LIFE EXTENSION.COM COMPANY C

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

December 2015

Sore Throat Prevention Strategy

Protect Your DNA Against Environmental Toxins

Exclusive Report On Healthy Food Choices

Shielding Cells
From RadiationInflicted Mutations



PLUS-

Block The Deadly Effects Of After-Meal Blood Sugar Metformin Plus Aspirin Inhibit Pancreatic Cancer Calorie Restriction Lowers Risk Of Age-Related Disease



FOR WINTER SEASON SUPPORT

Zinc stimulates the activity of about 300 enzymes¹ and fortifies the immune system.²

Taking the best zinc lozenge is important for maintaining the body's natural defense system during the cold season—and throughout the year.

Enhanced Zinc Lozenges is a special "ionic formula" that delivers on the original promise of seasonal immune support.

IMMUNE SUPPORT FOR SEASONAL CHANGES

Scientific evidence is accumulating that **zinc acetate** is the preferred zinc compound to provide enhanced seasonal immune support.

Zinc acetate releases **100%** of its zinc as *ionic zinc*—positively charged zinc ions. **Zinc acetate** lozenges have been shown to strongly support immune function.³⁻⁵

Life Extension® Enhanced Zinc Lozenges provide zinc acetate with no other ingredients that could reduce the delivery of these immune-supportive, positively charged zinc ions. Enhanced Zinc Lozenges come in a naturally flavored peppermint lozenge.

SUGGESTED USE

The suggested serving size of one vegetarian lozenge of **Life Extension® Enhanced Zinc Lozenges** provides:

ZINC (as zinc acetate)

18.75 mg

These special **zinc lozenges** are <u>not</u> meant for daily use. Only a few of these lozenges are usually required when extra support is needed for seasonal changes. Its good to keep a bottle close by at home or work for immediate access.



Zinc Lozenges

Item #01961 • 30 vegetarian lozenges

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$12 | \$9 |
| 4 bottles | | \$6 each |

To order Enhanced Zinc Lozenges, call 1-800-544-4440

or visit

www.LifeExtension.com

References

- 1. J Nutr. 2000 May;130(5S Suppl):1437S-46S.
- 2. Am J Clin Nutr. 2007 Mar;85(3):837-44.
- 3. Curr Ther Res. 1998;59:595-607.
- 4. Ann Intern Med. 2000;133:245-52.
- 5. J Infect Dis. 2008;197:795-802.

LifeExtension®

VOLUME TWENTY ONE / NUMBER TWELVE • December 2015

REPORTS

34 MITIGATING IMPACT OF POST-MEAL GLUCOSE

Blood sugar levels surge after most meals, creating advanced glycation end products that accelerate aging processes. A fat-soluble form of vitamin B1 called **benfotiamine** can impede glycation reactions, thus helping to protect against this destructive mechanism of elevated glucose.

46 INCREASED CANCER RISKS OF CT SCANS

The National Cancer Institute researchers estimate that the 72 million CT scans performed annually will result in about 29,000 future cancer cases each year! Up to 44% of these CT scans may be medically unnecessary. Nutrients used by health-conscious individuals may afford some protection, but avoiding unnecessary exposure to radiation-emitting medical diagnostics is the first line of defense.

58 PROTECT AGAINST ENVIRONMENTAL TOXINS

Every year, over 4 billion pounds of toxic, untested chemicals are released into the environment, many of which are known carcinogens. These chemicals surround us in the air we breathe, the food we eat, and the water we drink. Overcooked foods create similar mutagenic compounds. Chlorophyllin is a water-soluble form of chlorophyll that has been shown to neutralize toxic compounds and protect cellular DNA against mutations that can lead to cancer.

70 REPORT: THE 2015 AMERICAN SOCIETY FOR NUTRITION CONFERENCE

Amid conflicting media reports, Ben Best reports on a scientific conference he attended where world experts describe the effects of the ketogenic diet, dietary sugar, dietary starch, dietary fat, weight-loss strategies, muscle loss in the elderly, and more. He provides his personal interpretation and recommendation for each.



24 REDUCE RISK OF SORE THROAT A novel **probiotic** lozenge has

been found to markedly lower strep and other throat infections in humans. One of several controlled studies showed a significant 84% reduction in incidence of strep throat.

DEPARTMENTS



AS WE SEE IT: CONSUMER REPORTS RECOGNIZES HAZARDS OF CT SCANS

CT scans emit high amounts of ionizing radiation that damage DNA and create mutations that can lead to cancer. Life Extension® long ago cautioned against unnecessary X-ray exposure, but these early warnings were largely ignored. In 2015, Consumer Reports magazine published their own expose on radiation-emitting imaging procedures and the number of excess cancers they cause.

17 IN THE NEWS

NSAIDs damage kidneys; vitamin D linked to fewer polyps; metformin with aspirin inhibits cancer; calorie restriction blocks aging diseases; nutrients slow thymus aging; vitamin D prevents elderly falls; coffee lowers cardiovascular risk; metformin inhibits breast cancer, and more.

83 ASK THE DOCTOR

Dr. Chris Kleronomos takes a highly comprehensive approach to pain management. He explains how natural medicine, acupuncture, chiropractic, psychology, diet, biofeedback, functional nutrition, and other therapies can safely alleviate pain without addictive drugs.

91 WELLNESS PROFILE

Upon the advice of his coach, soccer star Michael Lahoud began supplementing with Life Extension®'s vitamin D. He explains the remarkable effects vitamin D had on his athletic performance and overall health.

97 SUPERFOODS: WALNUTS

Unlike most nuts, walnuts are rich in polyunsaturated fats, including beneficial omega-3 and omega-6 fatty acids. Exciting research shows walnuts can protect against cancers, cardiovascular disease, cognitive decline, oxidative damage, and microbiomic and metabolic disorders.



LifeExtension°



VOLUME TWENTY ONE / NUMBER TWELVE • December 2015

PUBLISHER • LE Publications, Inc.

CONNECT WITH LIFE EXTENSION ON THE WEB!



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.

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/Health Advisors



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

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

CHIEF MEDICAL OFFICER

Steven Joyal, MD

VICE PRESIDENT OF PRODUCT INNOVATION & SCIENTIFIC DEVELOPMENT

Luke Huber, ND, MBA

SCIENTIFIC ADVISORY BOARD

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

Ben Best • Michael Downey • Laurie Ezreich • Loretta Granham Andrea Levy • Edward R. Rosick, DO, MPH, DABIM • Kira Schmid, ND

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

Advisors: 800-226-2370 • Advisory email: advisory@LifeExtension.com

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

Please mail your comments to Life Extension Magazine®,

Attn: Letters to the Editor, PO Box 407198, Fort Lauderdale, FL 33340 or email us: LEmagazine@LifeExtension.com

LIFE EXTENSION (ISSN 1524-198X) Vol. 21, No.12 ©2015 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 for BOA 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 for placement of non-Life Extension branded products or services contained in Life Extension for placement of an advert

SUPPORT ARTERIAL **HEALTH**

With The Most Complete Vitamin K Formula

If **vitamin K** levels are less than ideal, the matrix Gla-protein that lines the vascular system allows **calcium** to infiltrate soft tissues, similar to the way calcium is absorbed into bone. Optimal vitamin K levels function like a control switch—and keep calcium out of the arteries.1

There are three forms of vitamin K that are key factors to promoting arterial health:2

- Vitamin K1
- Vitamin K2 (MK-4) (rapidly absorbed)
- Vitamin K2 (MK-7) (long-acting)

K1 is the kind of vitamin K found in green vegetables, but only a fraction is absorbed in the bloodstream. Supplementing ensures optimal K1 blood levels within normal range.

K2 is found in meat, dairy, and egg yolks. **MK-4** is the most rapidly absorbed kind of K2, while MK-7 boasts a long half-life. This makes the two together the perfect complement to any vitamin K regimen.3

Life Extension® Super K with Advanced K2 Complex is the world's most comprehensive vitamin K formulation. Take with meals that contain some fat for optimal absorption.

> Just one daily softgel of Super K formula provides:

Vitamin K2 (MK-7) 200 mcg Vitamin K2 (MK-4) 1,000 mcg Vitamin K1 1,500 mcg

To order **Super K with Advanced K2 Complex** or Once-Daily Health Booster, call 1-800-544-4440 or visit www.LifeExtension.com

with Advanced K2 formula.]

References

- 1. Br J Nutr. 2012 Nov 14;108(9):1652-7.
- 2. Asia Pac J Clin Nutr. 2013;22(3):492-6.
- 3. Blood. 2007 Apr 15;109(8):3279-83.

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 connection of your testing doctor, as your doctor may need to increase your dose of 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.



LifeExtension®

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

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

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

Anna M. Cabeca, DO, FACOG, ABAARM, is a board certified Gynecologist and Obstetrician, as well as board certified in Anti-Aging and Regenerative Medicine, an expert in Functional Medicine, and an expert in women's health. She specializes in bioidentical hormone replacement therapy and natural alternatives, successful menopause and age management medicine.

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

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

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

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

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

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

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

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

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.

Miquelangelo 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.Fls.

Prof. Francesco Marotta, MD, PhD,

Montenapoleone Medical Center, Milan, Italy. Gastroenterologist and nutrigenomics expert with extensive international university experience. Consulting Professor, WHO-affiliated Center for Biotech & Traditional Medicine, University of Milano, Italy. Hon. Res. Professor, Human Nutrition Dept, TWU, USA. 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 antiaging medicine in Italy where he also works as a journalist and a writer.

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

Lambert Titus K. Parker, MD, practices internal medicine at the Integrative Longevity Institute of Virginia in Virginia Beach, VA.

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 staffs of Pacific Alliance Medical Center, Los Angeles, and other area

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

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

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

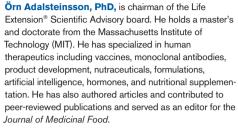
Michael D. Seidman, MD, FACS, is the director of otolaryngology-head and neck surgery for the Bloomfield satellite of Henry Ford Health System (HFHS), Detroit, MI, co-director of the Tinnitus Center, and co-chair of the Complementary/ Alternative Medicine Initiative for HFHS.

Ronald L. Shuler, BS, DDS, CCN, LN, is involved in immunoncology for the prevention and treatment of cancer, human growth hormone secretagogues, and osteoporosis. 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







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.



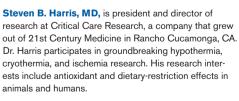
John Boik, PhD, is the author of two books on cancer therapy, Cancer and Natural Medicine (1996) and Natural Compounds in Cancer Therapy (2001), He obtained his doctorate at the University of Texas Graduate School of Biomedical Sciences with research at the MD Anderson Cancer Center, focusing on screening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



Frank Eichorn, MD, is a urologist specializing in prostate cancer for 10 years. He has a private practice in Bad Reichenhall, Germany, and is prostate cancer consultant at the Urologische Klinik Castringius, Planegg, Munich. In his integrative approach to prostate cancer he is working 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. Langsioen, MD. FACC, is a cardiologist specializing in congestive heart failure, primary and statin-induced diastolic dysfunction, and other heart diseases. A leading authority on coenzyme Q10, Dr. Langsjoen has been involved with its clinical application since 1983. He is a founding member of the executive committee of the International Coenzyme Q10 Association, a fellow of the American College of Cardiology, and a member of numerous other medical associations.



Dipnarine Maharaj MD, MB, ChB, FRCP (Glasgow), FRCP (Edinburgh), 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 boardcertified 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 nontoxic, hyper-effective antifreeze agents to fight unwanted ice formation in regenerative medicine and reduce mechanical icing.

BUILD BONE DENSITY AND INCREASE SKELETAL STRENGTH

With A Complete Combination of Critical Bone-Boosting Nutrients

Bone density loss is more than just a calcium deficiency—it also includes an insufficient intake of a host of other nutrients.

Bone Restore combines numerous bone-boosting nutrients into one superior, easy-to-take formula. In addition to 700 mg of highly-absorbable calcium, Bone Restore provides meaningful potencies of:

- Boron
- Vitamin D3
- Magnesium
- Manganese
- Zinc
- Silicon

These nutrients work together to keep aging bones strong.

Bone Restore also contains **vitamin K2**, which has been shown to play a critical role in maintaining healthy bone density by facilitating the transport of calcium from the bloodstream into the bone.

Who Should Take Bone Restore

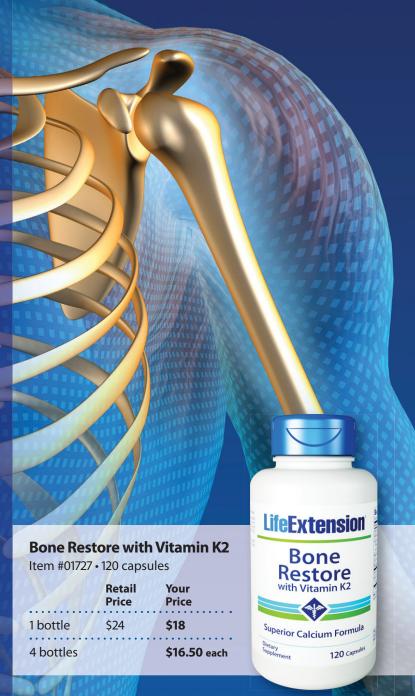
Progressive **loss of skeletal density** is a consequence of normal aging. Fortunately, an array of **nutrients** has been identified that can markedly impede and sometimes reverse this degenerative process.

Bone Restore is designed to provide the best forms of these nutrients in ideal potencies. It contains the expensive MK-7 form of vitamin K2. Many readers of this publication already obtain MK-7 in the Super K or Health Booster (formerly called Super Booster) formulas. For these individuals, Bone Restore is available without vitamin K2. The retail price for this formula of 120 capsules is \$22. If four bottles are purchased, the price is reduced to \$14.25 per bottle. (Item# 01726)

Non-GMO

Note: Those taking the anticoagulant drug Coumadin® (warfarin) should use BONE RESTORE without vitamin K2.

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



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

AS WE SEE IT

Consumer Reports Discovers Dangers Of CT Scans And X-Rays



BY WILLIAM FALOON

The March 2015 issue of *Consumer Reports* published an in-depth article about the cancer-causing effects of medical X-rays and CT scans. The <u>title</u> of this *Consumer Reports* article is:

"The Surprising Dangers Of CT Scans And X-rays."

We applaud **Consumer Reports** for publishing this expose. It will save human lives. We're taken aback, however, as to why the word "**surprising**" was used in the title.

There's nothing **surprising** about the striking number of **cancer deaths** caused by medical imaging procedures like **CT scans**. These imaging devices emit high amounts of **ionizing radiation** that damage DNA and create mutations that can lead to cancer.

The **National Cancer Institute** estimates that at least **15,000** cancer deaths will occur as a result of **CT scans** performed in **2007**.^{2,3}

This same report estimates **29,000** <u>new</u> cancer cases occurred in **2007** alone from **CT scan** exposure. Since exposure to radiation-emitting imaging procedures has risen approximately **6-fold**⁴ since **1980**, the number of <u>excess</u> **cancers** being caused is potentially astronomical.

What may have prompted **Consumer Reports** to publish their article was a recent study that looked at people who had CT scans as children and then

followed them for almost 10 years. This study found a **24%** increased **cancer risk** from just one **CT scan**, with each additional **CT** scan boosting cancer risk an additional **16%**. Cancer risk was greater with a younger age of exposure. While overall cancer rates in this group were low, the statistical significance was robust, meaning these frightening increases in cancer rates are unlikely to have occurred by chance.

Our earlier warnings about the dangers of radiation from medical diagnostic procedures like CT scans were ridiculed and largely ignored. We hope that the **Consumer Reports'** investigative analysis will encourage more patients to stand up to their doctors and question the necessity of exposing their body to cancer-causing radiation. As you'll read in this article, many X-rays and CT scans performed today should <u>not</u> be done.



COURTESY CONSUMER REPORTS® CONSUMERREPORTS.ORG

AS WE SEE IT

There is not a topic I have more hotly debated with physicians than the danger posed by ionizing radiation emitted from CT scans and X-ray imaging devices.

On a personal level, every doctor I have ever argued this point with has stated that CT scans and X-rays are 100% safe. I rarely violate rules of debate by stating that someone is categorically wrong, but I have never stepped back from declaring that exposing one's healthy cells to ionizing radiation from medical diagnostic imaging (like CT scans) increases cancer risk.

Life Extension® long ago published the frightening numbers of Americans who are contracting cancer as a result of prior CT scans or X-ray exposure. Medical authorities are now somewhat aware of this data, yet few who prescribe CT scans or X-ray imaging are paying attention to it.

One-Third Of Radiation Scans Are Unnecessary

Each year, there are **72 million** CT scans performed in the United States.⁶ This is up from **3 million** in 1980 when CT scans began to be aggressively marketed to doctors.⁷ CT scans provide superior imaging compared to conventional X-rays, but at the cost of vastly higher doses of **ionizing radiation**.⁷

A recent report found that as many as one-third of CT scans and other diagnostics that expose patients to high levels of radiation are being done too frequently. We at *Life Extension* believe that more than a third of radiation-emitting imaging procedures could be eliminated.

One obstacle we battle when it comes to this debate is physician prescribing practices that are very difficult to change. Doctors were



taught in school that radiation from medical diagnostic imaging was very <u>safe</u> and posed <u>no</u> long-term risk to their patients. Throughout their residency and into practice, the idea that **ionizing radiation** from medical diagnostic imaging is safe, and does not increase long-term cancer risk, is *consistently* reinforced to physicians, often by radiologists. In addition, physicians are often in the practice of *defensive medicine*, and order unnecessary imaging tests born out of the fear of litigation.

Especially worrisome is the fact that some physicians have a *financial investment* in the very medical diagnostic imaging centers to which patients are referred.

Consumer Reports magazine now urges patients to ask if their doctor has a financial interest in a diagnostic imaging center. It should not come as a surprise that when physicians invest in a CT scanner or other radiology equipment, they then have a financial incentive to refer more of their patients for CT scans and other imaging tests.

Consumer Reports urges all patients to question their doctor when a CT scan or X-ray is ordered, as some problems can be managed without powerful doses of radiation.⁸

Widespread Ignorance Of The Dangers

Consumer Reports conducted a survey and found that only **4%** of patients prescribed a **CT scan** had the knowledge to say "no" to their doctor.¹ This prompted one enlightened doctor to state that patients need to take the lead in questioning whether a CT scan or X-ray is necessary.

