol, TSH • Free T3 • Free T4 Reverse T3 • Insulin • Hemoglobin A1c • Vitamin D 25-hydroxy C-reactive protein (high sensitivity) • Ferritin

HEALTHY AGING PANEL-COMPREHENSIVE (LC100026)*

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 • Insulin

\$285 DIABETES MANAGEMENT PROFILE – COMPREHENSIVE (LC100040)

Hemoglobin A1C • Glucose • Insulin • Lipid Panel • Glycomark



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 is NOT a complete listing of LE blood test services. Call **1-800-208-3444** for additional information.

- * 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-Iysine
L-Taurine Powder
L-Tyrosine Powder
Super Carnosine

Blood Pressure & Vascular Support

Taurine

Advanced Olive Leaf Vascular Support with Celery Seed Extract
Arterial Protect
Blood Pressure Monitor Arm Cuff
Dual Action Blood Pressure
Endothelial Defense™ with Pomegranate
Complete and CORDIART™
Endothelial Defense™ with GliSODin®
Natural BP Management
NitroVasc with CORDIART™
Pomegranate Complete
Pomegranate Fruit Extract
Triple Action Blood Pressure AM/PM
VenoFlow™

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 CocoaMind™ 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 Memory Protect Migra-Éeze™ Neuro-Mag® Magnesium L-Threonate Neuro-Mag® Magnesium L-Threonate with Calcium and Vitamin D3 Optimized Ashwagandha Extract 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
Digest RC®
Effervescent Vitamin C - Magnesium Crystals
Enhanced Super Digestive Enzymes
Enhanced Super Digestive Enzymes
w/Probiotics
EsophaCool™
Esophageal Guardian

Extraordinary Enzymes
Fiber-Immune Support
Gastro-Ease™
Ginger Force®
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™
Optimized NAD+ Cell Regenerator™
with Resveratrol
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

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 OMEGA FOUNDATIONS® Provinal® Purified Omega-7

OMEGA FOUNDATIONS® Vegetarian DHA Organic Golden Flax Seed

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 InSea^{2®} and Crominex® 3+ Mega Benfotiamine Tri Sugar Shield®

Heart Health Aspirin (Enteric Coated)

Cardio Peak™ with Standardized Hawthorn and Arjuna Homocysteine Resist Optimized Carnitine with GlycoCarn® Super Ubiquinol CoQ10 With BioPQQ® Super Ubiquinol CoQ10 with BioPQQ® Super Ubiquinol CoQ10 with Enhanced Mitochondrial Support™ Super-Absorbable CoQ10 Ubiquinone with d-Limonene TMG Powder TMG Liquid Capsules

BioActive Folate & Vitamin B12 Caps

Hormone Balance

DHEA (Dehydroepiandrosterone) Inner Power Pregnenolone Triple Action Cruciferous Vegetable Extract with Resveratrol Triple Action Cruciferous Vegetable Extract

Immune Support

AHCC® Enhanced Zinc Lozenges Immune Modulator with Tinofend® Immune Protect with PARACTIN® Immune Senescence Protection Formula™ Kinoko® Gold AHCC Kinoko® Platinum AHCC Kyolic® Garlic Formula 102 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 Lozenaes

Inflammation Management

5-LOX Inhibitor with AprèsFlex®
Advanced Bio-Curcumin® with Ginger & Turmerones
Black Cumin Seed Oil
Black Cumin Seed Oil with Bio-Curcumin®
Boswella
Cytokine Suppress™ with EGCG
Serraflazyme
Specially-Coated Bromelain
Super Bio-Curcumin®
Zyflamend® Whole Body

Joint Support

Calcium D-Glucarate

Arthro-Immune Joint Support
ArthroMax® Advanced with UC-II® & AprèsFlex®
ArthroMax® with Theaflavins & AprèsFlex®
ArthroMax® Herbal Joint Formula
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 with HepatoProtection Complex

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

Ageless Cell™
Alpha-Lipoic Acid
AMPK Activator
AppleWise Polyphenol Extract
Berry Complete
Blueberry Extract
Blueberry Extract with Pomegranate

DNA Protection Formula Enhanced Berry Complete with Acai Essential Daily Nutrients Grapeseed Extract with Resveratrol & Pterostilbene Mediterranean Whole Food Blend Mega Green Tea Extract (decaffeinated) Mega Green Tea Extract (lightly caffeinated) Optimized Fucoidan with Maritech® 926 Optimized Resveratrol

Optimized Resveratrol with Nicotinomide . Riboside

pTeroPure®

Pycnogenol® French Maritime
Pine Bark Extract

Resveratrol with Pterostilbene RNA (Ribonucleic 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 Pomi-T® Prelox® Natural Sex for Men® Super MiraForte with Standardized Lignans Triple Strength ProstaPollen™ Ultra Natural Prostate

Minerals

Boron Extend-Release Magnesium Ionic Selenium Iron Protein Plus Magnesium (Citrate) Magnesium Caps Only Trace Minerals Optimized Chromium with Crominex® 3+ Sea-Iodine™ Se-Methyl L-Selenocysteine Vanadyl Sulfate Zinc Caps

Miscellaneous

Potassium lodide Solarshield® Sunglasses

Mood & Stress Management

5 HTP L-Theanine Natural Cortisol Balance 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™ 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 Xyliwhite Mouthwash

Pet Care

Cat Mix Dog Mix

Probiotics

Bifido GI Balance FLORASSIST* Balance FLORASSIST* GI with Phage Technology FLORASSIST® Heart Health FLORASSIST® Immune Health FLORASSIST® Mood FLORASSIST® Oral Hygiene FLORASSIST® Throat Health Jarro-Dophilus® for Women Theralac® Probiotics
TruFlora® Probiotics

Advanced Anti-Glycation Peptide Serum

Skin Care

Advanced Growth Factor 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-Redness & Adult Blemish Lotion Broccoli Sprout Cream Collagen Boosting Peptide Serum

DNA Repair Cream Essential Plant Lipids Reparative Serum Eye Lift Cream Face Rejuvenating Anti-Oxidant Cream

Fine Line-Less Healing Formula Healing Vitamin K Cream

Hvaluronic Facial Moisturizer Hyaluronic Oil-Free Facial Moisturizer Hydrating Anti-Oxidant Facial Mist Hydroderm

Lifting & Tightening Complex Melatonin Cream

Mild Facial Cleanser

Multi Stem Cell Skin Tightening Complex Neck Rejuvenating Anti-Oxidant Cream Resveratrol Anti-Oxidant Serum

Shade Factor™ Shade Factor™ Sunscreen Lotion Shade Factor™ Sunscreen Spray Skin Care Collection Anti-Aging Serum

Skin Care Collection Body Lotion Skin Care Collection Day Cream

Skin Care Collection Night Cream Skin Firming Complex Skin Lightening Serum

Skin Restoring Phytoceramides with Lipowheat® Skin Stem Cell Serum

Skin Tone Equalizer

Stem Cell Cream with Alpine Rose

Tightening & Firming Neck Cream Triple-Action Vitamin C Cream Ultimate MicroDermabrasion Ultra Eyelash Booster

Ultra Lip Plumper Ultra Wrinkle Relaxer

Under Eye Refining Serum Under Eye Rescue Cream

Vitamin C Serum Vitamin D Lotion Vitamin E-ssential Cream

Youth Serum

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 with CherryPure® Whey Protein Isolate (Chocolate and Vanilla Flavor)

Ascorbyl Palmitate Benfotiamine with Thiamine Beta-Carotene BioActive Complete B-Complex **Biotin** Buffered Vitamin C Powder Fast-C® with Dihydroquercetin Gamma E Mixed Tocopherol Enhanced with Sesame Lignans Gamma E Mixed 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 K with Advanced K2 Complex Vitamin B12 Vitamin B6 Vitamin C with Dihydroguercetin Vitamin D3 with Sea-Iodine™ Vitamin D3

Weight Management

Vitamins D and K with Sea-Iodine™

7-Keto® DHEA Metabolite Advanced Anti-Adipocyte Formula Advanced Natural Appetite Suppress CalReduce Selective Fat Binder DHEA Complete Garcinia HCA HCActive™ Garnicia 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 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

			YC	UR PRIC	CE				YO	UR PRIC	F
ITEM N	o. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each QTY Total	ITEM I	No. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each QTY Tot
	A					01008	BLAST™ • 600 grams of powder	26.95	20.21		
01524	ACETYL-L-CARNITINE • 500 mg, 100 veg. caps	34.00	25.50	22.50		02025	BLOOD PRESSURE (Dual Action) • 60 veg. tabs	44.00	33.00	28.00	
01874	ACETYL-L-CARNITINE ARGINATE ● 90 veg. caps	52.00	39.00	35.00		70000	BLOOD PRESSURE MONITOR (ACCUFIT™) • med/lg cuff	79.99	49.99		
01628	ADRENAL ENERGY FORMULA • 60 veg. caps	24.00	18.00	16.50		70004	BLOOD PRESSURE MONITOR • Digital wrist cuff	69.95	52.46		
01630	ADRENAL ENERGY FORMULA • 120 veg. caps	46.00	34.50	31.50		02024	BLOOD PRESSURE (Triple Action AM/PM) • 60 veg. tabs	44.00	33.00	28.00	
01828	ADVANCED LIPID CONTROL • 60 veg. caps	30.00	22.50	20.25		01214	BLUEBERRY EXTRACT • 60 veg. caps	22.50	16.88	15.00	
02119	AGELESS CELL™ • 30 softgels	40.00	30.00	27.00		01438	BLUEBERRY EXTRACT W/ POMEGRANATE • 60 veg. caps	30.00	22.50	20.25	
00681	AHCC ® • 500 mg, 30 caps	59.98	44.99			01506	BONE FORMULA (DR. STRUM'S INTENSIVE) • 300 caps	56.00	42.00	37.50	
24404	AHCC® (KINOKO® PLATINUM) • 750 mg, 60 veg. caps	84.95	63.71			01726	BONE RESTORE • 120 caps	22.00	16.50	14.25	
29727	AHCC® (KINOKO® GOLD) • 500 mg, 60 veg. caps	74.95	52.47			01727	BONE RESTORE W/VITAMIN K2 • 120 caps	24.00	18.00	16.50	
00457	ALPHA-LIPOIC ACID W/BIOTIN • 250 mg, 60 caps	37.00	27.75	24.00		01725	BONE STRENGTH FORMULA W/KOACT® • 120 caps	45.00	33.75	30.00	
01907	AMPK ACTIVATOR • 90 veg. caps	48.00	36.00	33.00		00313	BONE-UP® • 240 caps	28.95	21.71	20.41	
01509	ANTI-ADIPOCYTE FORMULA W/MERATRIM® & INTEGRA LEAN® (Advanced) • 60 veg. caps	39.00	29.25	27.00			BORON ● 3 mg, 100 veg. caps	5.95		3.94	
02140	ANTI-ALCOHOL w/HEPATOPRO COMPLEX • 60 caps	22.00	16.50	15.00		00202	BOSWELLA • 100 caps	38.00	28.50	22.50	
01625	APPLEWISE POLYPHENOL EXTRACT	21.00	15.75	14.25		01802	BRAIN SHIELD® GASTRODIN ◆ 300 mg, 60 veg. caps	33.00	24.75	22.50	
	600 mg, 30 veg. caps					01253	BRANCHED CHAIN AMINO ACIDS • 90 caps	19.50	14.63	12.75	
01039	ARGININE/ORNITHINE • 500/250, 100 caps	17.99	13.49			01942	BREAST HEALTH FORMULA • 60 caps	34.00	25.50	22.50	
00038	ARGININE/ORNITHINE POWDER ● 150 grams	22.95	17.21	14.25		00893	BRITE EYES III • 2 vials, 5 ml each	34.00	25.50	24.00	
01624	(L)-ARGININE CAPS • 700 mg, 200 veg. caps	26.50	19.88	17.44		01203	BROMELAIN (Specially-coated) 500 mg, 60 enteric coated tablets	21.00	15.75	14.25	
02004	ARTERIAL PROTECT • 30 veg. caps	44.00	33.00	29.00			C				
01617	ARTHROMAX® W/THEAFLAVINS & APRÈSFLEX® 120 veg. caps	44.00	33.00	30.00		01653	CALCIUM CITRATE W/VITAMIN D • 300 caps	24.00	18.00	15.94	
01618	ARTHROMAX® ADVANCED W/UC-II® & APRÈSFLEX®	36.00	27.00	24.00		01651	CALCIUM D-GLUCARATE • 200 mg, 60 veg. caps	18.00	13.50	11.25	
	60 caps					[†] 01823	CALREDUCE SELECTIVE FAT BINDER 120 mint chewable tablets	45.00	33.75	28.50	
	ARTHROMAX® HERBAL JOINT FORMULA ● 60 veg. caps	40.00		27.00		01700	CARDIO PEAK™ W/STANDARDIZED HAWTHORN & ARJUNA	36.00	27.00	24.00	
	ARTHRO-IMMUNE JOINT SUPPORT • 60 veg. caps		24.00	21.00		01700	120 veg. caps	30.00	27.00	24.00	
	ARTICHOKE LEAF EXTRACT ● 500 mg, 180 veg. caps	30.00	22.50	21.00		00916	CARNITINE W/GLYCOCARN® (Optimized) • 60 veg. caps	36.00	27.00	24.00	
01533	ASCORBYL PALMITATE ◆ 500 mg, 100 veg. caps	22.50	16.88	15.00		01532	L-CARNITINE • 500 mg, 30 veg. caps	15.00	11.25	9.90	
88800	ASHWAGANDHA EXTRACT (Optimized) ● 60 veg. caps	10.00	7.50	6.75		01829	CARNOSINE • 500 mg, 60 veg. caps	36.00	27.00	24.00	
01805	ASIAN ENERGY BOOST ● 90 veg. caps	24.00	18.00	16.50		02020	CARNOSINE (Super) • 500 mg, 60 veg. caps	40.00	30.00	27.00	
	ASPIRIN • 81 mg, 300 enteric coated tablets	6.00		4.00		01932	CAT MIX • 100 grams powder	14.00	10.50	8.25	
01923	ASTAXANTHIN WITH PHOSPHOLIPIDS • 4 mg, 30 softgels	16.00	12.00	10.50		01899	CHILDREN'S FORMULA LIFE EXTENSION MIX™ 100 chewable tablets	20.00	15.00	13.50	
00920	BENFOTIAMINE W/ THIAMINE • 100 mg, 120 veg. caps	19.95	14.96	13.95		00550	CHLORELLA • 500 mg, 200 tablets	23.98	17.99		
	BENFOTIAMINE (Mega) ◆ 250 mg, 120 veg. caps		22.50	20.25			CHLOROPHYLLIN • 100 mg, 100 veg. caps		18.00	15.00	
	BERRY COMPLETE • 30 veg. caps		15.75	14.00		01359	o, o .		26.25	10.00	
	BERRY COMPLETE W/ACAI (Enhanced) • 60 veg. caps		21.75	19.50			CHOL-SUPPORT™ • 60 liquid veg. caps		36.00	32.00	
	BETA-CAROTENE • 25,000 IU, 100 softgels		8.63				CHROMIUM W/CROMINEX® 3+ (Optimized)		6.75	6.00	
	BIFIDO GI BALANCE • 60 veg. caps	20.00	15.00	13.50			500 mcg, 60 veg. caps				
01873	BILBERRY EXTRACT • 100 mg, 90 veg. caps			24.00		01503	CINSULIN® W/INSEA2® AND CROMINEX® 3+ • 90 veg. caps	38.00	28.50	25.50	
	BIOACTIVE MILK PEPTIDES • 30 caps		13.50	12.00		01906	CISTANCHE (Standardized) ● 30 veg. caps	20.00	15.00	12.00	
	BIO-COLLAGEN W/PATENTED UC-II® • 40 mg, 60 small caps		27.00	24.00		01818	CITRIMAX® (Super) ● 180 veg. caps	40.00	30.00	28.50	
	BIOSIL™ • 5 mg, 30 veg. caps		15.99			00818	CLA BLEND W/SESAME LIGNANS (Super)	36.00	27.00	24.75	19.75
	BIOSIL™ • 1 fl oz		25.59			02102	120 softgels COCOAMIND™ • 14 packets	24.00	18.00	16.00	
	BIOTIN • 600 mcg, 100 caps	7.50		4.88						16.00	36.00
	BLACK CUMIN SEED OIL • 60 softgels		12.00	10.50			COGNITEX® W/BRAIN SHIELD® • 90 softgels		45.00	39.00	
	BLACK CUMIN SEED OIL W/BIO-CURCUMIN® • 60 softgels			22.50		01897	COGNITEX® W/PREGNENOLONE & BRAIN SHIELD® 90 softgels	62.00	46.50	39.75	37.50
	SUBTOTAL OF COLUMN 1						SUBTOTAL OF COLUMN 2				مألي
	SUBTOTAL OF COLUMN 1						DECEIVE 2506 OFF THE DET				