A **2012** study was done of medical personnel who worked with patients undergoing abdominal CT scans (which often emit the most radiation). This study found that less than **50**% understood that these scans could cause cancer.⁹

Another study revealed only **9%** of emergency room physicians said they knew that CT scans increased cancer risk.¹⁰

This widespread ignorance amongst professionals on the front lines of medical care is alarming.

Until doctors get up to speed on the risks posted by radiationemitting imaging devices, patients need to assert control and not capitulate to the exaggerated fears doctors instill to persuade patients to undergo unnecessary CT scans, X-rays, or other diagnostic imaging procedures involving ionizing radiation.

Defending Against Lawsuits

A study presented at the **2011** meeting of the **American Academy of Orthopedic Surgeons** provided clear evidence of why CT scans and other medical diagnostic imaging tests are being so overutilized.

It turns out that **35%** of imaging tests are being done by doctors out of fear of lawsuits.^{1,11-13} In other words, if sued by a patient (and zealous personal injury attorney) for malpractice, doctors need hard evidence showing the patient was aggressively <u>diagnosed</u>, as well as treated.

How To Reduce Radiation Exposure From Medical Diagnostic Tests

A partial solution to the widespread overexposure to ionizing radiation is to turn down the <u>amount</u> of radiation emitted from each scan. This can be done because most modern CT scanners can be *intensity modulated*. This means the <u>dose</u> of radiation needed to obtain a crisp picture of your insides can be greatly <u>reduced</u> based on your body mass and other factors.¹⁴

What hurried X-ray technicians have done too frequently is set the dose of radiation at the highest level for all patients, thereby eliminating the time needed to adjust the radiation dose to conform to each individual. This ensures great consistent images at the cost of many times the radiation dose required for most people. As a patient, you should insist that if a CT scan is needed, your body mass be evaluated and the <u>lowest</u> possible dose of radiation be used to obtain the needed images. ^{15,16}

A particularly disturbing trend pointed out by Consumer Reports is that children are too often being given adult-sized doses of radiation, which is many times what they need.1 The higher dose directly increases the child's cancer risk, vet rushed radiology technicians don't want to bother turning down the radiation intensity. The pressure to put patients on a fast-moving assembly line, with little regard for individualized care, is epidemic throughout today's hurried and increasingly depersonalized world of mainstream medical practice.

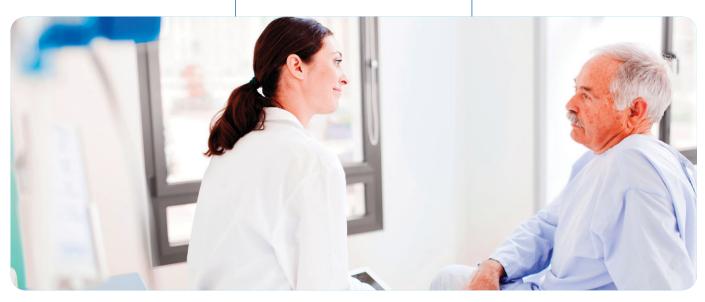
Modern Imaging Saves Lives!

It is important to point out that CT scans save more lives than the cancers they cause.¹⁴ The problem is they are being overused and wrongly used in too many cases.

For many diagnostics, **ultra-sound** devices can provide a clear internal image while emitting no ionizing radiation.¹⁷⁻¹⁹ **MRI** (magnetic resonance imaging) often provides a clearer picture than a CT scan for soft tissue,²⁰ but since MRIs cost much more, insurance companies increasingly are refusing to pay for them.²¹

Those who display outward symptoms of a **stroke** should insist on an immediate CT scan of their brain so that comprehensive stroke-reversal therapies can be promptly administered to reduce the risk of paralysis or death. CT scans can be done instantly compared to the much longer time it takes for magnetic resonance techniques. (Some hospital ER rooms are equipped to do quick MRI scans for stroke patients.)

When compared to the perils and pain of exploratory surgery, a CT scan is a breakthrough that should be utilized when appropriate.



AS WE SEE IT

For cancer patients, the use of multiple whole-body **PET/CT scans** with enhanced reading techniques provides enormous insight into whether treatments are working or need adjustment. For treatment of difficult and aggressive cancers, the long-term risk from exposure to the radiation from PET/CT scanners is less than the immediate risk posed by the aggressive cancer. 22,23

My Pain Of Saying No To Dentists

Dentists are adamant about doing annual X-rays and represent the strongest proponents of prophylactic X-rays. Not one dentist I have encountered has ever acknowledged there is any risk posed by these annual X-rays.

I have refused dental X-rays for most of my life despite dire

Startling **Numbers Of Excess Cancers**

In 2009, the journal Archives of Internal Medicine published findings showing the potential for adverse effects of ionizing radiation from medical imaging procedures like CT scans.

This study led by the **National Cancer Institute** used a model of radiation risk over time based on CT scans administered in the year 2007.

The results suggested that CT scans alone may contribute to 29,000 new cancer cases over time, and nearly 15,000 cancer deaths over time assuming 50% mortality.3

warnings by my dentists that there could be underlying tooth decay. When I developed some pain in my mouth two years ago, I consented to dental X-rays. No abnormality was revealed.

I then went to a medical specialist who said he could prescribe a CT scan of my entire jaw that might detect what was causing my considerable pain. This doctor also cautioned that some people develop a condition called idiopathic oral facial pain in which no underlying cause is detected by a CT scan. (Idiopathic means a disease of unknown origin.)

I declined the CT scan of my jaw and suffered fluctuating pain for almost a year and a half until the pain became so acute that I went back to the dentist for another dental X-ray, which this time revealed a single decayed tooth that was readily treated.

Had I opted for the CT scan, this decaying tooth would have likely been detected much earlier. Of course, very vulnerable parts of my head and neck would have also been exposed to high levels of radiation from the CT scan (much greater than typical dental X-rays).

So I paid a painful price for declining the CT scan of my jaw. I always like to relate real-world events so that readers understand the challenges of determining when to say "no" to a radiation-emitting imaging device. It's not always an easy decision.

Are Dental X-Rays Safe?

My dentist makes me sign a waiver of liability because I refuse to have annual X-ravs done. Other dentists refuse to treat me unless I capitulate to X-rays whenever they want to do them. There, of course. is considerable clinical value in dentists being able to view under your enamel.

My concern about dental X-rays was partially vindicated when a 2012 study published in the journal *Cancer* showed that people exposed to annual dental X-rays were twice as likely to develop a brain tumor called a meningioma.38 This type of tumor is usually benign and can be treated with radiation or surgery if needed, but who wants to go through this?





Experts Historically Underestimated Radiation Risks

When radiation was first discovered, doctors were wildly enthusiastic about being able to peer inside the body of a living human.

Sadly, radiation killed its early pioneers, who had no idea of its dangers.

Even as knowledge of radiation's lethal properties became apparent, experts consistently underestimated the risks.

One tragic example was an individual named Clarence Dally who intentionally exposed himself multiple times to ionizing radiation from Thomas Edison's "fluoroscope" invention. Within a few months, Dally began suffering debilitating fatigue, body aches, and multiple burn-like lesions on his hands. These lesions

turned out to be cancer that rapidly spread throughout his body. Dally lost both his arms to these malignant lesions, and died a painful death in 1904. Thomas Edison was said to be haunted for the rest of his life by Dally's cancer and death, and refused to have anything more to do with ionizing radiation.24

PHOTO BY LOUIS BACHRACH, BACHRACH STUDIOS, RESTORED BY MICHEL VUIJLSTEKE-UNITED STATES LIBRARY OF CONGRESS

In the 1950s, our federal government routinely conducted above-ground testing of **nuclear bombs** in the Nevada desert and claimed the radioactive fallout that

spread throughout much of the United States was "harmless." In 2002, the federal government admitted that the radiation emitted from these nuclear weapons tests caused 15,000 American cancer deaths.25 Critics claim this number grossly understates the actual number of cancer deaths.26-28

PHOTO COURTESY OF NATIONAL NUCLEAR SECURITY ADMINISTRATION / NEVADA SITE OFFICE



Operation Teapot **Nuclear Test**

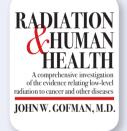
In the May 2002 edition of Life Extension® magazine, I wrote a tribute to Linus Pauling for his intellect and courage to challenge the federal government's claim that radioactive fallout was "safe."29 Dr. Pauling endured endless persecution from government



authorities for daring to interfere with the above-ground testing of nuclear bombs. Our government viewed Linus Pauling as a "communist," when this Nobel Prize-winning scientist was only seeking to protect Americans from the cancers he suspected would be caused by the radioactive fallout.

Dr. Pauling's tireless efforts eventually resulted in a ban against above-ground nuclear testing by both the United States and Russia, leading him to win his second Nobel Prize.30

Linus Pauling is the only person to ever win two undivided Nobel Prizes. New Scientist magazine ranks Dr. Pauling as



one of the 20 greatest scientists to ever live, an honor shared with such figures as Albert Einstein, Charles Darwin, and Isaac Newton.31

Yet here we are in 2015, and most of the medical establishment remains oblivious to the documented dangers of radiation-emitting imaging devices. This is despite irrefutable data provided by the late physicianscientist John Gofman, MD, PhD, who worked at the Los Alamos lab inventing nuclear weapons, went on to discover the existence of LDL cholesterol, and then campaigned relentlessly to limit unnecessary exposure to medical sources of ionizing radiation.³²⁻³⁷

John Gofman called me long ago and expressed gratitude that Life Extension regularly warned of the dangers posed by radiation-emitting medical imaging devices. He regretted so few doctors paid attention to the many books he authored documenting higher cancer rates in those exposed to medical radiation.

AS WE SEE IT

Modern dental X-rays emit less radiation than older devices, and thus may not pose as great a risk for meningioma. However, the authors concluded their 2012 study by warning:

"...there is little evidence to support the use of dental X-rays 'in search of occult pathoses in the asymptomatic patient' or 'routine dental radiographs at preset intervals for all patients. Although dental X-rays are an important tool in well-selected patients, efforts to moderate exposure to ionizing radiation to the head is likely to be of benefit to patients and health care providers alike."38

Said differently, these authors are suggesting that one minimize the number of dental X-rays they are exposed to, which contradicts what is being done in most dentist practices today.

Reducing And Repairing **DNA Damage** Inflicted By Radiation

When I reviewed the number of previous articles we have published about the dangers of ionizing radiation from medical imaging tests and CT scans, I was startled by how much hard data we had uncovered.

Some of our articles describe the potential protection one might obtain by having high doses of specific antioxidants in their body at the time a radiation-emitting imaging procedure is performed.³⁹ Other articles describe the potential for nutrients like **blueberry extract** to enhance **DNA repair** so that damage inflicted by medical radiation does not lead to future cancer.40

Some of you might remember an article I wrote last year about a strain of bacteria that was made resistant to 1.000 times the amount of radiation that would kill a human. The mechanism that enabled these bacteria to survive this onslaught of radiation-induced free radical attack was markedly enhanced **DNA repair**.41,42

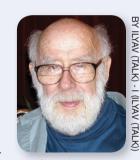
The data about the radiationprotecting effects of the nutrients

that many of you take daily continues to grow. This means that even if you have needed a CT scan or diagnostic X-ray procedure in the past, if you had these nutrients in vour body, vou may have gained some degree of protection. Regular intake of blueberry and other plant extracts may have facilitated enough **DNA repair** to offset

Tribute To Hero Who Uncovered Dangers Of CT-Scans And X-rays

John Gofman, MD, PhD, was a medical doctor, nuclear chemist. Manhattan Project scientist, co-discoverer of isotopes of uranium and protactinium, the first to separate plutonium in usable quantities, and an early member of the Life Extension Foundation.

Dr. Gofman fought to end policies that allow plutonium and other radioactivity from the nuclear power/weapons fuel chain to be dispersed into the environment. He repeatedly stood up to government pressure to suppress the truth about John Gofman, MD, PhD radiation health dangers.



Dr. Gofman's accomplishments extend to his groundbreaking research in cardiac medicine, which includes the identification and distinguishing of HDL cholesterol and LDL cholesterol. The Journal of Clinical Lipidology named him the "Father of Clinical Lipidology," honoring him for discoveries he made decades ago, which are now part of conventional cardiology.43

His tireless work to reduce unnecessary radiation exposure from medical procedures is finally taking hold with Consumer Reports disseminating data that emanated from work that Dr. Gofman initiated many decades ago.

Dr. Gofman firmly believed there is no safe threshold of ionizing radiation one should needlessly be exposed to. He used a linear no-threshold model of radiation risk and argued that far more cases of cancer and other diseases are caused by unnecessary exposure to medical radiation than what the "authorities" admit.44,45

Not everyone agrees with Dr. Gofman's "linear model of radiation **risk**" and a debate continues as to whether there is a low level of X-ray exposure that can be accepted as "safe."

Next time you review your blood test results and see LDL and HDL, remember it was Dr. Gofman's pioneering research in the 1950s that led to these lipids being included in standard blood test panels to evaluate vascular disease risk.

To read more about this brilliant physician-scientist, just type John Gofman into Google.

the known carcinogenic effects of the radiation. No one knows for certain.

In this month's issue, we describe some nutrients vou want to have in your body *prior* to being exposed to a radiation-emitting imaging procedure.

I want to again thank *Consumer* **Reports** for disseminating information that is almost identical to what we've been preaching for decades. While the mainstream media ignores most of what we publish, the **Consumer Reports** article generated press coverage that I believe will spare some humans from medical radiation-induced cancers.

For longer life,

William Faloon

References

- Available at: http://www.consumerreports. org/cro/magazine/2015/01/the-surprisingdangers-of-ct-sans-and-x-rays/index.htm. Accessed September 24, 2015.
- Available at: http://www.cancer.gov/aboutcancer/causes-prevention/hp-preventionoverview-pdg#section/all. Accessed September 24, 2015.
- Berrington de González A, Mahesh M, Kim K. et al. PRojected cancer risks from computed tomographic scans performed in the united states in 2007. Arch Intern Med. 2009 Dec 14;169(22):2071-7.
- Fazel R, Krumholz HM, Wang Y, et al. Exposure to low-dose ionizing radiation from medical imaging procedures. N Engl J Med. 2009 Aug 27;361(9):849-57.
- Mathews JD, Forsythe AV, Brady Z, et al. Cancer risk in 680,000 people exposed to computed tomography scans in childhood or adolescence: data linkage study of 11 million Australians. BMJ. 2013 May 21:346:f2360.
- Available at: http://www.cbsnews.com/ news/many-medical-imaging-tests-performed-in-u-s-are-unnecessary/. Accessed September 24 2015
- Brenner DJ, Hall EJ. Computed Tomography - An Increasing Source of Radiation Exposure. N Engl J Med. 2007 Nov 29;357(22):2277-84..
- Available at: http://pressroom.consumerreports.org/pressroom/2015/01/consumer-reports-warns-against-the-risks-of-radiationoverexposure-from-unnecessary-ct-scans. html. Accessed September 24, 2015.

- Uri IF. Lack of radiation awareness among referrers: implications and possible solutions. Int J Clin Pract. 2012 Jun;66(6):574-81.
- Lee CL Haims AH Monico EP et al Diagnostic CT scans: assessment of patient, physician, and radiologist awareness of radiation dose and possible risks. Radiology. 2004 May;231(2):393-8.
- 11. Available at: http://content.time.com/time/ health/article/0,8599,2053354,00.html. Accessed September 24, 2015.
- 12. Available at: http://www.sciencedaily.com/ releases/2009/02/090202175100.htm. Accessed September 24, 2015.
- Available at: http://www.aaos.org/news/ acadnews/2011/AAOS13_2_17.asp. Accessed September 24, 2015.
- McCollough CH, Primak AN, Braun N, Kofler J, Yu L, Christner J. Strategies for Reducing Radiation Dose in CT. Radiol Clin North Am. 2009 Jan;47(1):27-40.
- Hollingsworth C, Frush DP, Cross M, Lucaya J. Helical CT of the body: a survey of techniques used for pediatric patients. AJR Am J Roentgenol. 2003 Feb;180(2):401-6.
- Available at: http://www.fda.gov/MedicalDevices/Safety/AlertsandNotices/PublicHealthNotifications/ucm062185.htm#4. Accessed September 29, 2015.
- 17. Available at: http://www.fda.gov/Radiation-EmittingProducts/RadiationEmittingProductsandProcedures/MedicalImaging/ucm115357.htm. Accessed September 25, 2015.

Radiation Dose Comparison

Different types of medical diagnostic imaging tests pose varying degrees of risk from ionizing radiation exposure. A brief list of different medical diagnostic imaging tests that utilize ionizing radiation are provided to better identify scans that generate relatively high exposure in comparison to natural background radiation exposure.

| Diagnostic Procedure | Typical Effective Dose (mSv) ⁴⁶ | Number Of Chest X-rays (PA film) For Equivalent Effective Dose ⁴⁷ | Time Period For Equivalent Effective Dose From Natural Background Radiation ⁴⁸ |
|-----------------------------------|---|--|---|
| Chest X-ray (PA Film) | 0.02 | 1 | 2.4 days |
| Skull X-ray | 0.1 | 5 | 12 days |
| Lumbar Spine | 1.5 | 75 | 182 days |
| Intravenous (IV) Pyelogram | 3 | 150 | 1.0 year |
| Upper GI Exam (Barium Swallow) | 6 | 300 | 2.0 years |
| Lower GI Exam (Barium Enema) | 8 | 400 | 2.7 years |
| CT Head | 2 | 100 | 243 days |
| CT Abdomen | 8 | 400 | 2.7 years |

AS WE SEE IT

- 18. Michel F, Brevaut-Malaty V, Pasquali R, et al. Comparison of ultrasound and X-ray in determining the position of umbilical venous catheters. Resuscitation. Jun 2012:83(6):705-9.
- Bourcier JE, Paquet J, Seinger M, et al. Performance comparison of lung ultrasound and chest x-ray for the diagnosis of pneumonia in the ED. Am J Emerg Med. 2014 Feb;32(2):115-8.
- Available at: http://www.fda.gov/Radiation-EmittingProducts/RadiationEmittingProductsandProcedures/MedicalImaging/ucm200086.htm. Accessed September 25 2015
- 21. Available at: https://healthcarebluebook. com/. Accessed September 25, 2015.
- Available at: http://www.cancer.net/ navigating-cancer-care/diagnosing-cancer/ tests-and-procedures/integrated-pet-ctscan, Accessed September 29, 2015.
- 23. Gerth HU, Juergens KU, Dirksen U, Gerss J, Schober O, Franzius C. Significant benefit of multimodal imaging: PET/CT compared with PET alone in staging and follow-up of patients with Ewing tumors. J Nucl Med. 2007 Dec;48(12):1932-9.
- Kevies BH. Naked to the Bone: Medical Imaging in the Twentieth Century. New Brunswick, NJ: Rutgers University Press; 1997.
- 25. Available at: http://usatoday30.usatoday. com/news/nation/2002/02/28/usat-nuke. htm. Accessed September 25, 2015.
- 26. Available at: http://www.cnn.com/2002/ US/03/01/nuclear.fallout/index.html. Accessed September 25, 2015.
- 27. Available at: http://beforeitsnews.com/ alternative/2012/05/worst-scandal-inhistory-60-million-cancers-from-nuclearweapons-radiation-2136575.html. Accessed September 25, 2015.