ITEM N	lo. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each	QTY Total	ITEM No	o. PRODUCT
01421	COGNITEX® BASICS • 60 softgels	38.00	28.50		24.00		80107	FINE LINE-LESS • 1 oz
01659	COGNIZIN® CDP CHOLINE CAPS • 250 mg, 60 veg. caps	36.00	27.00	25.50			80137	HEALING FORMULA ALL-IN-ONE CREAM • 1 oz
01945	COMPLETE B-COMPLEX (BioActive) ◆ 60 veg. caps	12.00	9.00	8.00			80102	HEALING VITAMIN K CREAM • 1 oz
02198	COMPREHENSIVE NUTRIENT PACKS ADVANCED • 30 packs	90.00	67.50	61.50			80109	HYALURONIC FACIAL MOISTURIZER • 1 oz
01949	COQ10 w/d-LIMONENE (Super-Absorbable) • 50 mg, 60 softgels	25.00	18.75	16.50	15.00		80110	HYALURONIC OIL-FREE FACIAL MOISTURIZER • 1 oz
01948	COQ10 w/d-LIMONENE (Super-Absorbable)	46.00	34.50	28.00	26.25		80138	HYDRATING ANTIOXIDANT FACE MIST ◆ 4 0Z
21051	100 mg, 100 softgels	00.00	00.50	00.00			80103	LIFTING & TIGHTENING COMPLEX • 1 oz
01951	COQ10 w/d-LIMONENE (Super-Absorbable) 100 mg, 60 softgels	30.00	22.50	20.00			80135	MELATONIN CREAM • 1 oz
01929	COQ10 (Super Ubiquinol) • 100 mg, 60 softgels	56.00	42.00	36.00	33.00		80114	MILD FACIAL CLEANSER • 8 fl. oz
01733	COQ10 w/BIOPQQ® (Super Ubiquinol) • 100 mg, 30 softgels	54.00	40.50	33.00	30.00		80159	MULTI STEM CELL SKIN TIGHTENING COMPLEX $ullet$ 1 oz
01426	COQ10 w/ENH MITOCHONDRIAL SUPPORT™	62.00	46.50	39.00	36.00		80122	NECK REJUVENATING ANTIOXIDANT CREAM • 2 oz
	(Super Ubiquinol) • 100 mg, 60 softgels				H		80150	RENEWING EYE CREAM • 1/2 oz
01425	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super Ubiquinol) • 50 mg, 100 softgels	58.00	43.50	34.50	31.50		80142	RESVERATROL ANTI-OXIDANT SERUM • 1 oz
)1427	COQ10 w/ENH MITOCHONDRIAL SUPPORT™	20.00	15.00	12.00			80166	SKIN FIRMING COMPLEX • 1 fl. oz
	(Super Ubiquinol) • 50 mg, 30 softgels						80112	SKIN LIGHTENING SERUM • 1/2 oz
01431	COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super Ubiquinol) • 200 mg, 30 softgels	62.00	46.50	39.00	36.00		80130	SKIN STEM CELL SERUM • 1 0Z
0862	CRAN-MAX® • 500 mg, 60 veg, caps	17.50	13.13	11.25			80164	SKIN TONE EQUALIZER • 0.4 fl oz
1424	CRAN-MAX® WITH ELLIROSE™ (Optimized) • 60 veg. caps	18.00	13.50	12.00			80143	STEM CELL CREAM W/ALPINE ROSE • 1 oz
1529	CREATINE CAPSULES • 120 veg. caps	10.95	8.21	6.94			80148	TIGHTENING & FIRMING NECK CREAM • 2 oz
1746	CREATINE WHEY GLUTAMINE POWDER • 454 grams (vanilla)	30.00	22.50	19.50			80161	TRIPLE ACTION VITAMIN C CREAM • 1 oz jar
0407	CURCUMIN® (Super Bio) • 400 mg, 60 veg. caps	38.00	28.50	26.25			80162	ULTIMATE MICRODERMABRASION • 8 fl. oz
1924	CURCUMIN® W/GINGER & TURMERONES (Advanced Bio)	30.00	22.50	20.25			80160	ULTRA EYELASH BOOSTER • 0.25 oz (2 units \$39)
	30 softgels						80116	ULTRA LIP PLUMPER • 1/3 oz
1804	CYTOKINE SUPPRESS™ W/EGCG • 30 veg. caps	30.00	22.50	20.25			80101	ULTRA WRINKLE RELAXER • 1 oz
	COSMESIS						80113	UNDER EYE REFINING SERUM ● 1/2 oz
80157	ADVANCED ANTI-GLYCATION PEPTIDE SERUM ◆ 1 oz	53.00	39.75	34.50			80104	UNDER EYE RESCUE CREAM • 1/2 oz
80165	ADVANCED GROWTH FACTOR SERUM • 30 ml	65.00	48.75	42.75	_		80129	VITAMIN C SERUM • 1 oz
80154	ADVANCED LIGHTENING CREAM • 1 oz	65.00	48.75	42.75			80136	VITAMIN D LOTION ◆ 4 oz
80155	ADVANCED PEPTIDE HAND THERAPY • 4 0z	46.00	34.50	29.25			80145	VITAMIN E-ESSENTIAL CREAM • 1 oz
80152	ADVANCED TRIPLE PEPTIDE SERUM • 1 oz	65.00	48.75	42.75			80149	YOUTH SERUM ◆ 1 oz
80140	ADVANCED UNDER EYE SERUM W/STEM CELLS • .33 oz	49.00	36.75	31.50				D
80139	AMBER SELF MICRODERMABRASION • 2 oz	49.00	36.75	31.50			00658	7-KETO® DHEA METABOLITE • 25 mg, 100 caps
0158	ANTI-AGING FACE OIL • 1 oz	59.00	44.25	39.00			01479	7-KETO® DHEA METABOLITE • 100 mg, 60 veg. caps
80118	ANTI-AGING MASK • 2 oz	72.00	54.00	47.52			01640	DHA (Vegetarian) ◆ 30 veg. softgels
30151	ANTI-AGING REJUVENATING FACE CREAM • 2 oz	65.00	48.75	42.75			00607	DHEA • 25 mg, 100 tablets (Dissolve in mouth)
30153	ANTI-AGING REJUVENATING SCALP SERUM ● 2 oz	46.00	34.50	29.25			01478	DHEA COMPLETE • 60 veg. caps
30134	ANTI-GLYCATION SERUM W/BLUEBERRY & POMEGRANATE EXTRACTS • 1 oz	33.00	24.75	23.51			00335	DHEA • 25 mg, 100 caps
80133	ANTIOXIDANT FACIAL MIST • 2 oz	32.00	24.00	22.80				DHEA • 15 mg, 100 caps
80105	ANTI-REDNESS & ADULT BLEMISH LOTION • 1 oz	74.50	55.88	49.17				DHEA • 50 mg, 60 caps
80144	BROCCOLI SPROUT CREAM • 1 oz	46.00	34.50	29.25				DHEA • 100 mg, 60 veg. caps
80156	COLLAGEN BOOSTING PEPTIDE SERUM • 1 oz	59.00	44.25	39.00				
80141	DNA REPAIR CREAM • 1 oz		36.75	31.50			02021	DIGESTIVE ENZYMES (Enhanced Super) ● 60 veg. caps
80108		74.95		49.46				DIGESTIVE ENZYMES w/PROBIOTICS (Enhanced Super) ● 60 veg
	EYE LIFT CREAM • 0.5 fl oz		44.25	39.00			01671	D, L-PHENYLALANINE • 500 mg, 100 veg. caps
	FACE REJUVENATING ANTIOXIDANT CREAM • 2 oz		52.13	45.87			01540	DMAE BITARTRATE • 150 mg, 200 veg. caps
	SUBTOTAL OF COLUMN 3				L			SUBTOTAL OF COLUMN 4

10 Unit Each QTY Total

49.17

53.00 39.75 34.07 79.50 59.63 52.47 58.00 43.50 38.28 58.00 43.50 38.28 39.95 29.96 28.50 74.50 55.88 49.17 33.00 24.75 20.33 59.00 44.25 38.94 59.00 44.25 39.00

64.00 48.00 42.24 65.00 48.75 42.75 46.00 34.50 29.25 53.00 39.75 34.50 85.00 63.75 56.10 74.00 55.50 51.75 59.00 44.25 66.00 49.50 43.50 39.00 29.25 26.25 59.00 44.25

39.00

49.17

49.17

13.50

8.81

11.00

9.00

16.50

12.75

15.00

18.00

39.00 29.25 26.25 59.00 44.25 64.00 48.00 42.24 89.95 67.46 59.82 74.50 55.88

74.50 55.88

85.00 63.75 56.10 36.00 27.00 25.25 28.00 21.00 19.50 65.00 48.75 42.75

28.00 21.00 18.00 40.00 30.00 27.00 20.00 15.00

48.00 36.00 32.40 16.00 12.00

19.00 14.25 12.75 24.00 18.00

14.00 10.50

14.00 10.50

19.95 14.96

22.00 16.50

28.00 21.00

18.75 14.06 18.00 13.50 11.25

Retail Each \$ 74.50 55.88

TEM N	lo. PRODUCT	Retail	1	4	10	1	ITEM N	o. PRODUCT	Retail	1	UR PRIO	10	
		Each \$	Unit Each	Unit Each	Unit	QTY Tota			Each \$	Unit Each	Unit Each	Unit Each	Q
570	DNA PROTECTION FORMULA • 60 veg. caps	34.00	25.50	24.00			01541	GLUTATHIONE, CYSTEINE & C • 100 veg. caps	20.00	15.00	13.50		Į
931	DOG MIX • 100 grams powder	18.00	13.50	11.25			01669	GLYCINE • 1,000 mg, 100 veg. caps	12.00	9.00	8.10		
006	DOPA-MIND™ • 60 veg. tabs	44.00	33.00	28.00			01411		36.00	27.00	25.50		
321	DR. PROCTOR'S ADVANCED HAIR FORMULA • 2 oz	39.95	29.96	24.00			01620	100 mg, 60 veg. caps GREEN COFFEE EXTRACT COFFEEGENIC®	22.00	24.00	21.00		
320	DR. PROCTOR'S HAIR SHAMPOO • 8 oz	24.95	18.71	16.50			01620	400 mg, 90 veg. caps	32.00	24.00	21.00		
	E						00953	GREEN TEA EXTRACT (Mega)•lightly caffeinated,100 veg. caps	30.00	22.50	18.00		
1997	ENDOTHELIAL DEFENSE™ w/POMEGRANATE COMPLETE AND CORDIART™ • 60 softgels	68.00	51.00	46.50			00954	GREEN TEA EXTRACT (Mega)•decaffeinated, 100 veg. caps	30.00	22.50	18.00		
1997	ENDOTHELIAL DEFENSE™ w/GLISODIN® • 60 veg. caps	54.00	40.50	36.00				Н					
1937	· ·		15.00	13.50			01074	5 HTP • 100 mg, 60 caps	27.95	20.96			
2009	ESOPHACOOL™ • 120 chewable tablets		15.00	13.50			*02002	HAIR, SKIN & NAIL REJUVENATION FORM W/VERISOL® 90 tabs	32.00	24.00	22.00		
1737			27.00	24.00			01738	HCA (Garcinia) • 90 veg. caps	17.00	12.75	11.25		
1042	, <u>, , , , , , , , , , , , , , , , , , </u>		15.00	13.50			29754	HCACTIVE™ GARCINIA CAMBOGIA EXTRACT • 90 caps		22.50	11.23		
	600 mg, 30 veg. tabs	20.00	.0.00	10.00				HEPATOPRO • 900 mg, 60 softgels		37.50	34.50		
706	EXTRAORDINARY ENZYMES • 60 caps	26.00	19.50	18.00			02121	U, U		19.50	17.50		
2008	(CALIFORNIA ESTATE) EXTRA VIRGIN OLIVE OIL • 500 ml (16.9 fl. oz)	33.00	24.75	22.50				HUPERZINE A • 200 mcg, 60 veg. caps		30.00	27.00		
1514	EYE PRESSURE SUPPORT W/MIRTOGENOL® • 30 veg. caps	38.00	28.50	25.50				HYDRODERM® ◆ 1 oz			49.00		
	F						00001	I I I I I I I I I I I I I I I I I I I	79.93	39.90	49.00		
054	FACE MASTER® PLATINUM • Facial Toning System	199.00	199.00				01704	IMMUNE MODULATOR W/TINOFEND® ◆ 60 veg. caps	17.00	12.75	11.25		
965	FAST-ACTING JOINT FORMULA • 30 caps	39.00	29.25	27.00		-	00955	IMMUNE PROTECT W/PARACTIN® ◆ 30 veg. caps		22.13	19.91		
717	FAST-C® W/DIHYDROQUERCETIN • 120 veg. tabs	26.00	19.50	18.00		_		IMMUNE SENESCENCE PROTECTION FORMULA™ •60 veg. tabs			27.00		