- 28. Mangano JJ, Sherman JD. Elevated in vivo strontium-90 from nuclear weapons test fallout among cancer decedents: a case-control study of deciduous teeth. Int J Health Serv. 2011 41(1):137-58.
- Available at: http://www.lifeextension.com/ Magazine/2011/6/Optimize-Your-Internal-Defenses-Against-Radiation-Exposure/ Page-02. Accessed September 25, 2015.
- Available at: http://www.nobelprize.org/nobel_prizes/peace/laureates/1962/paulingbio.html. Accessed September 25, 2015.
- 31. Available at: https://www.newscientist. com/article/mg12416944.100-reviewfeet-of-clay--review-of-linus-pauling-a-man-and-his-science-by-anthonyserafini/. Accessed April 15, 2015.
- Available at: http://www.berkelev.edu/ news/media/releases/2007/09/04_GofmanObit.shtml. Accessed September 25, 2015.
- 33. Gofman JW JW, Lindgren F. The role of lipids and lipoproteins in atherosclerosis. Science. 1950 Feb 17;111(2877):166-71.
- Gofman JW. Serum lipoproteins and the evaluation of atherosclerosis. Ann NY Acad Sci. 1956 Nov 16;64(4):590-5.
- Gofman JW. Radiation-Induced Cancer from Low-Dose Exposure: An Independent Analysis. 1st ed. San Francisco, CA: Committee for Nuclear Responsibility;
- Gofman JW. Radiation from Medical Procedures in the Pathogenesis of Cancer and Ischemic Heart Disease: Dose-Response Studies with Physicians per 100,000 Population. 1st ed. San Francisco, CA: Committee for Nuclear Responsibility. 1999.
- Gofman JW, O'Connor E. Preventing Breast Cancer: the Story of a Major, Proven, Preventable Cause of this Disease. 2nd ed. San Francisco, CA: Committee for Nuclear Responsibility; 1996.

- 38. Claus EB, Calvocoressi L, Bondy ML, Schildkraut JM, Wiemels JL, Wrensch M. Dental x-rays and risk of meningioma. Cancer. 2012 Sep 15;118(18):4530-7.
- Available at: http://www.lifeextension.com/ Magazine/2011/6/Optimize-Your-Internal-Defenses-Against-Radiation-Exposure/ Page-01. Accessed September 25, 2015.
- Available at: http://www.lifeextension. com/Magazine/2015/4/Blueberries-Boost-Longevity-Beyond-Calorie-Restriction/ Page-01. Accessed September 25, 2015.
- 41. Available at: http://www.lifeextension.com/ magazine/2014/11/creating-immortalitygenes/page-01. Accessed September 25, 2015
- Byrne RT, Klingele AJ, Cabot EL, et al. Evolution of extreme resistance to ionizing radiation via genetic adaptation of DNA repair. Elife. 2014 Jan 1;3:e01322.
- 43. Brown W. From the Editor-in-Chief. J Clin Lipidol. 2007 May;1(2):97-9..
- Available at: http://www.ratical.org/radiation/CNR/synapse.html. Accessed April 20, 2015.
- Semendeferi I. Legitimating a nuclear critic: John Gofman, radiation safety, and cancer risks. Hist Stud Nat Sci. 2008 Spring;38(2):259-301.
- Mettler FA Jr, Huda W, Yoshizumi TT, Mahesh M. Effective doses in radiology and diagnostic nuclear medicine: a catalog. Radiology. 2008 Jul 248(1):254-63.
- Based on the assumption of an average "effective dose" from chest x ray (PA film) of 0.02 mSv.
- Based on the assumption of an average "effective dose" from natural background radiation of 3 mSv per year in the United



Fat-Soluble Nutrients Missing From Most Multi-Vitamin Formulas

Life Extension®'s **Health Booster** is a **cost-effective** formula that combines a variety of valuable nutrients in just one softgel. **Once-Daily Health Booster** provides the following nutrients:



Vitamin K1 is found in plants. It is often bound to plant fiber and requires intestinal conversion to transform into bioactive active **vitamin K2**.¹⁻³ Data supports value of K1 in addition to the K2 forms.4



Sesame lignans increases tissue levels of gamma tocopherol, which plays a pivotal role in quenching certain kinds of inflammation.14

Lycopene supports prostate health,





Vitamin K2 is the active form that keeps calcium in bone and out of arteries. MK-4 is rapidly absorbed, 1,8-9 while MK-7 provides 24-hour bioavailability of vitamin K2.9



guard against LDL oxidation. 15,16

Chlorophyllin offers protection against

environmentally induced DNA damage

protect against free radical activity, and



Trans-zeaxanthin, meso-zeaxanthin, and lutein supports eye health and healthy vision.



Critical Nutrients Not Found

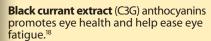
Most Multi-Vitamin Formulas

60 Softgels

from toxins like smoke, emission particles, and foods cooked at high temperatures.12



Gamma tocopherol is a form of vitamin E that quenches the damaging peroxynitrite free radical.¹⁰⁻¹¹ Those who take *alpha*-tocopherol should also take gamma tocopherol.







Blueberry extract boosts DNA function and sustain healthy blood sugar levels already within normal range. 12,1

Vitamin B12 helps maintain a healthy nervous system and metabolism.19 Vitamin B12 levels decrease with age.20



Each Bottle Of Health Booster Lasts Two Months

Super Cost Effective!

Just one softgel of the new **Health Booster** taken with a meal provides optimized potencies of fat-soluble vitamins, carotenoids, and other plant extracts. If these nutrients were taken separately, one would have to swallow many capsules and spend 2-3 times more money.

To order **Once-Daily** Health Booster, call **1-800-544-4440** or visit www.LifeExtension.com

Once-Daily Health Booster

Item #01981 • 60 softgels (two-month supply)

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$52 | \$39 |
| 4 bottles | | \$36 each |

One daily Health Booster softgel provides:

| Vitamin K1 | 1,000 mcg |
|-------------------------------------|-----------|
| Vitamin K2 (MK-4) | 1,000 mcg |
| Vitamin K2 (MK-7) | 200 mcg |
| Vitamin B12 | 300 mcg |
| Chlorophyllin | 100 mg |
| Gamma E Mixed Tocopherols | 359 mg |
| Gamma tocopherol 197.45 - 269.25 mg | • |
| Delta tocopherol 71.8 - 107.7 mg | • |
| Alpha tocopherol 30.52 - 43.08 mg | |
| Beta tocopherol < 17.95 mg | • |

| MacuGuard® Carotenoid Phospholipid Blend | 145 mg |
|---|--|
| Phospholipids, marigold extract (flower) | |
| [providing 10 mg free lutein, | |
| 4 mg meso-zeaxanthin & trans-zeaxanthin] | |
| C3G (Cyanidin-3-glucoside) | 2.2 mg |
| [from European black currant extract (fruit)] | ······································ |
| Lycopene (tomato extract) | 10 mg |
| Sesame Seed Lignan Extract | 20 mg |
| Wild Blueberry Whole Extract (fruit) | 100 mg |

- References
 1. Food Nutr Res. 2012;56.
- Food Nutr Res. 2012;56.

 J Biol Chem. 2008 Apr 25;283 (17):11270-9.

 J Nutr. 1998 128: 5 785-788.

 Am J Clin Nutr. 2012 Nov;96(5):1113-8.

 J Am Coll Nutr. 2009 Aug;28(4):369-79.

 Eur J Clin Nutr. 2005 Feb;59(2):196-204.

 J Nutr. 2014 May;144(5):743-50

 Int J Vitam Nutr Res. 1995;65(2):105-10.

 Nutr. 12012 Nov;12:11-01.

- 9. Nutr J. 2012 Nov 12;11:93 10.Nutr Rev. 1997 Oct;55(10):376-8. 11.Med Hypotheses. 2007;69(6):1367-70.
- 12.Int J Mol Sci. 2013;14(11):21447-62.
- 13. J Med Food. 2011 Dec;14(12):1511-8. 14. J Nutr. 2013 Jul;143(7):1067-73. 15. Nutr Cancer. 2009;61(6):775-83. 16. Lipids. 1998 Oct;33(10):981-4.
- 17. Toxicology. 2004 Mar 1;196(1-2):117-25. 18. J Biomed Biotechnol. 2004;2004(5):306-313.
- 19. Available at: http://www.ncbi.nlm.nih.gov/ pubmedhealth/PMHT0022013/. Accessed September 9, 2015. 20. Br J Haematol. 2010 Jan;148(2):195-204.

Contains soybeans.

Caution: if taking anticoagulant or antiplatelet medication, consult your health care provider before taking this product.

Tomat-O-Red® is a registered trademark of LycoRed, LTD. LuteinPlus® and Mz® are registered trademarks of NutriProducts Ltd., UK, licensed under U.S. Patent 8,623,428.



Blueberry Extract Boosts DNA Function

Studies show that blueberries delay the aging process through a variety of mechanisms, including **maintaining healthy DNA structure** and favorably modulating **genes** associated with aging. ¹²

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

- Enhance heart health³
- Maintain brain function^{4,5}
- Sustain healthy blood sugar levels already within normal range⁶
- Support smooth firm skin⁷
- Maintain a healthy weight⁸ and stable cholesterol levels already within normal range⁹

Blueberry extract is even more potent than the whole berry or juice, providing greater metabolic support throughout the entire body and without the excess sugar of raw fruit. Life Extension 's **Blueberry Extract Capsules** consists of only concentrated **extracts** from **wild blueberries**, which possess up to **10 times** the antioxidant capacity of cultivated berries.

Blueberry Extract Capsules

Item #01214 • 60 vegetarian capsules

LifeExtension

Blueberry

Extract

60 Vegetar

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

Suggested dose is <u>one</u> capsule daily, for most individuals.

References

- 1. Curr Pharm Des. 2013;19(34):6094-111.
- 2. Int J Mol Sci. 2013;14(11):21447-62.
- 3. PLoS One. 2009;4(6):e5954
- 4. Nutr Neurosci. 2005 Apr;8(2):111-20.
- Available at: http://www.scientificamerican.com/ article.cfm?id=your-brain-on-blueberries. Accessed January 28, 2015.
- 6. Georgian Med News. 2006 Dec;(141):66-72.
- 7. J Cosmet Dermatol. 2009 Jun;8(2):147-51.
- 8. Chem. 2010 Apr 14;58(7):3970-6.
- 9. J Agric Food Chem. 2005 May 4;53(9):3403-7.
- 10. J Agric Food Chem. 2010 Apr 14;58(7):3970-6.

Non-GMO

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

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



NSAIDs Increase Kidney Disease Risk

A study published in the journal Hypertension found that those with high blood pressure who regularly take nonsteroidal anti-inflammatory drugs

> (NSAIDs) have an elevated risk of developing chronic kidney disease.*

While some previous studies have linked NSAIDs to diminished kidney function, the outcome often ignored those who already had hypertension, which itself carries an increased risk for kidney damage.

After examining the data of over 30,000 patients with high blood pressure, researchers at the Institute of Population Health Sciences, National Health Research Institutes in Zhunan, Taiwan, found that those who'd been taking NSAIDs for at least three months were 32% more likely to have chronic kidney disease than those who didn't take NSAIDs. Those who used NSAIDs more than once a day had a 23% greater risk of developing chronic kidney disease than people who didn't.

Even taking NSAIDs for less than three months increased the risk of developing chronic kidney disease by 18%.

"Physicians should exercise caution when administering NSAIDs to people with hypertension and closely monitor renal function," said senior study author Hui-Ju Tsai.

Editor's Note: Taking NSAIDs may cause the kidneys to retain salt and water, increasing blood pressure and potentially making medications that lower hypertension ineffective.

* Hypertension. 2015 Sep;66(3):524-33.

Higher Vitamin D Levels And Lower Risk Of Colorectal Adenoma

The World Journal of Gastroenterology reported the results of a case-control study and meta-analysis which both found a lower risk of colorectal adenoma (polyps) in association with higher serum vitamin D levels.*

Researchers conducted a matched case-control study that included 112 Korean men and women with colorectal adenomas and 112 controls who underwent colonoscopy between August 2011 and September 2012. An association between higher vitamin D levels and a reduced risk of adenoma was found in women.

For the meta-analysis, 15 studies involving Western as well as Asian populations that examined serum or plasma vitamin D levels and the risk of colorectal adenoma were selected. The combined studies included 5,454 subjects with colorectal adenomas and 6,656 controls. When highest to lowest categories of vitamin D were compared, a 32% average reduction in the risk of adenoma was uncovered in the *highest* vitamin D group.

Editor's Note: While many adenomas remain benign, some are a precursor to colorectal cancer, one of the more common human malignancies.

* World J Gastroenterol. 2015 Aug 7;21(29): 8868-77.



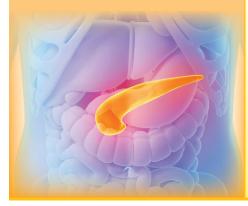
Metformin Plus Aspirin Inhibit Pancreatic Cancer

A recent study published in Oncotarget investigated the effects of metformin and aspirin, alone and in combination. on pancreatic cancer cell lines.* In a xenograft mouse model, the scientists demonstrated that the metformin/aspirin combination inhibited pancreatic tumor growth. For the study, randomized mice with tumors were injected with 200 mg/kg metformin, 60 mg/kg aspirin, or both three times a week for 28 days.

The researchers found that metformin combined with aspirin, at relatively low concentrations, demonstrated a synergistic effect on cell proliferation. Compared to each drug alone, the combination had significantly stronger effects on the inhibition of colony formation and cell migration, as well as the modulation of key molecular targets in AMPK and other signaling pathways. Furthermore the combination led to apoptosis through downregulation of anti-apoptotic proteins and the upregulation of pro-apoptotic proteins.

Editor's Note: "At the cellular level, metformin stimulates AMPK activation by disrupting mitochondrial respiratory chain complex I and decreasing ATP synthesis," say study authors. "Recently, aspirin was also shown to inhibit the dephosphorylation of AMPK thus activating ÂMPK.

* Oncotarget. 2015 Aug 28;6(25):21208-24.



Calorie Restriction Lowers Risk Of Age-Related Diseases

A two-year study supported by the National Institutes of Health found that calorie restriction lowered certain risk factors of age-related diseases.* In the study, published in the Journal of Gerontology, 218 healthy normal-weight and moderately overweight men and women were randomized to a reduced-calorie diet that was 25% below their normal calorie consumption. The calorie restriction group was given a weight-loss target of 15.5% in the first year and weight stability over the second year. Weight loss was expected to be achieved by reducing calorie intake 25% below their regular intake at baseline. The other participants maintained their regular baseline diets over the course of the study.

Although the weight loss by the calorie restriction group was the largest sustained weight loss reported in any clinical trial of nonobese participants, weight loss fell short of the target. The intervention arm only reached 12% caloric restriction instead of the trial's 25% goal but did maintain calorie restriction over the two-year period.

Calorie restriction significantly reduced several predictors of cardiovascular disease compared to the control group, including decreasing total cholesterol by 6%, increasing HDL levels, and lowering average blood pressure by 4%. Calorie restriction led to a 47% reduction in levels of **C-reactive protein** and markedly decreased insulin resistance.

Editor's Note: "It's important to find out whether these reductions would yield longterm benefits," said NIH director and paper author Dr. Evan Hadley. "It also would be useful to discover if calorie restriction over longer periods has additional effects on predictors of health in old age, and compare its effects with exercise-induced weight loss.

* J Gerontol A Biol Sci Med Sci. 2015 Sep;70(9):1097-104.

Nutrients Reduce Aging Of Thymus

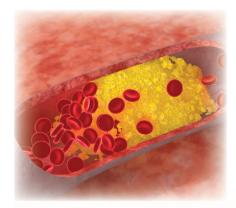
An article appearing in *Cell Reports* describes a role for nutrients in slowing aging of the thymus, a gland responsible for the production of immune cells known as T lymphocytes. The thymus reaches its peak size at adolescence and subsequently begins to atrophy.*

The researchers examined gene activity in the thymus' stromal (connective tissue) cells and lymphoid cells. They discovered that stromal cells were deficient in the body's enzyme catalase, making them subject to increased damage from reactive oxygen species.

To help confirm the benefit of this nutrient protection, mice were provided with drinking water enhanced with the nutrients N-acetylcysteine or vitamin C from the time of weaning. In comparison with mice that received plain water, thymus glands from mice that received either nutrient were larger after 10 weeks than those of control animals of the same age.

Editor's Note: In another experiment, mice that were genetically modified to overexpress mitochondrially targeted catalase had thymus glands that were significantly protected from thymus atrophy at six months of age compared to those of normal control animals.

* Cell Rep. 2015 Aug 18;12(7):1071-9.



Alpha-Lipoic Acid **Prevents Atherosclerosis**

An article published in Cell Reports reveals a protective effect for supplementation with alpha-lipoic acid on telomere length and vascular health in mice given a high-fat diet.*

"The effects of chronic diseases such as atherosclerosis and diabetes on blood vessels can be traced back to telomere shortening," noted senior author R. Wayne Alexander, MD, PhD.

Previous research, reported in the journal Circulation, found an inhibitory effect for lipoic acid against high-fat diet-induced atherosclerosis in mouse models of the disease. In the current report, Dr. Alexander and colleagues confirmed that alpha-lipoic acid stimulates peroxisome proliferator-activated receptor gamma co-activator 1-alpha (PGC1-alpha), which is known to control some aspects of how muscles respond to exercise. "What's new here is that we show that PGC1-alpha is regulating telomerase, and that has real beneficial effects on cellular stress in a mouse model of atherosclerosis," first author Shigin Xiong stated.

Editor's Note: Telomeres are segments of genetic material that cap and protect the ends of chromosomes. Aging-associated reduction in telomere length is considered a marker and cause of cellular aging. Telomere length is maintained by an enzyme known as telomerase. While telomerase is activated in proliferating cells, including those which are cancerous, Dr. Alexander's team has not observed an increased incidence of malignancies in animals that received alpha-lipoic acid.

* Cell Rep. 2015 Sep 1;12(9):1391-9.

Diet Alone Fails To Provide Vitamin E

The journal PLOS One published findings derived from NHANES data that reveal a high prevalence of suboptimal alpha-tocopherol (vitamin E) levels among subjects for whom food alone was their only source of the vitamin.*

Michael I. McBurnev and associates analyzed information from 7,922 NHANES participants with available measurements of serum alpha-tocopherol. A striking 87% of subjects aged 20 to 30 years and 43% of those aged 51 years and older had vitamin E levels lower than 30 micromoles per liter, which was categorized as inad-



equate based on Estimated Average Requirement and the individuals with the lowest mortality rates in the Alpha-Tocopherol Beta-Carotene study. Serum alpha-tocopherol levels among subjects whose only source of vitamin E was food were below adequate levels on average, at **24.9** micromoles per liter. In comparison, those whose vitamin E was derived from both food and supplements had an average level of 33.7 micromoles per liter.

Editor's Note: "We propose that for many Americans, especially those relying exclusively upon food sources, that serum alpha-tocopherol concentrations may not be adequate," Dr. Mc-Burney and colleagues write.

* PLoS One. 2015 Aug 19;10(8):e0135510.

Vitamin D Helps The Homebound

An article published on August 16, 2015, in the Journal of the American Geriatrics Society reports a successful outcome for home delivery of a vitamin **D** supplement to older individuals in Forsyth County, North Carolina.*

The study included 68 homebound Meals-on-Wheels participants. Subjects received a supplement providing 100,000 IU vitamin D or a placebo delivered with their meal once per month for five months. Serum 25-hydroxyvitamin D levels were assessed at the beginning and end of the study.

While 57% of subjects had serum vitamin D levels of less than 20 ng/mL prior to treatment, just one of 34 participants who received vitamin D3 had levels that were this low after five months, in comparison to 18 of the 34 participants randomized to the placebo group. After adjustment for several factors, a lower rate of falls was found among those randomized to vitamin D3 in comparison with the placebo group.

Editor's Note: Improved vitamin D status could aid in the prevention of falls that can be a complication of muscle weakness, osteoporosis, or impaired balance and/or vision that may occur in older men and women.

* J Am Geriatr Soc. 2015 Aug 16.





Symposium Highlights Coffee's Cardiovascular **Benefits**

Findings from studies presented at a Satellite Symposium on Coffee and Cardiovascular Disease Mortality held during the European Association for Cardiovascular Prevention And Rehabilitation's 2015 Congress add evidence to an association between drinking coffee and a reduced risk of cardiovascular disease and related mortality.*

Most notable were the results of a meta-analysis published in the American Journal of Epidemiology which concluded that, compared to not drinking coffee, consuming three cups per day was associated with up to a 21% reduction in the risk of dying from cardiovascular disease among 997,464 subjects. For all-cause mortality, the greatest protective effect was found in association with four cups daily. Another meta-analysis published in 2014, which appeared in Circulation, suggests that an optimal amount for protection against cardiovascular disease is three cups per day.

Coffee drinking has also been associated with protection against diabetes, a disease that significantly increases cardiovascular disease risk.

Editor's Note: Protective mechanisms for coffee against the risk of cardiovascular mortality remain uncertain; however, the beverage's antioxidant and anti-inflammatory effects are likely to

European Association for Cardiovascular Prevention & Rehabilitation's 2015 Congress. 2015 May 14.

Exciting Advances Regarding Metformin And Breast Cancer Prevention

Exciting advances in breast cancer prevention were made in 2015, with the focus on the diabetes drug metformin. 1-4 A clinical trial demonstrated that metformin can reduce proliferation of some types of breast cancer cells by over **60%**, compared with placebo.¹

There are three key pieces of evidence that explain why an antidiabetic drug would be an effective cancer preventive agent.

- 1. Insulin resistance increases risk for breast cancer and worsens prognosis. Metformin reduces insulin resistance. 5-7
- 2. Obese people are at higher risk for different types of cancers including breast cancer.8 Metformin fights obesity.9-11
- 3. High levels of insulin promote tumor formation. ¹² Metformin counteracts insulin resistance.5,10

And large-scale studies have shown substantial reduction in cancer rates and deaths (especially for breast cancer) among diabetics using metformin.¹³

One exciting finding is that metformin activates a "master signaling molecule" called AMPK.1 AMPK is involved and plays an important role in systemic energy balance, insulin signaling, and the metabolism of glucose and fats.1

Another of metformin's mechanisms is its unique ability to change cancer cells' metabolism. Studies show that this effect promotes killing of cancer stem cells in breast, prostate, colon, and brain cancers by producing an energy crisis in these rapidly dividing cells.

Metformin has been shown to induce cell death by energy starvation in breast cancer stem cells. This effect made the otherwise treatment-resistant stem cells highly vulnerable to standard chemotherapy.4 Chemotherapy works by inducing DNA damage in stem cells, and metformin blocks the normal DNA repair mechanisms by reducing available energy levels.4

While promising, these studies require validation from a controlled human trial. Such a study was recently conducted among a group of 200 women scheduled for surgery to remove invasive breast cancer lesions. Subjects were assigned to receive a placebo or metformin at 1,700 mg daily for four weeks preceding their surgery. At surgery, biopsies were taken to determine the effects of metformin. The researchers used a known marker of cancer cell proliferation, a protein called **Ki-67**, to detect evidence of cell replication, which would be evidence of cancerous progression.¹⁵

The study found that proliferation of cells in the precancerous lesions called *ductal carcinoma in situ* (DCIS) among metformin recipients was reduced compared with that in placebo patients in the group of women with several known markers of aggressive breast cancer including the cancer growth factor human epidermal growth factor receptor 2 (HER2), a marker of aggressive cancer seen in 20-30% of breast cancer victims).¹⁶ and the estrogen receptor molecule expressed on the precancerous cells.1

Women with DCIS and HER2 had a 40% reduction in cell proliferation, while those who had both HER2 and estrogen receptor markers had over **60%** reduction in proliferation.¹

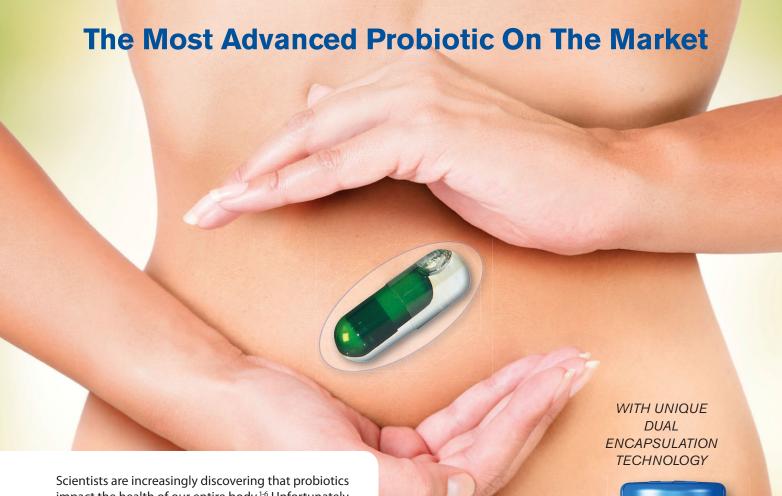
These most recent findings add to the evidence base that favors use of metformin in cancer-prevention efforts. The metformin story is a classic example of how closely related all of our bodies' metabolic systems are to one another, and how simple preventive approaches trump complex "designer drugs" aimed at fixing problems after they arise.

References

- Cancer Prev Res (Phila). 2015 Aug 14.

- Canter Frev Res (Frital), 2013 Aug 14.
 FEBS J. 2015 Aug 3.
 Prostate Cancer Prostatic Dis. 2015 Jul 28.
 Apoptosis. 2015 Oct;20(10):1373-87.
 Front Neurosci. 2015 Jul 14;9:235.
 World J Hepatol. 2015 Jun 28;7(12):1652-9.
 J Clin Oncol. 2011 Jan 1;29(1):7-10. Lancet. 2008 Feb 16;371(9612):569-78.
- 9. Obes Facts. 2012;5(5):753-65

- 7. Oves racts. 2012;3(5):755-65.
 10. Diabetes Care. 2009 Sep;32(9):1743-5.
 11. Hum Reprod Update. 2015 Sep;21(5):560-74.
 12. Cancer Discov. 2012 Sep;2(9):778-90.
 13. Cancer Prev Res (Phila). 2010 Nov;3(11):1451-61.
 14. Circ Res. 2007 Feb 16;100(3):328-41.
 15. J Cell Physiol. 2000 Mar;182(3):311-22.
- 16. Chemother Res Pract. 2012;2012:743193.



Scientists are increasingly discovering that probiotics impact the health of our entire body. ¹⁻⁶ Unfortunately, most commercial probiotics are destroyed by the stomach's natural digestive acids <u>before</u> they reach their destination.⁷

FlorAssist® Probiotic Liquid Vegetarian Capsules with "dual encapsulation" technology delivers maximum probiotic protection to your small intestines.

FlorAssist® Probiotic Liquid Vegetarian Capsules:

- Contain probiotic strains that are stomach acid resistant
- Have dual encapsulation technology, which keeps the capsule intact longer and ensures that the probiotic reaches the small intestine
- Provide 15 billion CFU—Colony Forming Units per capsule
- Contain 6 varieties of beneficial bacteria

FlorAssist® Probiotic

Item #01825 • 30 liquid veg. capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$32 | \$24 |
| 4 bottles | | \$21 each |

Non-GMO



To order FlorAssist® Probiotic Liquid Vegetarian Capsules, call 1-800-544-4440 or visit www.LifeExtension.com

FlorAssist™ contains the following bacterial strains:

- 1. Lactobacillus acidophilus LA-14 2. Lactobacillus rhamnosus LR-32
- 3. Lactobacillus paracasei LPC-37 4. Bifidobacterium longum BL-05
- 5. Bifidobacterium lactis BL-04 6. Bifidobacterium bifidum/lactis BB-02

These potent strains of probiotic bacteria adhere to the soft lining of the intestinal tract and help maintain a healthy surface and aid in support for the digestive system.

References

- 1. Eur J Clin Nutr. 2013 Feb;67(2):161-7.
- 2. Curr Top Microbiol Immunol. 2013;358:273-89.
- 3. Br J Nutr. 2013 May 28;109(10):1866-72.
- 4. Nutr Hosp. 2011 Jan-Feb;26(1):228-35.
- 5. Fur I Cancer Prev. 2013 Jan:22(1):46-51.
- 6. Pediatr Int. 2012 Oct;54(5):682-7
- 7. Microbiology. 2007 Oct;153(Pt 10):3563-71.

Support Healthy Metabolic Factors

Omega-7

With Highly Purified Provinal® Fish Oil

As a Life Extension® customer, you most likely take an omega-3 supplement. Scientists are increasingly excited about the newly discovered effects of another family of beneficial fats—omega-7 fatty acids.

Omega-7 palmitoleic acid powerfully complements omega-3 fatty acids with cellular-signaling properties that support healthy metabolic factors associated with:

- Superior cardiovascular health^{1,2}
- A healthy inflammatory response^{1,3,4}
- Optimum normal metabolism of glucose and insulin1,5,6,
- Beneficial management of hunger and satiety^{7,8}

Scientific studies show that palmitoleic acid promotes healthy levels of triglycerides, total cholesterol, LDL, and **HDL** for those already in normal range—after just one month of supplementation at 210 mg.9

Further studies have found within normal range, palmitoleic acid can help balance C-reactive protein levels, 10 optimize insulin sensitivity, 1,10,11 and regulate hunger-promoting gastric hormones.^{7,8}

Life Extension®'s omega-7 supplement contains **Provinal**®—a highly refined anchovy and/or menhaden oil that is **non-GMO**. Conventional processing methods result in products with only 25% palmitoleic acid. Life Extension®'s Provinal® Purified Omega-7 is concentrated to 50% palmitoleic acid.



References

- Lipids Health Dis. 2011;10:120.
- Biosci Biotechnol Biochem. 2011;75(12):2401-3.
- PLoS One. 2012;7(6):e39286.
- Eur J Nutr. 2011 Aug;50(5):323-30.
- 5. Biochem J. 2006 Nov1;399(3):473-81. 9. Effect of Two Dosage Level of Provi- 10. J Clin Endocrinol Metab. 2013
- 6. Diabetes. 2001 Jan;50(1):69-76. Appetite. 2013 Jun;65:1-7.
- 8. Am J Physiol Gastrointest Liver Physiol. 2012 Aug 1;303(3):G367-76.
- nal® on serum lipid and c-reactive protein profiles in humans: Tersus Pharmaceuticals, LLC

Item #01812 • 30 softgels

1 bottle

4 bottles

Retail

Price

\$27

Your

Price

\$20.25

\$18 each

- Jan:98(1):E40-50.
- 11. Diabetes Care. 2010 Feb;33(2):405-7.

Provinal

urified Omega-7

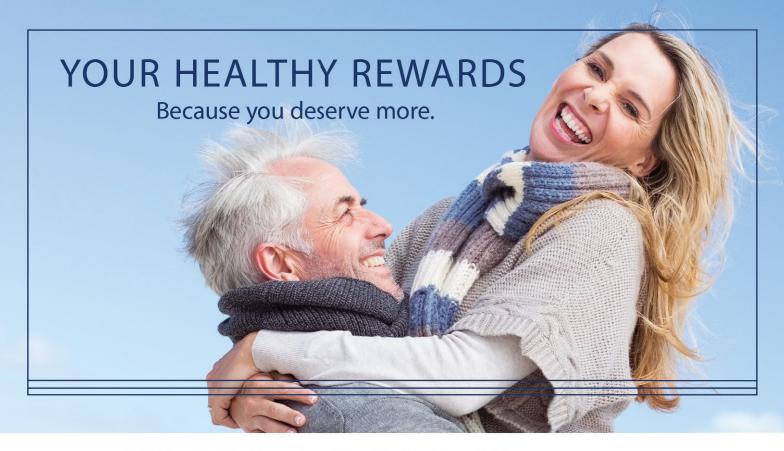
m Highly Purified Fish Oil

Healthy Heart and

Inflammatory Response Support

30 Softgels

To order Provinal® Purified Omega-7, call 1-800-544-4440 or visit www.LifeExtension.com



With Your Healthy Rewards and Your Healthy Rewards *Premier*, everyone earns LE Dollars back on nearly everything you buy ... and don't forget: Life Extension® members earn **DOUBLE LE Dollars back** from September 1, 2015 until the end of their existing LE membership!

Earn 2% LE Dollars Earn 2% LE Dollars back on every product or blood test you

buy from Life Extension. (4% for current LE members)*

Exclusive LE Perks Enjoy 25% – 60% off retail prices, complimentary access to

expert Health Advisors, and a FREE monthly subscription to

Life Extension Magazine.**

Want to earn even more rewards?

Join Your Healthy Rewards *Premier* and get a \$50 LE Dollar sign up bonus, earn DOUBLE LE Dollars (4%) back, plus a FREE

year of **CHOICE** unlimited shipping (a \$19.95 value)![†]

With a low annual fee of just \$49.95, **Your Healthy Rewards** *Premier* more than pays for itself. Upgrade today!

(\$59.95 for international customers)

(\$39.93 for international customers)

To learn more about Your Healthy Rewards, call toll-free

1-888-224-8239 • www.LifeExtension.com/Rewards
Mention Code YRH537A







A Novel Probiotic To Reduce The Risk Of STREP THROAT

Strep throat (known to physicians as **streptococcal pharyngitis**) is caused by a group of bacteria known as type A streptococcus. Responsible for up to about **15%** of sore throat infections in adults and up to about 30% in children, strep throat is characterized commonly by fever, painful swallowing, and tender, enlarged lymph nodes in the neck.1

Although most of the time in adults the cause of a sore throat is viral in origin, and therefore antibiotics are of no use, in the case of confirmed strep throat, antibiotic treatment is important to prevent dangerous consequences, like rheumatic heart disease or retropharyngeal abscess.1,2

At a time when antibiotics were not available, many people suffered as a consequence of inadequate treatment of strep throat.

While antibiotic treatment of confirmed **strep throat** is important to prevent the potential for rheumatic heart disease and other complications, an opportunity exists to reduce the risk of strep infection in the first place.

The intriguing news is that scientists have identified a **novel probiotic**, known as **S.** salivarius K12, and studies suggest that this targeted probiotic lozenge may reduce the risk of strep throat. In addition, this probiotic also appears capable of reducing the incidence of non-strep, viral sore throats as well.

Sore Throats: Significant Adult Problems

There are few "minor" illnesses as irritating as a sore throat.