064	FEMMENESSENCE MACAPAUSE® • 120 veg. caps	34.99	26.24					INNERPOWER™ • 530 grams powder		31.50			
2007	FIBER-IMMUNE SUPPORT (Apple Cinnamon) • 235 grams	34.00	25.50	23.50			01674	INOSITOL CAPSULES • 1,000 mg, 360 veg. caps		46.50	43.50		
125	FLORASSIST® GI w/PHAGE TECHNOLOGY•30 liquid veg. caps	33.00	24.75	22.50		_		INTEGRA-LEAN® AFRICAN MANGO IRVINGIA		21.00	18.00		
821	FLORASSIST® HEART HEALTH • 60 veg. caps	32.00	24.00	21.00				150 mg, 60 veg. caps					
2124	FLORASSIST® IMMUNE HEALTH • 30 veg. caps	26.00	19.50	18.00		-	30731	IONIC SELENIUM • 2 oz, 300 mcg	13.69	10.27			
2120	FLORASSIST® ORAL HYGIENE • 30 lozenges	20.00	15.00	13.00		_	01677	IRON PROTEIN PLUS • 300 mg, 100 caps	28.00	21.00	19.50		
825	FLORASSIST® BALANCE • 30 liquid veg. caps	32.00	24.00	21.00			01492	IRVINGIA W/PHASE 3TM CALORIE CONTROL COMPLEX	56.00	42.00	36.00		
2000	FLORASSIST® MOOD • 60 caps	33.00	24.75	22.50		-		(Optimized African Mango) • 120 veg. caps J, K, L					
	FLORASSIST® THROAT HEALTH • 30 lozenges		15.00			_	52142	JARRO-DOPHILUS® PROBIOTIC FOR WOMEN	27.95	20.96			
913	FOLATE HIGH POTENCY (Optimized) • 5,000 mcg, 30 veg. tablets	18.00	13.50	12.00			02112	30 enteric-coated veg. caps	2,,00	20.00			
939	FOLATE (Optimized) • 1,000 mcg, 100 veg. tablets	15.00	11.25	10.00			00056	JARRO-DOPHILUS EPS® • 60 veg. caps	23.95	17.96			
1842	FOLATE + VITAMIN B12 (BioActive) • 90 veg. caps	12.00	9.00	8.00			01834	K W/ADVANCED K2 COMPLEX (Super) ● 90 softgels	30.00	22.50	20.25		
1544	FORSKOLIN • 10 mg, 60 veg. caps	16.00	12.00	10.50		_	01600	KRILL HEALTHY JOINT FORMULA • 30 softgels	32.00	24.00	21.75		
513	FUCOIDAN W/MARITECH® 926 (Optimized) • 60 veg. caps	36.00	27.00	24.75			01050	KRILL OIL (Jarrow) ● 60 softgels	33.95	25.46			
	G						00316	KYOLIC® GARLIC FORMULA 102 • 200 veg. caps	27.45	20.59			
2070	GAMMA E MIXED TOCOPHEROL/TOCOTRIENOLS • 60 softgels	40.00	30.00	27.00		-	00789	KYOLIC® RESERVE • 600 mg, 120 caps	28.95	21.71			
2075	GAMMA E MIXED TOCOPHEROL w/ENHANCED SESAME LIGNANS • 60 softgels		24.00			-	01681	LACTOFERRIN ● 60 caps	44.00	33.00	30.00		
1394	, , ,		18.71				00020	LECITHIN • 16 oz granules	18.00	13.50	12.00		
2100	GASTRO-EASE™ • 60 veg. caps	44.00	33.00	30.00			02155	LIFE EXTENSION MIX [™] • 315 tablets	80.00	60.00	52.00	43.75	
1122	GINGER FORCE® ● 60 liquid caps	34.95	26.21				02157	LIFE EXTENSION MIX™ W/EXTRA NIACIN • 315 tablets	80.00	60.00	52.00	43.75	
658	GINKGO BILOBA CERTIFIED EXTRACT™ 120 mg, 365 veg. caps	50.00	37.50	33.00			02154	LIFE EXTENSION MIX™ • 490 caps	90.00	67.50	58.00	47.50	
756	GLA WITH SESAME LIGNANS (Mega) • 60 softgels	19.50	14.63	13.50			02156	LIFE EXTENSION MIX™ POWDER • 14.81 oz	80.00	60.00	52.00	43.75	
)345	(L-) GLUTAMINE CAPSULES • 500 mg, 100 veg. caps		11.21	10.13			02165	LIFE EXTENSION MIX [™] • 315 tablets w/o copper	80.00	60.00	52.00	43.75	
0141	(L-) GLUTAMINE CAPSULES • 500 mg, 100 veg. caps (L-) GLUTAMINE POWDER • 100 grams		16.50	15.00			02164	LIFE EXTENSION MIX™ • 490 caps w/o copper	90.00	67.50	58.00	47.50	
	GLUCOSAMINE/CHONDROITIN CAPSULES • 100 caps		28.50				02166	LIFE EXTENSION MIX™ POWDER • 14.81 oz w/o copper	80.00	60.00	52.00	43.75	
JZZ	accordanine, offendroffin our outer • 100 caps	30.00	20.00	24.00			01608	LIVER EFFICIENCY FORMULA • 30 veg. caps	18.00	13.50	12.00		