Unfortunately, there's no good way to discern whether an infectious sore throat is caused by a bacteria or virus based solely on clinical symptoms in isolation. Sore throat caused by a potentially dangerous organism like Streptococcus pyogenes (strep throat), however, tends to be more severe, and accompanied by (high) fever, but not always.1

In the majority of cases a sore throat is caused by a virus.3

Antibiotics are of no use in viral infections. Unfortunately, overprescribing antibiotic treatment in the face of any sore throat continues, though to a lesser degree over the past several years since most physicians and patients are aware of the unnecessary problems antibiotics cause.4

Bacterial infection with pathogenic organisms like Streptococcus pyogenes is the only real indication for antibiotic use. This particular organism produces a painful sore throat ("strep throat") that can last three to five days, and can also cause long-term complications such as rheumatic heart disease, retropharyngeal abscess, and post-streptococcal glomerulonephritis (kidney disease).^{1,2} These conditions are rare today in comparison with the era prior to antibiotics, but concerns remain high, with overprescribing of antibiotics a consequence.3

Overprescribed antibiotics are a known public health problem, particularly in adults with upper respiratory tract infections such as sore throat.^{5,6} These practices contribute to the emergence of "superbugs" that are difficult to treat and have the potential to

produce dangerous infections.⁷⁻⁹ And indiscriminate antibiotic use also creates problems by producing an imbalance of the gastrointestinal (GI) microbiome, the population of healthy organisms that lives in our GI tract and has tremendous impact on overall health.¹⁰

And because it is impossible to tell at home if a sore throat is caused by strep, many people with sore throats visit their physicians, and research suggests that potentially up to 75% of adult patients receive antibiotic treatment for what ultimately turns out to be nonbacterial in origin.3

What this all adds up to is that we need a way to reduce the risk of any kind of sore throat in adults. That will help reduce the incidence of genuine, antibioticrequiring cases of sore throat caused by bacteria, such as strep throat, and also minimize the number of physician visits for sore throat that might result in inappropriate antibiotic prescribing.

And that's where the beneficial, probiotic **S.** salivarius K12 comes in, as we'll now see.

Novel Probiotic Fights Strep

There is a new method for fighting and reducing the risk of strep throat infection.

The beneficial S. salivarius probiotic strain was discovered in the 1980s in cultures from individuals who seemed resistant to developing sore throats. 12,13 That search yielded a specific strain of S. salivarius called **K12**, which is found in only about **2**% of individuals who show a natural resistance to strep throat.¹⁴ With sufficient numbers of this bacterial strain, it can keep disease-causing organisms such as Streptococcus pyogenes under control. 12,15





One of its key properties is that S. salivarius K12 produces compounds called *lantibiotics*. ¹⁶ These lantibiotics strongly inhibit many strains of disease-producing *Streptococcus pyogenes* responsible for causing strep throat in humans. 15 As a result, disease-causing organisms may be prevented.

Salivaricin A2, a type A lantibiotic produced by S. salivarius K12, works like a drill that forms molecular pores in the membranes of target organisms, permitting them to leak out their contents into surrounding fluid and then break apart and die.15-19

But that's not where the benefits of *S. salivarius* K12 end. While the mechanisms are still being investigated, there is now evidence that S. salivarius K12 colonization of the throat reduces secretion of the inflammatory signaling molecule (cytokine) IL-8, and increases concentrations of the antiviral compound interferon-gamma. 18 These properties may account for observation that children treated with S. salivarius K12 had a significant reduction in both viral and bacterial sore throats.18

Clinical Evidence

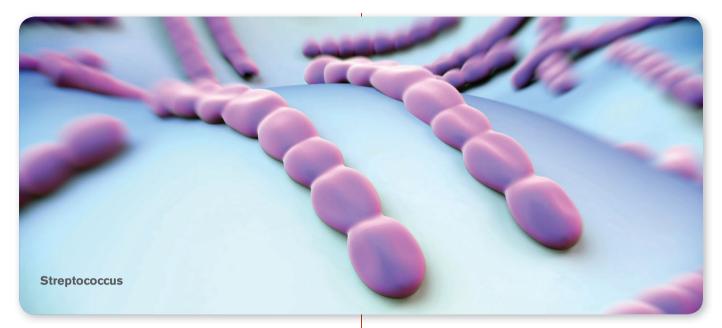
A growing body of evidence demonstrates the effectiveness of the probiotic S. salivarius K12 in helping to reduce the risk of sore throats of all kinds.

A randomized, controlled clinical trial was performed among a group of adults to determine the effects of S. salivarius K12 oral therapy on the number and severity of episodes of proven strep throat. Subjects were people with a history of recurrent strep throat or tonsillitis (four or more proven episodes in the preceding year).²¹

Subjects were free of symptoms at enrollment in the study. They were started on either a daily lozenge containing S. salivarius K12 organisms, or given no treatment. Treatment was given daily for 90 days. Subjects were then followed for an additional six months with no further treatment.

Stop Sore Throats Without Dangerous Drugs

- Sore throats are a major problem in adults as well as children.
- Even strep throat, considered a "childhood illness," is known to occur in adults, though less frequently.
- While most sore throats are caused by viruses and are merely uncomfortable, strep throat can produce dangerous complications like rheumatic heart disease, retropharyngeal abscess, and kidney disorders.
- Instead of waiting for a sore throat to develop and taking your chances with or without antibiotics, you can proactively work to prevent sore throats of all kinds, including potentially dangerous strep.
- Probiotic bacteria S. salivarius K12 produces potent, locally acting lantibiotics, compounds that target the organisms that cause strep throat.
- Clinical studies demonstrate a significant reduction in strep throat infections in people supplementing with this probiotic in the form of a daily oral lozenge.
- Remarkably, studies also show that S. salivarius K12 contributes to a reduction in viral sore throats as well. This property is attributed to the probiotic's impact upon certain signaling molecules (cytokines), including interferongamma.
- Regular use of probiotic S. salivarius K12 may help you dodge the upcoming winter sore throat season.



Treated adults who completed the 90-day course of supplementation had a significant **84**% <u>reduction</u> in their incidence of strep throat or tonsillitis compared with the previous year.

Even during the six-month, no-treatment follow-up period, patients who underwent the original 90-day supplement experienced a significant **62**% reduction in episodes of strep throat or tonsillitis, compared with the untreated group.

This study therefore demonstrated that *preventive* treatment with *S. salivarius* K12 was effective at significantly reducing the rates of recurrent strep disease. That's a first.

In a similar study targeting school-age children with recurrent strep infections, treatment with *S. salivarius* K12 was given for 90 days.²² Again, the subjects were followed for six additional months after the supplementation period.

At the 90-day mark, supplemented children demonstrated a **92**% reduction in strep throat, compared with the previous year. Again, there was a significant reduction (this time **66**%) in the rate of strep throat infections in the six months following the end of the supplementation period.²²

In an important additional finding, the rate of acute ear infections in this group of children was also significantly reduced by about **40**% in the supplemented group, compared with the previous year.²¹ This suggests that not only *Streptococcus pyogenes*, but also potentially other disease-causing organisms, are susceptible to this preventive treatment.

In a third study, children with recurrent strep throat were supplemented for 90 days with *S. salivarius* K12 (1 billion organisms per lozenge), or given no therapy.¹⁸

Supplemented children showed a significant **97%** reduction in strep throats, from a baseline of 3.1 infections per child in the year prior to the study to just 0.1 infection per child in the year of the study, with no significant change seen in the unsupplemented group.¹⁸

Remarkably, the supplemented patients also had a significant (and unprecedented) **80**% decrease in the incidence of oral and throat *viral* infections. The researchers speculated that this reduction might be related to a known ability of *S. salivarius* K12 to reduce inflammatory **IL-8** and increase anti-viral **interferongamma**. ^{18,23}

Summary

Sore throats affect most everyone, causing loss of work, sleep, and quality of life.

Strep throats caused by pathogenic type A streptococcus bacteria strains can strike both children and adults.

The novel probiotic supplement *S. salivarius* **K12** now available in a once-daily lozenge, produces potent, natural, locally acting *lantibiotics*, a class of compounds that may reduce the risk of throat infection.

S. salivarius K12 has now been clinically documented to reduce the incidence of **strep throat** infections in both adults and children. In an intriguing finding, people who supplemented with this probiotic also appeared to be protected against **viral** throat infections as well.

Viral infections are the leading cause of sore throats, and although viruses are unfazed by antibiotics, antibiotic treatment is often prescribed, contributing to their inappropriate use. •

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

References

- Shulman ST, Bisno AL, Clegg HW, et al. Clinical practice guideline for the diagnosis and management of group A streptococcal pharyngitis: 2012 update by the Infectious Diseases Society of America. Clinical infectious diseases: an official publication of the Infectious Diseases Society of America. Clin Infect Dis. 2012 Nov 15;55(10):e86-102.
- Karmarkar MG, Hule GP, Cameron A, et al. Antibodies to group A streptococcal virulence factors, SIC and DRS, increase predilection to GAS pyoderma. BMC Infect Dis. 2015;15:113.
- Perone N, Humair JP. Diagnosis and management of pharyngitis. Rev Med Suisse. 2007 Jan 31;3(96):286-90.
- West JV. Acute upper airway infections. Br Med Bull. 2002;61: 215-30.
- Dekker AR, Verheij TJ, van der Velden AW. Inappropriate antibiotic prescription for respiratory tract indications: most prominent in adult patients. Fam Pract. 2015 Aug;32(4):401-7.
- Gulliford MC, Dregan A, Moore MV, et al. Continued high rates of antibiotic prescribing to adults with respiratory tract infection: survey of 568 UK general practices. BMJ Open. 2014;4(10):e006245.
- Special Report. The rise of superbugs. Consum Rep. 2015 Aug:80(8):20-6.
- Blondeau J. Gram-negative superbugs: inappropriate antimicrobial therapy and mortality. Expert Rev Clin Pharmacol. 2013 Jul;6(4):347-9.
- Dawson MP, Smith AJ. Superbugs and the dentist: an update. Dent Update. 2006 May;33(4):198-200, 02-4, 07-8.
- 10. Modi SR, Collins JJ, Relman DA. Antibiotics and the gut microbiota. J Clin Invest. 2014 Oct;124(10):4212-8.
- 11. Available at: http://www.niaid.nih.gov/topics/streptococcal/Pages/ Default.aspx. Accessed September 21, 2015.

- 12. Fantinato V, Jorge AOC, Shimizu MT. Production of bacteriocinlike inhibitory substances (BLIS) by Streptococcus salivarius strains isolated from the tongue and throat of children with and without sore throat. Revista de Microbiologia. 1999;22(30):332-34.
- 13. Dempster RP, Tagg JR. The production of bacteriocin-like substances by the oral bacterium Streptococcus salivarius. Arch Oral Biol. 1982:27(2):151-7.
- 14. Burton J, Chilcott C, Wescombe P, et al. Extended safety for the oral cavity probiotic streptococcus salivarius K12. Probiotics & Antimicro. Prot. 2010;2:135-44.
- 15. Wescombe PA, Upton M, Dierksen KP, et al. Production of the lantibiotic salivaricin A and its variants by oral streptococci and use of a specific induction assay to detect their presence in human saliva. Appl Environ Microbiol. 2006 Feb;72(2):1459-66.
- 16. Hyink O, Wescombe PA, Upton M, et al. Salivaricin A2 and the novel lantibiotic salivaricin B are encoded at adjacent loci on a 190-kilobase transmissible megaplasmid in the oral probiotic strain Streptococcus salivarius K12. Appl Environ Microbiol. 2007 Feb;73(4):1107-13.
- 17. Walls T, Power D, Tagg J. Bacteriocin-like inhibitory substance (BLIS) production by the normal flora of the nasopharvnx: potential to protect against otitis media? J Med Microbiol. 2003 Sep;52(Pt 9):829-33.
- 18. Di Pierro F, Colombo M, Zanvit A, et al. Use of Streptococcus salivarius K12 in the prevention of streptococcal and viral pharvngotonsillitis in children. Drug Healthc Patient Saf. 2014;6:15-20.
- 19. Gillor O, Etzion A, Riley MA. The dual role of bacteriocins as antiand probiotics. Appl Microbiol Biotechnol. 2008;81(4):591-606.
- 20. Draper LA, Cotter PD, Hill C, et al. Lantibiotic resistance. Microbiol Mol Biol Rev. 2015 Jun;79(2):171-91.
- 21. Di Pierro F, Adami T, Rapacioli G, et al. Clinical evaluation of the oral probiotic Streptococcus salivarius K12 in the prevention of recurrent pharyngitis and/or tonsillitis caused by Streptococcus pyogenes in adults. Expert Opin Biol Ther. 2013 Mar;13(3):339-43.
- 22. Di Pierro F, Donato G, Fomia F, et al. Preliminary pediatric clinical evaluation of the oral probiotic Streptococcus salivarius K12 in preventing recurrent pharyngitis and/or tonsillitis caused by Streptococcus pyogenes and recurrent acute otitis media. Int J Gen Med. 2012;5:991-7.
- 23. Wescombe PA, Hale JD, Heng NC, et al. Developing oral probiotics from Streptococcus salivarius. Future Microbiol. 2012 Dec;7(12):1355-71.





LIFE EXTENSION'S BEST-SELLING **DHEA** PRODUCT!

DHEA

Supports Total-Body Health

DHEA is a critically important hormone, but its production declines sharply as we age. By the time you reach 70, your DHEA levels are likely to be **75-80%** *lower* than when you were at your peak.¹⁻⁴

Scientists are discovering numerous health benefits when aging people restore their **DHEA** to youthful ranges. **DHEA** therapy has been shown to:

- Support healthy arterial structure and function^{5,6}
- Support endothelial health by helping with blood flow⁷
- Promote insulin sensitivity8,9
- Benefit the normal aging brain 10-12
- Improve mood and alleviate melancholy¹³
- Protect hip bone and spine bone mineral density¹⁴
- Enhance the increases in muscle mass and strength in the elderly with resistance exercise¹⁵
- Boost a broad array of immune system cells and signaling molecules¹⁶

By supplementing with DHEA, you can get your levels back to youthful values. An ideal daily dose of **DHEA** for many healthy aging people is **25 mg**. Life Extension's®

convenient, economical **25 mg** capsules are a popular way to consume the **precise amount** of DHEA your body may need.



DHEA 25 mg

Item #00335 • 100 capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$18 | \$13.50 |
| 4 bottles | | \$11.25 each |



Each bottle lasts a typical user over three months!

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

CAUTION: Do not use DHEA if you are at risk for or have been diagnosed as having any type of hormonal cancer, such as prostate or breast cancer.

Non-GMO

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.

SUPPORT THROAT HEALTH WITH NOVEL PROBIOTIC LOZENGE

FlorAssist® Throat Health is an oral probiotic that provides novel beneficial bacteria to colonize the throat and is clinically shown to support throat health.¹⁻³

Just one lozenge of FlorAssist® Throat Health contains 20 mg of BLIS K12®, a proprietary oral probiotic known as *S. salivarius* K12, which delivers 2 billion colony-forming units. These organisms survive naturally in the throat, maximizing their potential to promote throat health by helping to regulate inflammation and reduce the damage caused by germs that originate there.

FlorAssist® Throat Health—naturally flavored with spearmint and cherry—can significantly protect throat health, ¹⁻³ ease inflammation, and help maintain good health. ⁴ And because these unique compounds act locally in the throat, clinical study participants supplementing with *S. salivarius* K12 have reported excellent tolerability without systemic side effects. ⁵

Contains milk.

BLIS K12° is the registered trademark of BLIS Technologies Limited.

References

- 1. Oral Microbiol Immunol. 2009 Apr;24(2):152-61.
- 2. Appl Environ Microbiol. 2006 Feb;72(2):1459-66.
- 3. Revista de Microbiologia. 1999;22(30):332-4.
- 4. Drug Healthc Patient Saf. 2014;6:15-20.
- 5. *Microbiol Mol Biol Rev.* 2015 Jun;79(2):171-91.

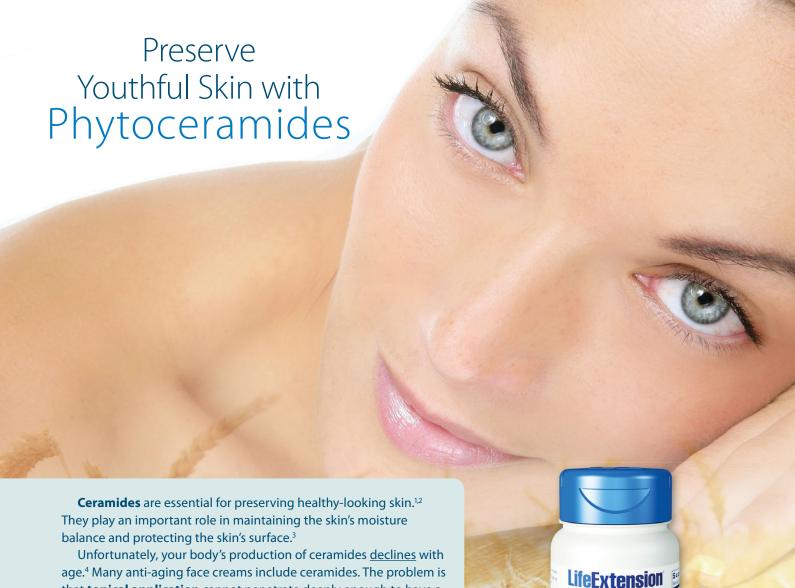
FlorAssist® Throat Health

Item #01920 • 30 lozenges

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$20 | \$15.00 |
| 4 bottles | | \$13.50 each |



To order FlorAssist® Throat Health, call 1-800-544-4440 or visit www.LifeExtension.com



age.4 Many anti-aging face creams include ceramides. The problem is that topical application cannot penetrate deeply enough to have a long-term impact on your skin's appearance.

Restoring Youthful Ceramide Levels

Researchers have discovered that the ceramides naturally produced by young skin are identical to those present in wheat and that these wheat-derived oils can be taken orally.

Skin Restoring Phytoceramides with Lipowheat® can reach the deepest layers of skin all over the body—not just where creams are applied—where it can offset the visible impact of the body's gradual decline in ceramides. The hydrating action of Lipowheat® ceramides has proved effective in clinical trials.

Just One Capsule Daily

Life Extension® has brought together these skin-nourishing oils in a concentrated oral formula called Skin Restoring Phytoceramides with Lipowheat®.

References

- 1. Biophys Chem. 2010 Aug;150(1-3):144-56.
- 2. Chemistry and Physics of Lipids. Apr 2007; 146(2):67-75.
- 4. Baran R, Maibach H, eds. Textbook of Cosmetic Dermatology. 3rd ed. Taylor & Francis:2005:177

3. Int J Cosmet Sci. 2010 July 14.

Non-GMO

Contains wheat. Gluten-free.