			YC	UR PRIC	CE					YO	UR PRIC	E	
ITEM N	o. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each Q	ITY Total	ITEM N	o. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each (QTY Tota
01639	5-LOX INHIBITOR W/APRÈSFLEX® • 100 mg, 60 veg. caps	22.00	16.50	15.00			01551	NATURAL SLEEP® w/ MELATONIN (Enhanced) • 30 caps	22.00	16.50	15.00		
01678	L-LYSINE • 620 mg, 100 veg. caps	9.00	6.75	6.00	ш		01511	NATURAL SLEEP® W/O MELATONIN (Enhanced) ● 30 caps	20.00	15.00	13.50		
00455	LYCOPENE (Mega) • 15 mg, 90 softgels	35.00	26.25	22.50			01445	NATURAL SLEEP® MELATONIN • 5 mg, 60 veg. caps	18.00	13.50	12.00		
	M						00987	NATURAL STRESS RELIEF • 30 veg. caps	28.00	21.00	18.00		
01992	MACUGUARD® OCULAR SUPPORT w/SAFFRON• 60 softgels	25.00	18.75	17.50	ш		01603	NEURO-MAG® MAGNESIUM L-THREONATE ● 90 veg. caps	40.00	30.00	27.00		
01993	MACUGUARD® OCULAR SUPPORT w/SAFFRON & ASTAXANTHIN• 60 softgels	44.00	33.00	30.00			01602	NEURO-MAG® MAGNESIUM L-THREONATE w/CALCIUM & VITAMIN D3 • 225 grams • Lemon flavor	40.00	30.00	27.00		
01459	MAGNESIUM CAPS • 500 mg, 100 veg. caps	12.00	9.00	7.50			01990	NITROVASC w/CORDIART™ • 30 veg. caps	18.00	13.50	12.00		
01682	MAGNESIUM (CITRATE) • 160 mg, 100 veg. caps	12.00	9.00	7.50			01903	NK CELL ACTIVATOR™ • 30 veg. tablets	45.00	33.75	31.50		
02107	(EXTEND-RELEASE) MAGNESIUM • 60 veg. caps	13.00	9.75	8.75			00373	NO FLUSH NIACIN • 800 mg, 100 caps	19.00	14.25	12.75		
01908	MEDITERRANEAN TRIM WITH SINETROL™-XPUR 60 veg. caps	18.00	13.50	12.00			01824	OLIVE LEAF VASCULAR SUPPORT W/CELERY SEED EXTRACT	26.00	27.00	24.00		
02109	MEDITERRANEAN WHOLE FOOD BLEND ● 90 veg. caps	44.00	33.00	30.00			01024	(Advanced) • 60 veg. caps	30.00	27.00	24.00		
01668	MELATONIN • 300 mcg, 100 veg. caps	5.75	4.31	3.75			01988		45.00	33.75	31.50	24.75	
01083	MELATONIN ● 500 mcg, 200 veg. caps	18.00	13.50	12.00				OLIVE EXTRACT, KRILL & ASTAXANTHIN (SUPER) • 120 softgels					
00329	MELATONIN ● 1 mg, 60 caps	5.00	3.75	3.47			01983	OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 60 softgels	18.00	13.50	12.00	9.38	
	MELATONIN • 3 mg, 60 veg. caps	8.00	6.00	5.16			01982	OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 120 softgels	32.00	24.00	21.00	17.05	
00331	MELATONIN • 10 mg, 60 veg. caps	28.00		18.00			01984		34.00	25.50	23.25	18.00	
	MELATONIN ◆ 3 mg, 60 veg. lozenges	8.00		5.16				OLIVE EXTRACT (Super) • 120 enteric coated softgels					
	MELATONIN (Fast-Acting Liquid) • 2 fl. oz (Citrus-Vanilla)	12.00	9.00	8.25			01985		20.00	15.00	13.50	10.50	
01787	MELATONIN TIMED RELEASE • 300 mcg, 100 veg. tabs	12.00		8.25			01006	OLIVE EXTRACT (Super) • 60 enteric coated softgels OMEGA 3 EPA/DHA w/SESAME LIGNANS &	22.00	24.00	21.00	17.05	
	MELATONIN TIMED RELEASE • 750 mcg, 60 veg. tablets	8.00	6.00	5.25			01900	OLIVE EXTRACT (Super) • 240 small softgels	32.00	24.00	21.00	17.23	
	MELATONIN TIMED RELEASE • 3 mg, 60 veg. tabs	12.00	9.00	8.25			01991	ONCE-DAILY HEALTH BOOSTER • 60 softgels	54.00	40.50	38.00		
	MEMORY PROTECT • 36 day supply	24.00		16.00			02113	ONE-PER-DAY • 60 tablets	22.00	16.50	15.00		
	METHYLCOBALAMIN • 1 mg, 60 veg. lozenges (vanilla)	9.95	7.46	6.00			01328	ONLY TRACE MINERALS • 90 veg. caps	15.00	11.25	9.38		
	METHYLCOBALAMIN • 5 mg, 60 veg. lozenges (vanilla)		24.00	18.75	17.25			P					
	MIGRA-EEZE™ (Butterbur) • 60 softgels		24.75	22.00			01789	PALMETTOGUARD® SAW PALMETTO W/BETA-SITOSTEROL	15.00	11.25	10.50	9.00	
	MILK THISTLE (European) ● 60 veg. caps		25.50	22.50			04700	30 softgels	00.00	04.00	40.50	40.00	
	MILK THISTLE (European) ● 60 softgels		21.00	18.75			01790	PALMETTOGUARD® SAW PALMETTO/ NETTLE ROOT W/BETA-SITOSTEROL • 60 softgels	28.00	21.00	19.50	18.00	
	MILK THISTLE (European) • 120 softgels		33.00	30.00			*00342	PECTA SOL-C® MODIFIED CITRUS PECTIN • 454 grams powder	109.95	93.46			
01940	MIRAFORTE w/STANDARDIZED LIGNANS (Super) ● 120 veg caps	62.00	46.50	42.00			*01080	PECTA SOL-C® MODIFIED CITRUS PECTIN • 270 veg. caps	79.95	67.96			
	MITOCHONDRIAL BASICS W/BIOPQQ® • 30 caps		33.00	30.00			01811	PEONY IMMUNE • 60 veg. caps	36.00	27.00	24.00		
	MITOCHONDRIAL ENERGY OPTIMIZER w/BIOPQQ®●120 caps		54.00	48.00			**00673	PGX® PLUS MULBERRY (WellBetX®) ● 180 veg. caps	34.95	26.21			
00065	MK-7 • 90 mcg, 60 softgels	28.00	21.00	18.75			01953	POMEGRANATE COMPLETE • 30 softgels	24.00	18.00	15.75		
00451	MSM (Methylsulfonylmethane) • 1,000 mg, 100 caps	14.00	10.50	8.96			00956	POMEGRANATE FRUIT EXTRACT • 30 veg. caps	19.50	14.63	13.16		
0450	N AOFTVI I OVOTEINE - COO CO	44.00	10.50	0.05			~ 01837	POMI-T® • 60 veg. caps	35.00	26.25	24.00		
	N-ACETYL-L-CYSTEINE • 600 mg, 60 veg. caps		10.50	9.25			00577	POTASSIUM IODIDE • 130 mg, 14 tabs	6.95	5.21	3.94		
	NAD + CELL REGENERATOR™ • 100 mg, 30 veg. caps		16.50	15.00			01500	PQQ CAPS W/BIOPQQ® ◆ 10 mg, 30 veg. caps	24.00	18.00	13.50	12.00	
02144	NAD + CELL REGENERATOR™ NICOTINAMIDE RIBOSIDE 250 mg, 30 veg. caps	42.00	31.50	28.00			01647	PQQ CAPS W/BIOPQQ® • 20 mg, 30 veg. caps	40.00	30.00	24.00	21.00	
02145	NAD + CELL REGENERATOR™ W/RESVERATROL (Optimized) 30 veg. caps	50.00	37.50	34.00			00302	PREGNENOLONE ◆ 50 mg, 100 caps	26.00	19.50	16.50		
01807	NATURAL APPETITE SUPPRESS (Advanced) ◆ 60 veg. caps	38.00	28.50	25.50			00700	PREGNENOLONE • 100 mg, 100 caps	30.00	22.50	20.25		
00984	NATURAL BP MANAGEMENT • 60 tablets	44.00	33.00	30.00			**01373	PRELOX® NATURAL SEX FOR MEN® • 60 tablets	52.00	39.00	36.00		
02012	NATURAL CORTISOL BALANCE • 30 veg. caps	45.00	33.75	30.00			00525	PROBOOST™ THYMIC PROTEIN A • 30 packets	66.60	49.95			
01892	NATURAL ESTROGEN ● 60 veg. tabs	38.00	28.50	25.50			01441	PROGESTA-CARE® ◆ 4 oz cream	36.39	27.29	25.72		
01626	NATURAL SEX FOR WOMEN® 50+ (Advanced) • 90 veg. caps	59.00	44.25	34.00			01928	PROSTATE FORMULA (Ultra Natural) • 60 softgels	38.00	28.50	26.25	24.00	
01444	NATURAL SLEEP® ◆ 60 veg. caps	13.00	9.75	7.50			01909	PROSTAPOLLEN™ (Triple strength) • 30 softgels	28.00	21.00	18.75		
	SUBTOTAL OF COLUMN 7							SUBTOTAL OF COLUMN 8					

	o. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each	QTY	Total
1742	PROTEIN-ISOLATE (Whey) Vanilla • 403 grams		22.50	19.50			
1743	PROTEIN-ISOLATE (Whey) Chocolate • 437 grams	30.00	22.50	19.50			
	PROTEIN CONCENTRATE (New Zealand Whey) Vanilla 500 grams	30.00	22.50	19.95			
	PROTEIN CONCENTRATE (New Zealand Whey) Chocolate 640 grams	30.00	22.50	19.95			
1812	PROVINAL® PURIFIED OMEGA-7 • 30 softgels	27.00	20.25	18.00			
1676	PS CAPS (Phosphatidylserine) • 100 mg, 100 veg. caps	54.00	40.50	36.00			
1508	PTEROPURE® Pterostilbene● 50 mg, 60 veg. caps	32.00	24.00	22.50			
1209	PUMPKIN SEED EXTRACT (Water-soluble) • 60 veg. caps	20.00	15.00	13.50			
	PYCNOGENOL® FRENCH MARITIME PINE BARK EXTRACT 100 mg, 60 veg. caps	64.00	48.00	45.00			
	PYRIDOXAL 5'-PHOSPHATE • 100 mg, 60 veg. caps	22.00	16.50	14.85			
	Q, R						
	QUERCETIN (Optimized) • 250 mg, 60 veg. caps		16.50	15.00			
	RED YEAST RICE (Bluebonnet) • 600 mg, 60 veg. caps		13.56				
	REGIMINT • 60 enteric-coated caps		14.96	14.00			
1708	REISHI EXTRACT MUSHROOM COMPLEX • 60 veg. caps		22.50	20.25			
	RESVERATROL W/PTEROSTILBENE • 100 mg, 60 veg. caps		27.00	24.00			
	RESVERATROL W/NICOTINAMIDE RIBOSIDE (Optimized) • 30 veg. caps	42.00	31.50	27.00			
2030	RESVERATROL (Optimized) • 60 veg. caps	46.00	34.50	31.00			
0889	RHODIOLA EXTRACT • 250 mg, 60 veg. caps	14.00	10.50	9.00			
	RIBOGEN™ FRENCH OAK WOOD EXTRACT 200 mg, 30 veg. caps	36.00	27.00	24.75			
0972	(D) RIBOSE POWDER • 150 grams	27.50	20.63	18.56			
1473	(D) RIBOSE TABLETS • 100 veg. tabs	32.00	24.00	21.00			
1609	RICH REWARDS® BREAKFAST GROUND COFFEE • 12 oz. bag	13.00	9.75				
	RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Mocha • 12 oz. bag	15.00	11.25	10.50			
	RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Vanilla • 12 oz. bag	15.00	11.25	10.50			
	RICH REWARDS® BREAKFAST BLEND WHOLE BEAN COFFEE 12 oz. bag		9.75				
	RICH REWARDS® DECAFFEINATED ROAST GROUND COFFEE 12 oz. bag		10.50				
	R-LIPOIC ACID (Super) • 240 mg, 60 veg. caps		36.75	33.75			
	RNA CAPSULES • 500 mg, 100 caps	17.95	13.46	12.12			
	SAFFRON W/SATIFDFAL® (Outining It as Source and	00.00	07.00	04.00			
1935	SAFFRON W/SATIEREAL® (Optimized) • 60 veg. caps SAMe (S-ADENOSYL-METHIONINE)		27.00 18.75	24.00 16.50			
1933	200 mg, 30 enteric coated tablets SAMe (S-ADENOSYL-METHIONINE) 400 mg, 30 enteric coated tablets	36.00	27.00	24.00			
1934	SAMe (S-ADENOSYL-METHIONINE) 400 mg, 60 enteric coated tablets	66.00	49.50	45.00			
	SEA-IODINE™ • 1,000 mcg, 60 veg. caps	8.00	6.00	5.40		۱	
	SE-METHYL L-SELENOCYSTEINE • 200 mcg, 90 veg. caps	11.00		7.50		۱	
	SERRAFLAZYME • 100 tablets		13.50	12.00		۱	
	SHADE FACTOR™ • 120 veg. caps		33.00	30.00		۱	
1938	o .og. oupo	50	- 5.50	_ 3.30			