Lipowheat® is a registered trademark of Arco, Robertet Group, France.



Skin Restoring

Phytoceramides

with Lipowheat®

Helps Maintain Healthy Skin Hydration

Item #01596 • 30 liquid vegetarian capsules

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

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

Healthy Holiday

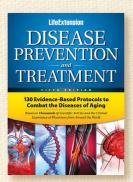
BOOK SALE

Great holiday gifts Provide lifesaving information for vour loved ones.

Powerhouse titles by knowledgeable doctors, best-selling authors, and cultural icons.

> Order by December 18, 2015 to receive in time for the holidays!

SAVE UP TO 80%!



Item #DPT05

≈ SPECIAL OFFER ≈

Disease Prevention and Treatment — Updated 5th Edition Your ultimate medical reference ... only \$19.99!

Featuring 1,400 pages of breakthrough information that bridges the gap between cutting-edge science and mainstream medicine, addressing 130 different health concerns. A \$69.95 value!

 When you order four or more copies of Disease Prevention and Treatment, the price drops to \$15 per book.

∞OTHER TITLES ≈



The Supplement Pyramid by Michael A. Smith, M.D.

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| \$24.95 | \$7.99 | 67% |

Item #33864

| Retail Price | Sale Price | Save |
|--------------|------------|------|
| \$24.95 | \$7.99 | 67% |

Healthy Skin

Your Guide to Healthy Skin the Natural Way

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| | | |

Item #33838

by Gary Goldfaden, M.D.

| il Price | Sale Price | Save | |
|----------|------------|------|--|
| 26.00 | \$3.99 | 84% | |



Heart: An American Medical Odyssey by Dick Cheney and Jonathan Reiner, M.D.

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| \$28.00 | \$13.99 | 50% |



Item #33670

A Primer on **Prostate Cancer** S.B. Strum, M.D., and D. Pogliano

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| \$28.95 | \$11.49 | 60% |



Cheating Death by Sanjay Gupta, M.D.

| Retail Price | Sale Price | Save |
|--------------|------------|------|
| \$24.99 | \$12.49 | 50% |

Item #33814



Pharmocracy by William Faloon

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| \$24.00 | \$4.99 | 79% |

Item #33835

∞MORE ON SALE ≈

Popular titles by **Suzanne Somers**

Written by the pioneering health and wellness advocate, and New York Times best-selling author, these books make perfect gifts for anyone trying to stay healthy, avoid the pitfalls of perimenopause, or just fight fat over 40!



I'm Too Young for This! by Suzanne Somers

| Retail Price | Sale Price | Save |
|--------------|------------|------|
| \$26.00 | \$14.29 | 45% |

Item #33862



Item #33830

Sexy Forever: How to Fight Fat after Forty by Suzanne Somers

| Retail Price | Sale Price | Save |
|---------------------|------------|------|
| \$25.00 | ¢12.00 | 460/ |



Knockout: Interviews with Doctors Who Are **Curing Cancer**

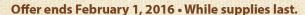
by Suzanne Somers

| Retail Price | Sale Price | Save |
|--------------|------------|------|
| \$25.99 | \$13.99 | 46% |

EVEN MORE BEST SELLER TITLES ~ AVAILABLE HERE ~

www.LifeExtension.com/BookSale Or call toll-free 1-866-626-7455 to order

You must mention marketing code AVK503A to get these special savings.







Block Deadly Effects Of After-Meal Blood Sugar

Diabetics and prediabetics are prone to accelerated aging due to the systemic damage caused by excess blood sugar. Unfortunately, as humans age, blood sugar tends to rise. What researchers are discovering is that people don't have to be a diabetic to suffer from the extensive health consequences of high blood sugar. ²⁻⁹

The problem is that after a meal, one's blood sugar (whether a person is diabetic or not) spikes and the result is the production of deadly byproducts called *advanced glycation end products* (AGEs).^{10,11} **AGEs** are rogue molecules that damage cells, tissues, and organs.¹¹⁻¹³

They also trigger inflammation and oxidation, generating even more of these molecules in a vicious, health-robbing cycle.¹⁴

Life Extension® has long advised customers about the importance of maintaining a fasting glucose level between **70** to **85 mg/dL**.

Compelling data now confirms that a unique form of vitamin B1 called **benfotiamine** can protect against the ravages of elevated blood sugar.^{15,16}

In the process of helping neutralize the adverse impact of elevated after-meal blood glucose, **benfotiamine** can help prevent Alzheimer's disease, vision impairment, cardiovascular disease, kidney damage—and possibly even aging itself!⁶

Blood Sugar And Tissue Damage

The browning that takes place when foods are cooked at high temperatures (known as the Maillard **Reaction**) is the result of a sugar-related molecular change known as *glycation*. This damaging process takes place when a reducing sugar and an amino acid react in the presence of heat. 17,18 Put simply, the structure of normal proteins becomes warped.

This same dysfunctional chemical change is occurring every day *inside* the body.

Within every cell, the sugar that the body uses for fuel is busy at work at every moment reacting with proteins and fats. 17,18 The byproducts of this chemical reaction⁴⁻⁷ are called *advanced glycation end products*.¹⁹

Ultimately, these *advanced glycation end products* deactivate enzymes, disrupt cell signaling, and damage the body's DNA. This results in intracellular damage and apoptosis (programmed cell death),20 inflammation, 9,21 oxidative stress, 9,22-24 nerve-cell component damage.²⁵ a depletion of nitric oxide^{24,26} (which is vital to vascular health), and high cholesterol.28

These **effects** are implicated in the progression of numerous age-related diseases,28 ranging from neuropathy23 and vision problems29 to kidney30 and cardiovascular disease^{24,27,31}—to *even aging itself*.^{2,32}

With advancing age, even **normal** blood sugar levels eventually trigger deadly advanced glycation end products. Studies clearly demonstrate links between major health risks and levels of AGEs in the blood of nondiabetics.33,34

Scientists have speculated that AGE-related changes ultimately contribute to a limitation of the human life span³⁵—suggesting that blocking the pathways of glycation could extend human life expectancy.



Supercharge The Body's Natural Defenses

One of the natural defenses against advanced glycation end products is vitamin B1, known as thia**mine**. Vitamin B1 prevents glycation molecules from becoming fully formed in the first place. 22,36

The problem is that glucose quickly depletes the body's limited thiamine supplies.³⁷ And although it might seem logical to use thiamine supplements to block advanced glycation end products, that's problematic as well.

Because thiamine is water soluble, it has difficulty penetrating the lipid molecules that primarily make up cell membranes. Also, the body has a very *low* upperdose limit, which means that taking larger doses of thiamin wouldn't do any good because it would get excreted in the urine. 38,39 Ultimately, tissues simply can't absorb nearly enough thiamine to prevent cumulative damage caused by advanced glycation end products. 40,41

Researchers realized there was an urgent need for a better means of delivering thiamine to the cellsand that's where the unique form of thiamine called **benfotiamine** comes in. The major difference is that **benfotiamine** is a *fat-soluble* derivative of thiamine. Because it is fat soluble, benfotiamine substantially boosts thiamine absorption into cells and throughout the body. 15,40,41

Administration of benfotiamine resulted in a 10 to **40% higher** thiamine incorporation into the liver and heart—and a remarkable 5- to 25-fold higher thiamine incorporation into muscle and brain!³⁹ A small clinical study showed that this enhanced bioavailability of benfotiamine caused it to slash the intracellular formation of AGEs in subjects' blood cells by 40%!42

Benfotiamine has been found to block sugar-induced damage in a number of ways:

- It activates a vital enzyme called *transketolase*. which converts toxic glucose-induced metabolites into harmless compounds.^{3,16}
- It helps prevent inflammation by inhibiting the activation of nuclear factor-kappaB (NF-kB), a trigger for dangerous inflammatory reactions in the body.16
- A 2012 study showed that benfotiamine boosts glucose breakdown in hyperglycemic (highglucose) conditions by 70% and up to 49% in normoglycemic (normal-glucose) conditions!43

Benfotiamine is superior to thiamine alone in mitigating the spectrum of diseases such as neurodegenerative diseases and cancer,44 as well as the kidney, eye,

and nerve damage seen in diabetic patients. 7 Let's now examine the evidence demonstrating benfotiamine's potent protection against several of the most common sugar-induced disorders.

Benfotiamine Reverses The Pathways **And Memory Loss of Alzheimer's**

Recent studies have indicated that benfotiamine could potentially enhance memory and contribute to the delay in the progression of Alzheimer's disease.

When benfotiamine treatment was used for eight weeks in a mouse model of Alzheimer's disease, researchers found that it reduced both amyloid plaque



numbers and **phosphorylated tau** protein levels in the brain. It also helped prevent amyloid plaque from forming in the first place by reducing enzyme activities related to amyloid precursor proteins. 45 These changes were validated by a dose-dependent enhancement in memory.45

The study authors concluded that "benfotiamine may be beneficial for clinical Alzheimer's disease treatment."45

Then in 2012, scientists determined that benfotiamine was able to do more than just **reduce** the plaque and tau formation—it could regress it. One of the ways it produced this effect was through a reduction in advanced glycation end products. The study authors wrote that "the use of benfotiamine could provide a safe intervention to reverse biological and clinical processes of Alzheimer's disease progression."46

Defense Against Sugar-Induced Vision Loss

The same blood sugar effects that are associated with blindness in diabetics can cause cataract and retinal damage in nondiabetics as well. 47,48 Because of its high vascularity, eye tissue is especially susceptible to damage from advanced glycation end products caused by elevated glucose levels.

In an animal study, benfotiamine supplementation was found to **reverse** retinal accumulation of AGEs. as well as increasing their urinary excretion.⁴⁹ This can lead to a number of beneficial effects on the eyes.

Elevated blood sugar and the formation of AGEs can trigger cell death within the retina. However, when researchers applied benfotiamine to retinal blood vessel cells in culture, they found that the benfotiamine was

Block The Aging Effects Of High Blood Sugar

- Glycation-the deadly binding of sugars to proteins-produces deadly byproducts called advanced glycation end products (AGEs).
- AGEs damage cells, tissues, and organs, and cause premature aging.
- Thiamine is the body's natural defense against AGEs, but since it is water soluble, the body doesn't absorb it well.
- Benfotiamine, a fat-soluble derivative of thiamine (vitamin B1), substantially boosts intracellular thiamine levels, inhibiting the AGEs that cause sugar-related damage to tissue.
- Numerous studies demonstrate that benfotiamine blocks the pathways of Alzheimer's disease, vision loss, cardiovascular disease, and kidney damage.
- Benfotiamine is extremely safe and appears to slow the chronic disease and aging effects of glycation in nondiabetic as well as diabetic individuals.

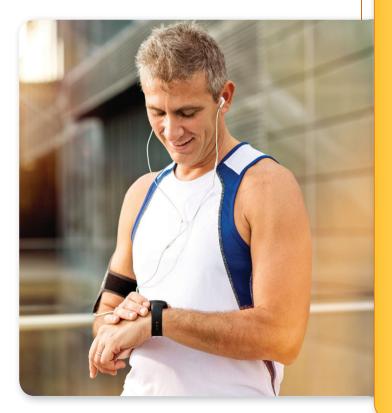
able to prevent this dangerous process. It was also able to reduce DNA damage.50

In the eye, advanced glycation end products increase enzymes called *matrix metalloproteinases* (MMPs). These protein-dissolving enzymes cause substantial damage to retinal tissue. Scientists found that benfotiamine lowered MMP production to normal levels and increased production of certain proteins that specifically inhibit their activity.51

Inflammatory changes associated with aging, elevated blood glucose, and infections can affect the eves. Uveitis, a serious inflammation of the middle layer of the eye, is responsible for 10% of all cases of blindness in the United States⁵² and is characterized by infiltrates of white blood cells, proteins, and inflammatory cytokines that inflict oxidative stress, leading to further inflammation.⁵³ Benfotiamine was found to suppress this process and the expression of inflammatory marker molecules.53

Benfotiamine Protects Against Cardiovascular Damage

The thin layer of endothelial cells that lines blood vessels constantly regulates blood pressure and flow. Advanced glycation end products damage these cells, which can lead to heart attacks, heart failure, and stroke.54,55



Advanced glycation end products damage blood vessels in three ways:

- 1. They reduce endothelial cell replication. a vital process to keep arteries healthy.⁵⁶
- 2. They trigger an increased level of death among endothelial cells.⁵⁷
- 3. They produce toxic **polyol compounds**, which disrupt the functioning of vascular and endothelial cells.58

Remarkably, studies indicate that benfotiamine protects endothelial cells from all three of these damaging effects of AGEs. 56-58 This protection ultimately improves the functioning of blood vessels and heart muscle. 59,60

An interesting study on humans revealed just what a dramatic impact these effects would have on heart health. In the study, subjects took 1,050 mg of benfotiamine per day for three days. The scientists wanted

Conditions Linked With Elevated Blood Glucose And Advanced Glycation End Products

- Neuronal dysfunction⁶⁹
- Alzheimer's disease^{69,70}
- Cardiovascular disease^{24,27,31} and stroke⁷¹
- Eye cataracts²⁹
- Reduced muscle function¹⁹
- Kidney disease^{2,30}
- Neuropathy²³
- Liver damage⁷²
- Stiffening of connective tissues in joints 17,73
- Damage to collagen and elastin, which ages the skin¹⁹
- DNA damage⁷⁴ and an increased risk of cancer⁶⁸
- Enlarged prostate, or benign prostatic hyperplasia (BPH)75
- Greater incidence and severity of obstructive sleep apnea²⁸
- Accelerated aging^{2,32}



to determine the effect benfotiamine would have on the changes in endothelial function and blood flow that occur as a result of consuming a meal rich in advanced glycation end products.⁵⁹ The striking outcome of supplementation revealed that benfotiamine *completely* prevented endothelial and blood flow impairment!^{26,61}

Benfotiamine also counters the negative effects of oxidative stress on blood vessels. In a study in which rodents were experimentally induced with vascular endothelial dysfunction, supplemental benfotiamine resulted in improved endothelial integrity and function by enhancement of production of *nitric oxide*, a compound that helps blood vessels to relax.^{53,73}

Blood sugar and advanced glycation end products greatly contribute to the diminished heart-muscle function known as heart failure, which can be caused by a heart attack or untreated high blood pressure. Scientists have demonstrated that benfotiamine helps normalize the heart muscle's ability to contract, salvaging damaged heart muscle and improving its capacity to effectively pump blood.⁶⁰

Benfotiamine Inhibits Sugar-Related Kidney Disease

Kidney disease, or *nephropathy*, is a known side effect of "natural" aging, but it is also a common and very serious complication of excess blood sugar and especially of diabetes.

Like the eyes, the kidneys are rich in tiny blood vessels known as *capillaries*. The kidneys are also the site of intense *metabolic activity*. These facts make the kidneys particularly vulnerable to the damaging effects of glucose and advanced glycation end products.³⁰

The good news is that evidence indicates that benfotiamine helps protect against sugar-related kidney damage by reducing pathological increases in advanced glycation end products.^{49,62} In one study, benfotiamine activated the important enzyme *transketolase*, which rapidly cleared AGEs from the blood before they could damage kidney tissue.⁶²

Another study demonstrated that benfotiamine could help reduce sugar-induced kidney damage as efficiently as the prescription drug *fenofibrate*.⁶³ The two in combination demonstrated beneficial synergistic effects.⁶³

Benfotiamine has also been found to be helpful during various forms of *dialysis* treatments.

Hemodialysis, which is a last-resort treatment for patients whose kidneys have failed, quickly depletes the body's store of thiamine, the *very* vitamin required to prevent further glucose-related damage. Benfotiamine, which is vastly more bioavailable than thiamine, boosts thiamine levels over **4 times higher** than supplementation with thiamine in dialysis patients.⁶⁴

Peritoneal dialysis is somewhat less stressful to the body than hemodialysis, but it is associated with substantial damage to the delicate tissues lining the abdominal cavity, believed to be caused by glucose and AGEs. ⁶⁵ By reducing the level of AGEs during peritoneal dialysis, benfotiamine decreases markers of inflammation and abnormal new blood vessel formation in the abdominal cavity. ⁶⁵ This protects the delicate abdominal lining, prolonging its usefulness as a dialysis site.

Both types of dialysis also substantially damage DNA throughout the body, increasing the risk of cancer. Benfotiamine treatment has been shown to significantly reduce this DNA damage in dialysis patients by reducing the circulating advanced glycation end products.⁶⁶⁻⁶⁸

These results provide dramatic evidence of the high degree of kidney protection that benfotiamine offers against blood sugar increases and advanced glycation end products.

Summary

Just over a decade ago, the health-devastating and faster-aging effects of high blood sugar were associated almost exclusively with diabetics. But healthy individuals are also susceptible to the chronic, agerelated diseases triggered by high blood sugar.

Something as simple as eating a heavy meal can ultimately lead to the production of deadly byproducts called advanced glycation end products that damage cells, tissues, and organs—and even cause premature

Fortunately, the discovery of a relative of thiamine (vitamin B1) called **benfotiamine** has provided a way to powerfully boost intracellular levels of thiamine, the body's natural defense against the AGEs that cause sugar-related tissue damage.

Numerous studies confirm that benfotiamine blocks the pathways of Alzheimer's disease, vision loss, cardiovascular disease, and kidney damage.

Benfotiamine, with its impressive safety record, has shown itself to be a powerful factor in slowing glycation-induced aging—potentially maximizing life span—in both diabetic and nondiabetic individuals.

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

References

- Ko GT, Wai HP, Tang JS. Effects of age on plasma glucose levels in non-diabetic Hong Kong Chinese. Croat Med J. 2006 Oct;47(5):709-
- Monnier VM, Sell DR, Nagaraj RH, et al. Maillard reaction-mediated molecular damage to extracellular matrix and other tissue proteins in diabetes, aging, and uremia. Diabetes. 1992 Oct;41 Suppl 2:36-41.
- Obrenovich ME, Monnier VM. Vitamin B1 blocks damage caused by hyperglycemia. Sci Aging Knowledge Environ. 2003 Mar 12;2003(10):PE6
- Cameron NE, Gibson TM, Nangle MR, et al. Inhibitors of advanced glycation end product formation and neurovascular dysfunction in experimental diabetes. Ann NY Acad Sci. 2005 Jun:1043:784-92
- Giusti C, Gargiulo P. Advances in biochemical mechanisms of diabetic retinopathy. Eur Rev Med Pharmacol Sci. 2007 May;11(3):155-63.
- Head KA. Peripheral neuropathy: pathogenic mechanisms and alternative therapies. Altern Med Rev. 2006 Dec;11(4):294-329.
- Karachalias N, Babaei-Jadidi R, Ahmed N, Thornalley PJ. Accumulation of fructosyl-lysine and advanced glycation end products in the kidney, retina and peripheral nerve of streptozotocin-induced diabetic rats. Biochem Soc Trans. 2003 Dec;31(Pt 6):1423-5.
- Ahmed N, Thornalley PJ. Advanced glycation endproducts: what is their relevance to diabetic complications? Diabetes Obes Metab. 2007 May;9(3):233-45.
- Vasdev S, Gill V, Singal P. Role of advanced glycation end products in hypertension and atherosclerosis: therapeutic implications. Cell Biochem Biophys. 2007;49(1):48-63.
- 10. Bjornholt JV, Erikssen G, Aaser E, et al. Fasting blood glucose: an underestimated risk factor for cardiovascular death. Results from a 22-year follow-up of healthy nondiabetic men. Diabetes Care. 1999 Jan; 22(1): 45-9.
- 11. McGlothin, P, Averill M. The CR Way: Using the Secrets of Calorie Restriction for a Longer, Healthier Life. NY: HarperCollins; 2008:





- 12. Thorpe SR, Baynes JW. Role of the Maillard reaction in diabetes mellitus and diseases of aging. Drugs Aging. 1996 Aug;9(2):69-77. 2.
- Matthews CE, Sui X, LaMonte MJ, Adams SA, Hebert JR, Blair SN. Metabolic syndrome and risk of death from cancers of the digestive system. Metabolism. 2010 Aug;59(8):1231-9.
- 14. Sick E, Brehin S, Andre P, et al. Advanced glycation end products (AGEs) activate mast cells. Br J Pharmacol. 2010 Sep;161(2):
- 15. Balakumar P, Rohilla A, Krishan P, Solairaj P, Thangathirupathi A. The multifaceted therapeutic potential of benfotiamine. Pharmacol Res. 2010 Jun:61(6):482-8.
- 16. Hammes HP, Du X, Edelstein D, et al. Benfotiamine blocks three major pathways of hyperglycemic damage and prevents experimental diabetic retinopathy. Nat Med. 2003 Mar;9(3):294-9.
- 17. van Boekel MA. The role of glycation in aging and diabetes mellitus. Mol Biol Rep. 1991 May;15(2):57-64.
- Sztanke K, Pasternak K. The Maillard reaction and its consequences for a living body. Ann Univ Mariae Curie Sklodowska Med. 2003;58(2):159-62.
- 19. Haus J, Carrithers J, Trappe S, Trappe T. (2007). Collagen, crosslinking, and advanced glycation end products in aging human skeletal muscle. J Appl Physiol. 2007;103(6):2068-76.
- Shaikh S, Nicholson LF. Advanced glycation end products induce in vitro cross-linking of alpha-synuclein and accelerate the process of intracellular inclusion body formation. J Neurosci Res. 2008 Jul;86(9):2071-82.
- 21. Uribarri J, Cai W, Sandu O, Peppa M, Goldberg T, Vlassara H. Diet-derived advanced glycation end products are major contributors to the body's AGE pool and induce inflammation in healthy subjects. Ann NY Acad Sci. 2005 Jun;1043:461-6.
- 22. Booth AA, Khalifah RG, Todd P, Hudson BG. In vitro kinetic studies of formation of antigenic advanced glycation end products (AGEs). Novel inhibition of post-Amadori glycation pathways. J Biol Chem. 1997 Feb 28;272(9):5430-7.
- 23. Sugimoto K, Yasujima M, Yagihashi S. Role of advanced glycation end products in diabetic neuropathy. Curr Pharm Des. 2008;14(10):953-61.
- Goldin A, Beckman JA, Schmidt AM, Creager MA. Advanced glycation end products: sparking the development of diabetic vascular injury. Circulation. 2006 Aug 8;114(6):597-605.
- 25. Thornalley PJ. Glycation in diabetic neuropathy: characteristics, consequences, causes, and therapeutic options. Int Rev Neurobiol. 2002;50:37-57.

- 26. Verma S, Reddy K, Balakumar P. The defensive effect of benfotiamine in sodium arsenite-induced experimental vascular endothelial dysfunction. Biol Trace Elem Res. 2010 Oct;137(1): 96-109.
- 27. Urberg M, Rajdev K. A correlation between serum cholesterol and glycosylated hemoglobin in nondiabetic humans. J Fam Pract. 1989 Mar;28(3):269-74.
- 28. Tan KC, Chow WS, Lam JC, et al. Advanced glycation endproducts in nondiabetic patients with obstructive sleep apnea. Sleep. 2006:29(3):329-33.
- 29. Gul A, Rahman MA, Hasnain SN. Role of fructose concentration on cataractogenesis in senile diabetic and non-diabetic patients. Graefes Arch Clin Exp Ophthalmol. 2009 Jun;247(6):809-14.
- 30. Hall PM. Prevention of progression in diabetic nephropathy. Diabetes Spect. 2006 Jan;19(1):18-24.
- 31. Simm A, Wagner J, Gursinsky T, Nass N, Friedrich I, Schinzel R, Czeslik E, Silber RE, Scheubel RJ. Advanced glycation endproducts: a biomarker for age as an outcome predictor after cardiac surgery? Exp Gerontol. 2007 Jul;42(7):668-75.
- 32. Vlassara H, Palace MR. Glycoxidation: the menace of diabetes and aging. Mt Sinai J Med. 2003 Sep;70(4):232-41.
- 33. Yamagishi S, Adachi H, Takeuchi M, et al. Serum level of advanced glycation end-products (AGEs) is an independent determinant of plasminogen activator inhibitor-1 (PAI-1) in nondiabetic general population. Horm Metab Res. 2007 Nov;39(11):845-8.
- 34. Hartog JW, Voors AA, Schalkwijk CG, et al. Clinical and prognostic value of advanced glycation end-products in chronic heart failure. Eur Heart J. 2007 Dec;28(23):2879-85.
- 35. Robert L, Robert AM, Fülöp T. Rapid increase in human life expectancy: will it soon be limited by the aging of elastin? Biogerontology. 2008 Apr;9(2):119-33.
- 36. Booth AA, Khalifah RG, Hudson BG. Thiamine pyrophosphate and pyridoxamine inhibit the formation of antigenic advanced glycation end-products: comparison with aminoguanidine. Biochem Biophys Res Commun. 1996 Mar 7;220(1):113-9.
- 37. Beltramo E, Berrone E, Tarallo S, Porta M. Effects of thiamine and benfotiamine on intracellular glucose metabolism and relevance in the prevention of diabetic complications. Acta Diabetol. 2008 Sep;45(3):131-41.
- 38. Geyer J, Netzel M, Bitsch I, et al. Bioavailability of water- and lipid-soluble thiamin compounds in broiler chickens. Int J Vitam Nutr Res. 2000 Dec;70(6):311-6.

- 39. Hilbig R, Rahmann H. Comparative autoradiographic investigations on the tissue distribution of benfotiamine versus thiamine in mice. Arzneimittelforschung. 1998 May;48(5):461-8.
- Stracke H, Hammes HP, Werkmann D, et al. Efficacy of benfotiamine versus thiamine on function and glycation products of peripheral nerves in diabetic rats. Exp Clin Endocrinol Diabetes. 2001;109(6):330-6.
- 41. Volvert ML, Seyen S, Piette M, et al. Benfotiamine, a synthetic S-acyl thiamine derivative has different mechanisms of action and a different pharmacological profile than lipid-soluble thiamine disulfide derivatives. BMC Pharmacol. 2008 Jun 12;8:10.
- 42. Available at: http://benfotiamine.org/Benfo600Study.pdf. Accessed September 15, 2015.
- 43. Fraser DA, Hessvik NP, Nikoli N, et al. Benfotiamine increases glucose oxidation and downregulates NADPH oxidase 4 expression in cultured human myotubes exposed to both normal and high glucose concentrations. Genes Nutr. 2012 July;7(3):459-69.
- 44. Coy JF, Dressler D, Wilde J, Schubert P. Mutations in the transketolase-like gene TKTL1: clinical implications for neurodegenerative diseases, diabetes and cancer. Clin Lab. 2005;51(5-6):257-73.
- 45. Pan X, Gong N, Zhao J, et al. Powerful beneficial effects of benfotiamine on cognitive impairment and beta-amyloid deposition in amyloid precursor protein/presenilin-1 transgenic mice. Brain. 2010 May;133(Pt 5):1342-51.
- 46. Gibson GE, Hirsch JA, Cirio RT, et al. Abnormal thiamine-dependent processes in Alzheimer's disease. Lessons from diabetes. Mol Cell Neurosci, 2013 Jul:55:17-25.
- 47. Pokupec R, Kalauz M, Turk N, Turk Z. Advanced glycation endproducts in human diabetic and non-diabetic cataractous lenses. Graefes Arch Clin Exp Ophthalmol. 2003 May;241(5):378-84.
- 48. Kandarakis SA, Piperi C, Topouzis F, Papavassiliou AG. Emerging role of advanced glycation-end products (AGEs) in the pathobiology of eye diseases. Prog Retin Eye Res. 2014 Sep;42:85-102.
- 49. Karachalias N, Babaei-Jadidi R, Rabbani N, Thornalley PJ. Increased protein damage in renal glomeruli, retina, nerve, plasma and urine and its prevention by thiamine and benfotiamine therapy in a rat model of diabetes. Diabetologia. 2010 Jul;53(7):1506-16.
- 50. Beltramo E, Nizheradze K, Berrone E, Tarallo S, Porta M. Thiamine and benfotiamine prevent apoptosis induced by high glucoseconditioned extracellular matrix in human retinal pericytes. Diabetes Metab Res Rev. 2009 Oct;25(7):647-56.
- 51. Tarallo S, Beltramo E, Berrone E, Dentelli P, Porta M. Effects of high glucose and thiamine on the balance between matrix metalloproteinases and their tissue inhibitors in vascular cells. Acta Diabetol. 2010 Jun;47(2):105-11.
- 52. Ke Y, Jiang G, Sun D, Kaplan HJ, Shao H. Anti-CD3 antibody ameliorates experimental autoimmune uveitis by inducing both IL-10 and TGF-beta dependent regulatory T cells. Clin Immunol. 2011 Mar;138(3):311-20.
- 53. Yadav UC, Subramanyam S, Ramana KV. Prevention of endotoxininduced uveitis in rats by benfotiamine, a lipophilic analogue of vitamin B1. Invest Ophthalmol Vis Sci. 2009 May;50(5):2276-82.
- 54. Thomas MC, Baynes JW, Thorpe SR, Cooper ME. The role of AGEs and AGE inhibitors in diabetic cardiovascular disease. Curr Drug Targets. 2005 Jun;6(4):453-74.
- 55. Hanssen NM, Beulens JW, van Dieren S, et al. Plasma advanced glycation end products are associated with incident cardiovascular events in individuals with type 2 diabetes: a case-cohort study with a median follow-up of 10 years (EPIC-NL). Diabetes. 2015 Jan;64(1):257-65.
- 56. Pomero F, Molinar Min A, La Selva M, Allione A, Molinatti GM, Porta M. Benfotiamine is similar to thiamine in correcting endothelial cell defects induced by high glucose. Acta Diabetol. 2001:38(3):135-8.
- 57. Beltramo E, Berrone E, Buttiglieri S, Porta M. Thiamine and benfotiamine prevent increased apoptosis in endothelial cells and pericytes cultured in high glucose. Diabetes Metab Res Rev. 2004 Jul-Aug;20(4):330-6.
- 58. Berrone E, Beltramo E, Solimine C, Ape AU, Porta M. Regulation of intracellular glucose and polyol pathway by thiamine and benfotiamine in vascular cells cultured in high glucose. J Biol Chem. 2006 Apr 7;281(14):9307-13.

- 59. Stirban A, Negrean M, Stratmann B, et al. Benfotiamine prevents macro- and microvascular endothelial dysfunction and oxidative stress following a meal rich in advanced glycation end products in individuals with type 2 diabetes. Diabetes Care. 2006 Sep:29(9):2064-71.
- 60. Cevlan-Isik AF, Wu S, Li Q, Li SY, Ren J. High-dose benfotiamine rescues cardiomyocyte contractile dysfunction in streptozotocininduced diabetes mellitus. J Appl Physiol. 2006 Jan;100(1):150-6.
- 61. Balakumar P, Sharma R, Singh M. Benfotiamine attenuates nicotine and uric acid-induced vascular endothelial dysfunction in the rat. Pharmacol Res. 2008 Nov-Dec;58(5-6):356-63.
- 62. Babaei-Jadidi R, Karachalias N, Ahmed N, Battah S, Thornalley PJ. Prevention of incipient diabetic nephropathy by high-dose thiamine and benfotiamine. Diabetes. 2003 Aug;52(8):2110-20.
- 63. Balakumar P, Chakkarwar VA, Singh M. Ameliorative effect of combination of benfotiamine and fenofibrate in diabetes-induced vascular endothelial dysfunction and nephropathy in the rat. Mol Cell Biochem. 2009 Jan; 320(1-2):149-62.
- 64. Frank T, Bitsch R, Maiwald J, Stein G. High thiamine diphosphate concentrations in erythrocytes can be achieved in dialysis patients by oral administration of benfotiamine. Eur J Clin Pharmacol. 2000 Jun;56(3):251-7.
- 65. Kihm LP, Muller-Krebs S, Klein J, et al. Benfotiamine protects against peritoneal and kidney damage in peritoneal dialysis. J Am Soc Nephrol. 2011 May;22(5):914-26.
- 66. Schmid U, Stopper H, Heidland A, Schupp N. Benfotiamine exhibits direct antioxidative capacity and prevents induction of DNA damage in vitro. Diabetes Metab Res Rev. 2008 Jul-Aug;24(5):371-7.
- 67. Schupp N, Schmid U, Heidland A, Stopper H. New approaches for the treatment of genomic damage in end-stage renal disease. J Ren Nutr. 2008 Jan;18(1):127-33.
- 68. Schupp N, Dette EM, Schmid U, et al. Benfotiamine reduces genomic damage in peripheral lymphocytes of hemodialysis patients. Naunyn Schmiedebergs Arch Pharmacol. 2008 Sep;378(3):283-91.
- 69. Sasaki N, Fukatsu R, Tsuzuki K, et al. Advanced glycation end products in Alzheimer's disease and other neurodegenerative diseases. Am J Pathol. 1998 Oct;153(4):1149-55.
- 70. Srikanth V. MacZurek A. Phan T. et al. Advanced glycation endproducts and their receptor RAGE in Alzheimer's disease. Neurobiology of Aging. 2011;32(5):763-77.
- 71. Zimmerman GA, Meistrell M III, Bloom O, et al. Neurotoxicity of advanced glycation endproducts during focal stroke and neuroprotective effects of aminoguanidine. Proc Natl Acad Sci (USA). 1995 Apr 25;92(9):3744-8.
- 72. Succurro E, Arturi F, Grembiale A, et al. One-hour post-load plasma glucose levels are associated with elevated liver enzymes. Nutr Metab Cardiovasc Dis. 2011 Sep;21(9):713-8.
- 73. Rosenbloom AL, Silverstein JH. Connective tissue and joint disease in diabetes mellitus. Endocrinol Metab Clin North Am. 1996 Jun;25(2):473-83.
- 74. Suji G, Sivakami S. DNA damage during glycation of lysine by methylglyoxal: assessment of vitamins in preventing damage. Amino Acids. 2007 Nov;33(4):615-21.
- 75. Kim WT, Yun SJ, Choi YD, et al. Prostate size correlates with fasting blood glucose in non-diabetic benign prostatic hyperplasia patients with normal testosterone levels. J Korean Med Sci. 2011 Sep;26(9):1214-8.

POWERFUL PROTECTION FOR A LOW PRICE

GREEN TEA EXTRACT

EGCG FOR OPTIMAL HEALTH

Nearly 6,000 studies have been published on the broad-spectrum health benefits of **green tea**.

Research shows that green tea favorably influences cardiovascular health, lipid clearance, glucose tolerance, healthy body weight, DNA support, prostate and breast health, and healthy cell division. Focientists have identified the polyphenol **EGCG** as the key compound for green tea's multimodal health benefits.

Life Extension® has created a standardized 98% polyphenol green tea extract. These highly concentrated Mega Green Tea Extract Capsules contain 725 mg of either lightly caffeinated or decaffeinated 98% standardized green tea extracts.

Each bottle will last over **three months** at the typical dose of <u>one</u> capsule daily.

Non-GMO

References

- 1. Harv Heart Lett. 2012 Dec;23(4):7.
- PLoS One. 2014 May 1;9(5):e96884
 J Nutr Biochem. 2014 May;25(5):557-64.
- 4. Endocrinology. 2000 Mar;141(3):980-7.
- Endocrinology. 2000 Mar;141(3):980
 Nutr. Res. 2002 Oct;22(10):1143-50
- 6. Int J Cancer. 2004 Jan 1;108(1):130-5.
- 7. J Agric Food Chem. 2007 May 2;55(9):3378-85.
- 8. Genes Nutr. Mar 2010; 5(1): 75–87.

Mega Green Tea Extract Decaffeinated

Item #00954 • 100 vegetarian capsules

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



Mega Green Tea Extract Lightly Caffeinated

Item #00953 • 100 vegetarian capsules

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



To order Mega Green Tea Extract, call 1-800-544-4440 or visit www.LifeExtension.com

Advanced Defense Against Cellular Aging

NAD+ Cell Regenerator

If you are experiencing fatigue and lack of motivation, it may be due to the age-related decline in NAD+ levels¹ and subsequent impairment of healthy cellular metabolism.²

NAD+ (*nicotinamide adenine dinucleotide*) promotes systemic youthful functions and is found in every cell in the body.³ In addition, **NAD+** plays an essential role in regulating **genes**^{4,5} that control aging.⁶

How To Boost NAD+ Levels Within Your Cells

Newly patented **nicotinamide riboside** increases cellular levels of NAD+ in the body.7,8

For the first time, aging humans have an effective and affordable method to boost the critical **NAD+** enzyme for refreshed vitality.