			YO	UR PRIC	E		
ITEM N	o. PRODUCT	Retail Each	1 Unit	4 Unit	10 Unit		
20112	AULDE FLOTOR'N AUMONDERN ARRAY A G	\$	Each	Each	Each	QTY	Total
02118	SHADE FACTOR™ SUNSCREEN SPRAY • 6 fl. oz	22.00	16.50	14.25			
01884	SILYMARIN • 100 mg, 90 veg. caps		10.50	9.50			
01249	SINUS CLEANSER • 4 oz. bottle		18.75	07.50			
02129	SKIN CARE COLLECTION ANTI-AGING SERUM • 1.75 fl. oz		45.00	37.50			
02132	SKIN CARE COLLECTION BODY LOTION • 6 oz		21.00	18.00			
02130	SKIN CARE COLLECTION DAY CREAM • 1.65 fl. oz		37.50	33.00			_
02131	SKIN CARE COLLECTION NIGHT CREAM • 1.65 fl. oz	39.00		27.00			_
01596	SKIN RESTORING PHYTOCERAMIDES 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	76.00	57.00	50.00			
01649	SOY ISOFLAVONES (SUPER ABSORBABLE) • 60 veg. caps	28.00	21.00	18.75			
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			
01778	SUPER SELENIUM COMPLEX • 200 mcg, 100 veg. caps	14.00	10.50	9.00	8.25		
	т						
02023	TART CHERRY W/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	TRUFLORA® 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		Î	
02116	TWO-PER-DAY • 60 tablets	10.50	7.88	7.13		Î	
02115	TWO-PER-DAY • 120 tablets	20.00	15.00	13.50		Î	
02114	TWO-PER-DAY ◆ 120 caps	22.00	16.50	15.00		Î	
00326	L-TYROSINE • 500 mg, 100 tablets	13.50	10.13				
	SUBTOTAL OF COLUMN 10						

			YO	UR PRIC	Œ		
ITEM N	o. PRODUCT	Retail Each	1 Unit	4 Unit	10 Unit		
	U, V	\$	Each	Each	Each	QTY	Total
01921	URIC ACID CONTROL • 60 veg. caps	24.00	18.00	16.50			
00213	VANADYL SULFATE • 7.5 mg, 100 veg. tablets		11.25	9.38			
02102			39.00	36.00			
00408			14.21	12.00			
01327			13.50	10.50			
00372		7.65	5.74	4.99			
02028	VITAMIN B5 • 500 mg, 100 veg. caps (Pantothenic Acid)	11.00	8.25	7.50			
01535	VITAMIN B6 • 250 mg, 100 veg. caps	12.50	9.38	8.25			
00361	VITAMIN B12 • 500 mg, 100 lozenges	8.75	6.56	5.44			
01634	VITAMIN C w/DIHYDROQUERCETIN	10.00	7.50	6.75			
01004	1,000 mg, 60 veg. tablets	10.00	7.50	0.75			
00927	VITAMIN C w/DIHYDROQUERCETIN 1,000 mg, 250 veg. tablets	27.00	20.25	18.00			
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 • 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						
	X-R SHIELD • 90 veg. caps	15.00	11.25	9.75			
00409	XYLIWHITE™ MOUTHWASH • 16 oz	10.00	7.50				
01010	ZINA WAN PATENAY TO BE	7.05	5.00	- 0-			
	ZINC HIGH POTENCY • 50 mg, 90 veg. caps	7.95	5.96	5.25			
	ZINC LOZENGES • 60 veg. lozenges	9.00	6.75	6.00			
	ZINC LOZENGES (Enhanced) • 30 veg. lozenges	12.00	9.00	6.00			
**01051	ZYFLAMEND® WHOLE BODY • 120 liquid veg. caps	72.95	54.71				
33998	THE RIGHT TO TRY by Darcy Olsen • 2016	26.99	20.24				
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				
33877	THE TRUTH ABOUT MEN AND SEX by Abraham Morgentaler, MD, FACS • 2015	16.99	12.74				
33875	DOCTORED: THE DISILLUSIONMENT OF AN AMERICAN PHYSICIAN • by Sandeep Jauhar • 2015	26.00	19.50				
	SUBTOTAL OF COLUMN 11						

			Y0	UR PRIC	E	1	
ITEM N	o. PRODUCT	Retail Each \$	1 Unit Each	4 Unit Each	10 Unit Each	QTY	Total
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				
33867	THE COMPLETE MEDITERRANEAN DIET by Michael Ozner, MD • 2014	19.95	14.96				
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				
	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.
- † † These products are not 25% off retail price. Due to license restrictions this product is not for sale to custumers outside of the USA.



LifeExtension®

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** $^{\rm tt}$ Customers enrolled in Premier receive free unlimited standard delivery in the U.S., excluding U.S. territories, and do not have to pay the \$5.50 postage and handling fee. \$5.50** POSTAGE & HANDLING (Any size order, in the U.S, includes Alaska & Hawaii) C.O.D.s (ADD \$7 FOR C.O.D. ORDERS) 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 SHIPPING OTHER INTERNATIONAL AIR WILL BE ADDED. 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

Our premium rewards program.

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 unlimited FREE shipping. Anywhere in the U.S. ... including Alaska and Hawaii.*

4% BACK

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

\$50 BONUS

Instant \$50 LE Dollar Bonus. Offsets the cost of Premier, and can be redeemed simultaneously with enrollment.[†]

Experience Premier Today.

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[®]

- * 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 grift cards. Offer not available to international customers serviced by distributors of Life Extension products.
- **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.

 ¹ Can be redeemed simultaneously with Premier purchase, as long as you also make a product purchase totaling \$50 or more.

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.



For full product description and to order, call 1-800-544-4440 toll-free • www.LifeExtension.com

AlaskOmega® is a registered trademark of Organic Technologies. IFOS™ certification mark is a registered trademark of Nutrasource Diagnostics, Inc.

These products have been tested to the quality and purity standards of the IFOS™ program conducted at Nutrasource Diagnostics, Inc.



PO BOX 407198 FORT LAUDERDALE, FLORIDA 33340-7198

WHAT'S INSIDE

Visit us at www.LifeExtension.com

LifeExtension°

Magazine



7 NEW RESVERATROL DOSE

Resveratrol activates **sirtuin** proteins that promote longevity. **NAD+** is required for **sirtuins** to <u>function</u>. Optimal benefits can occur in response to **higher**-dose **NAD+** precursors with modest **resveratrol** intake.



40 IMPEDE UPPER-RESPIRATORY INFECTIONS

Scientists have identified a targeted **probiotic cocktail** that boosts the body's immune defenses to reduce colds, flu, and upper-respiratory complications.



64 REVERSING A ROOT CAUSE OF GLAUCOMA

Glaucoma is a common cause of blindness. A human study demonstrates **eye pressure** reduced by **24%** using two **plant extracts.**



30 BEN STILLER ADVOCATES PROSTATE CANCER SCREENING

Millions of men are skipping annual **PSA blood** tests. Actor **Ben Stiller** is a beneficiary of early diagnosis and has become a passionate advocate for **PSA screening**.



52 RELIEVE URINARY-TRACT SYMPTOMS

Prostate enlargement can lead to nighttime urinary frequency and weak stream. A combination of **natural extracts** helps alleviate urinary discomfort in men.



75 POMEGRANATE IMPROVES MITOCHONDRIA

Swiss researchers have identified a molecule produced in the body from **pomegranates** that protects mitochondrial function.