Nicotinamide riboside represents an innovative advance to combat aging that functions via unique mechanisms not found in typical dietary supplements. It has emerged as a front-line weapon in Life Extension®'s ongoing war against premature aging.

The name of this new nicotinamide riboside formulation is NAD+ Cell Regenerator™.

Multiple Benefits Of Increasing NAD+ Cellular Levels

Nicotinamide riboside has been documented to help replenish cellular **NAD+** and in the process:

- Promote sirtuin (SIRT1 and SIRT3) gene activation,6
- Enhance growth and efficiency of mitochondria—supporting energy levels and physical performance,6
- Favorably modulate metabolism,⁶
- Contribute to neuronal health—supporting cognitive function during aging,9-11
- Promote insulin activity—supporting healthy blood sugar in those within the normal range.6

References

- PLoS One. 2012;7(7):e42357.
- Cell. 2013 Dec 19;155(7):1624-38.
- PLoS One. 2008 May 21;3(5):e2267. Cell. 2007 May 4;129(3):473-84.
- Nature. 2000 Feb 17;403(6771):795-800.
- Cell Metab. 2012 Jun 6;15(6):838-47. Curr Opin Clin Nutr Metab Care. 2013
- Nov;16(6):657-61
- EMBO Mol Med. 2014 Apr 6;6(6):721-31. Neurobiol Aging. 2013 Jun;34(6):1581-8. 10. J Neurosci. 2006 Aug 16;26(33):8484-91.
- 11. Front Biosci. 2007 Jan 1;12:1863-88.

Life Extension® **NAD+ Cell Regenerator**™ contains the patented ingredient NIAGEN®, the first commercially available form of nicotinamide riboside.

The suggested daily dose of one NAD+ Cell **Regenerator**[™] vegetarian capsule provides:

NIAGEN® Nicotinamide Riboside 100 mg

NAD+ Cell Regenerator™

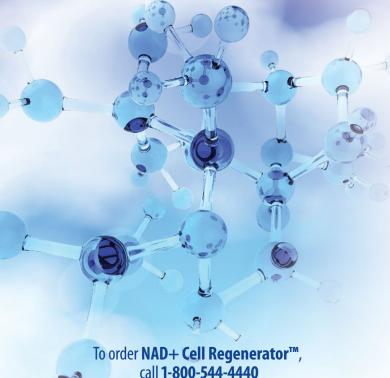
Item #01904 • 30 vegetarian capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$34 | \$25.50 |
| 4 bottles | | \$19.50 each |

Non-GMO

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





or visit www.LifeExtension.com

MEGA BENFOTIAMINE

PROTECT AGAINST SYSTEMIC GLYCATION

Scientists have known that when blood sugar combines with fats and proteins the result is known as *glycation*, and it produces *accelerated aging*.¹ Even those with blood sugar levels within normal range experience the impact of systemic *glycation* on a daily basis.²

benfotiamine, a unique form of vitamin B1 (thiamine) that supports healthy blood sugar metabolism and protects against *glycation*.³⁻⁵ What makes benfotiamine especially effective is that unlike ordinary vitamin B1, it is fat soluble and can easily penetrate the inside of cells.⁶ Regular vitamin B1 is water soluble and has a short life span in the body.⁷

Mega Benfotiamine helps inhibit the formation of *advanced glycation end products* (AGEs) to maintain healthy endothelial, retinal, kidney, and nerve cell function.⁸⁻¹²

Each capsule provides **250 mg** of **benfotiamine** and **10 mg** of vitamin B1 (as thiamine HCl).

References

- 1. Drugs & Aging. 1996 Aug;9(2):69-77.
- 2. J Biol Chem. 2011 Dec 30; 286(52): 44350-6.
- 3. J Biochem. 1954;41:219-44.
- 4. Genes Nutr. 2012 Jul;7(3):459-69.
- 5. Diabetes Metab Res Rev. 2004 Jul-Aug;20(4):330-6.
- 6. Acta Diabetol. 2001;38(3):135-8.
- 7. Nutr Res. 2010 Mar;30(3):171-8.
- 8. Nat Med. 2003 Mar;9(3):294-9.
- 9. Diabetes Care. 2006 Sep;29(9):2064-71.
- 10. Ann N Y Acad Sci. 2005 Jun;1043:777-83.
- 11. Pharmacol Res. 2010 Jun;61(6):482-8.
- 12. *Ann N Y Acad Sci.* 2005 Jun;1043:784-92.



To order **Mega Benfotiamine**, call **1-800-544-4440** or visit **www.LifeExtension.com** The Mitochondrial Energy Optimizer formula provides **150 mg** of benfotiamine, which when combined with other antiglycation agents like carnosine and pyridoxal-5-phosphate, provides substantial protection at different checkpoint opportunities in the glycation process. Item #01868



CANCER RISKS Of CT SCANS

Every year, Americans are exposed to potentially unsafe levels of DNA-altering **radiation** through medical imaging such as CT scans.

An astonishing **72 million** CT scans are performed annually in the United States, which is about one scan for every four people in the country each year.¹

Why is this so troubling? Because the radiation in that CT scan can <u>increase</u> your risk of cancer.

National Cancer Institute researchers now estimate that those 72 million CT scans could account for roughly 29,000 future cancer cases each year! Another way of looking at this figure is that for roughly every 400 to 2,000 routine chest CT scans, one new case of cancer occurs.

And, by some estimates, up to **44%** of CT scans done in this country each year are medically <u>un</u>necessary.²

Despite these dangers, modern imaging techniques have been a tremendous boon to health care, allowing physicians to spot cancers, fractures, aneurysms, abscesses, and other risks that would otherwise go undetected.

But what if there was a way to get the benefits of a CT scan while reducing your risk of DNA mutation?

Three natural compounds have been shown to have properties that may help counteract the most dangerous consequences of radiation exposure.

If your doctor orders an X-ray or CT scan, taking these compounds at least <u>five</u> days prior may help protect against cellular damage inflicted by the **ionizing radiation**.

In addition, regular consumption of **blue-berries** or **blueberry extract** enhances **DNA repair**, which may help reverse some of the genetic damage inflicted by ionizing radiation.

CT Scans Boost Cancer Risk

After years of overlooking the potential risks of medical imaging, mainstream physicians are finally beginning to acknowledge the dangers of these radiation-based diagnostic tools.

In 2013, a scientific consensus was reached that even just one **CT scan** in childhood is linked to the risk of developing future cancers. Of course, cancer risk estimates for individuals must also include other criteria like specifics of exposure, age at exposure, and absorbed dose to certain tissues.3

This consensus was likely triggered by a deeply disturbing 2013 study published in the prestigious British Medical Journal (BMJ).4

The researchers behind that study followed approximately 11 million individuals from birth in the 1980s into young adulthood, identifying over 680,000 who had at least one CT scan during that period. They compared the cancer rate in this extremely large group of patients with an equally large, matched group who were never exposed to a CT scan.

The results were alarming.

The study found that those who had undergone CT scans in their childhood had a 24% increased risk for developing any cancer, compared with those who didn't have scans. 4 They also found that the more scans a person had, the greater the risk of developing cancer.



The risk persisted for years after the original scan was completed, producing cancer risks, compared with unexposed individuals, which were:4

- **35%** higher in the first four years following exposure
- **25**% higher at five to nine years
- **14**% higher at 10 to 14 years

Even 15 or more years following the first exposure to this level of ionizing radiation, cancer risks remained stubbornly elevated by 24%.

Virtually every kind of cancer was documented to occur in excess in the CT-exposed group, including solid tumors (digestive organs, skin, ovary/uterus, urinary tract, brain, and thyroid), leukemia (blood cancer), lymphomas, and other cancers.4

Let's take a quick look now at exactly *how* the radiation in CT scans (and other X-ray-based technologies) raises your cancer risk and threatens your longevity. Understanding this will help you and your doctor make a more rational decision as to whether a particular CT scan is scientifically warranted.

How Ionizing Radiation Causes Cancer And Other Deadly Threats

For all the potential good they can do, CT scans expose us to surprisingly high levels of ionizing radiation. On a cellular level, ionizing radiation can strip electrons from the atoms that make up our tissues. producing energetic chemical ions that damage tissue and impose potent genetic stresses.

One of the most dangerous outcomes of exposure to ionizing radiation is damage to the DNA contained in the nucleus of the cells in your body. 5 Damaged DNA is an open invitation to cancer development. The problem that damaged DNA presents is that it removes the effective controls our bodies use to regulate the cell reproductive cycle and keep cell growth and replication in check.6

Ionizing radiation's effects on tissues are both immediate and long lasting. When human tissue is irradiated, a number of DNA-breaking compounds form. Together these are known as "clastogenic factors." 7-9 Most clastogenic factors that form in living tissue are associated with reactive oxygen species whose unpaired electrons actively react with DNA molecules, inducing breaks in DNA strands that can lead to cancer. 10-12

Unfortunately, cancer-inducing clastogenic factors have a very long life within the body. Studies show that survivors of the atomic bomb attacks in Japan continued to have such factors detectable in their blood more than 30 years later.7

Ionizing radiation poses a large range of other threats to one's health as well. It can directly impair cell functions, leading to the loss of proper tissue and organ operation as well as killing cells.¹³ One victim of ionizing radiation is the fat-laden cell membrane. Under the effects of ionizing radiation, membrane-bound fats undergo oxidation and form toxic breakdown products.14

Recent studies also document that X-rays and other forms of ionizing radiation activate inflammatory pathways and produce early cell death.¹⁵ And, ever since the first nuclear physicists began assembling atomic bomb components, it has been clear that ionizing radiation causes potentially lethal but silent bone marrow damage. This impairs the development of the white blood cells that protect us from infections and cancer, the red blood cells that carry oxygen to all of our tissues, and the platelets that help us stop bleeding after an injury. 16,17 And bone marrow is where hematological cancers like leukemia originate.

What You Need To Know CT Scans And Cancer Risk Exposure to ionizing radiation is growing rapidly in the US, partly the result of an explosion in the use of X-ray-based imaging studies, particularly CT scans. Ionizing radiation produces reactive compounds that directly damage cells that cause DNA strand breaks leading to cancer formation. Radiation exposure also suppresses bone marrow production of infection- and cancer-fighting white blood cells.

Protection From Ionizing Radiation

Fortunately, it may be possible to derive the benefits of modern imaging techniques while obtaining some protection from the radiation exposure.

Three specific nutrients have been found to help counteract the toxic effects of ionizing radiation. They include the following:

- **1.** *Lemon balm*, which prevents the formation of damaging chemicals triggered by ionizing radiation, 18,19
- **2.** *Ginkgo biloba*, which protects fragile DNA from the resulting cancer-causing damage,7,8
- 3. Spirulina, which stimulates the immune system, particularly bone marrow, to make and maintain levels of white blood cells, whose production is impaired by ionizing radiation.²⁰

Let's look at each individually.

You may help protect yourself ahead of a scheduled CT scan or other study involving ionizing radiation by taking specific nutrients on a timely basis.

test is completed.

Studies show that exposure to CT scans can increase the risk of cancer by at least 24%,

and that those risks last for years after the

- Lemon balm reduces production of reactive chemical compounds by radiation, helping cells retain their integrity.
- Ginkgo biloba reduces DNA damage that can lead to cancer.
- Spirulina supports bone marrow production of white and red blood cells, protecting against the risk of infection.
- By having these ingredients on hand, one can initiate potential protective measures if the need for a CT scan or exposure to other radiation sources arises.

Lemon Balm Extract

Lemon balm extract is known for its ability to preserve foods, particularly meats, from oxidant stress that induces spoilage. This same mechanism helps protect against a similar form of chemical stress induced in the body by ionizing radiation. In fact, one particular study found that lemon balm extract has numerous protective mechanisms:

- Lemon balm can boost levels of the *superoxide dismutase* (SOD), an essential component of the body's native ability to protect itself from the effects of ionizing radiation and other major chemical stresses.¹⁹
- Lemon balm defends lipid cell membranes in living organisms, as shown by a sharp reduction in the lipid peroxidation that is a measure of direct cell damage following radiation exposure.¹⁹
- Lemon balm also protects DNA, as shown by a reduction in the amount of the plasma marker 8-OH-dG, a product of oxidized DNA damage.^{19,21}

A human study documents the benefits of lemon balm in radiation technologists, who are exposed to persistent low-level radiation during their routine work despite taking precautionary measures. For the study, the radiation technologists consumed lemon balm tea (1.5 grams/100 mL) twice daily for 30 days.¹⁹

CT Scans and X-Rays Are Not The Only Radiation Risk

Exposure to radiation released from **nuclear** accidents can be more deadly than a single CT scan.

Most health-conscious people keep a supply of **potassium iodide** on hand in case of a nuclear emergency. When properly ingested, potassium iodide travels to the thyroid gland and saturates it with iodine, thus blocking entrance into the thyroid gland of **radioactive iodine** (e.g. the radionuclide iodine-131).

The thyroid is the most sensitive part of the body to radioactive iodine, but bone marrow and other tissues are also adversely affected and <u>not</u> protected by potassium iodide.

Those concerned with adverse nuclear events may want to keep nutrients close by that have been shown to provide more systemic protection against radiation damage.



The lemon balm tea produced a beneficial *increase* in the activity of natural enzyme systems that fight chemical/oxidant stress, including:

- A **71**% increase in superoxide dismutase (SOD) activity,
- A **12**% increase in glutathione peroxidase (GPX, another native antioxidant molecule) activity,
- A 61% increase in catalase (CAT) activity.

In addition, the lemon balm tea produced a beneficial *decrease* in numerous markers of cellular and DNA damage, including:

- A **31**% decrease in the activity of myeloperoxidase (MPO, an indicator of fat oxidation)
- A 29% decrease in markers of lipid peroxidation (LPO), indicating cell membrane damage,
- A **10**% decrease in 8-OH-dG, a marker of DNA damage.

In other words, prior to supplementation, these technologists, despite careful shielding and limitation of their exposure, were walking around with evidence of the impact of ionizing radiation in their bodies, which was reduced by lemon balm supplementation.

Ginkgo Biloba

Ginkgo biloba is a well-known botanical capable of scavenging the reactive oxygen species that make up the bulk of clastogenic (DNA-breaking) factors produced by radiation.²² Ginkgo biloba reduces the levels of DNA strand breaks that lead directly to cancer.

Lab studies show that irradiation of whole blood from healthy volunteers produced an average of **18** *abnormal* chromosomes per **100** cells, which indicates very high-level clastogenic activity. However, when similar specimens were treated with ginkgo biloba extract, the number of abnormal chromosomes fell to **7.3** per **100** cells—a significant reduction of nearly **60**%.8

The protective effects of ginkgo biloba extract were demonstrated in a unique, if tragic, setting: the 1986 radiation disaster at Chernobyl, in what is now Ukraine.

A 1994 study evaluated blood samples from Armenian workers involved in the initial clean-up of the nuclear reactor, finding an average of 17.9 per 100 cells with chromosomal damage, compared with just **5.7** per **100** in control samples. The same group of researchers treated 30 of those workers with ginkgo biloba extract, containing bioactive flavonoids and terpenoids, at a dose of **40 mg** three times daily (total dose 120 mg per day) for a two-month period.⁷

At the end of the treatment period, the clastogenic (DNA-breakage-inducing) activity of the subjects' plasma fell to control levels. The benefits persisted for at least seven months, but by one year, 33% of the workers again showed elevated clastogenic factors, demonstrating the persistency of the cancer-inducing risk of radiation damage to DNA.

A more recent study demonstrated the protective effects of ginkgo biloba extract in patients undergoing radioactive iodine treatments for Graves' disease (an autoimmune condition involving abnormally high thyroid activity).²³

Radioactive iodine therapy increased chromosomal damage in placebo patients, which peaked at 21 days. In patients supplemented with ginkgo, that early rise in chromosomal damage was followed by a rapid return towards baseline levels. This study revealed a significant increase in DNA damage in the placebo group overall. Interestingly, clastogenic factors never rose significantly above baseline in the ginkgo supplemented subjects.²³

Since radioactive iodine therapy is provided as implanted material, this study demonstrates the potent radioprotective effect of ginkgo biloba extract even in the face of continuous radiation exposure within the body.





Spirulina Extract

Spirulina extract enhances white blood cell production following exposure to ionizing radiation.

The bone marrow is a major site of immediate toxicity from ionizing radiation. Those exposed to very high doses of radiation often end up with fatal, overwhelming infections.^{20,24} Spirulina extract promotes the production of bone marrow-stimulating growth factors such as granulocyte macrophage colony-stimulating factor (GM-CSF).20,24

Another way spirulina helps protect the immune system is by increasing the production of antibodies.²⁵ Antibodies are complex proteins that bind to invading organisms, and flag them for destruction.²⁶ In a study of mice exposed to gamma-irradiation, spirulina polysaccharides containing a molecule called C-phycocyanin stimulated the recovery of white blood cells and bone marrow cell counts suppressed by the radiation exposure.27 In addition, radiation-induced anemia was also suppressed.

These effects translate directly into humans, as shown by a study using lymphocytes (white blood immune system cells) from nuclear power plant workers, which showed that pretreatment of the cells with C-phycocyanin from spirulina stimulated the cells' natural antioxidant systems, protecting them from destruction.²⁸

In a compelling study of youngsters exposed to radioactive fallout from the Chernobyl nuclear plant explosion, a daily 4 gram dose of spirulina (containing the active *C-phycocyanin*) for 21 days produced marked increases in the white and red blood cell counts that had been suppressed by the **radiation**.²⁹ Additionally, elevated levels of inflammatory eosinophils were restored back into the normal range, and anemia, as defined by low hemoglobin levels, was also corrected.

Summary

The use of medical imaging technology, especially CT scans, enables doctors to noninvasively see injuries and diseases deep inside our bodies. But that same technology brings with it a serious threat: *ionizing radiation*.

A recent study showed a single CT scan in youth can increase one's long-term risk for subsequent cancer by **24%**.

Fortunately, specific nutrients with complementary actions may help provide protection against radiation ahead of time. Lemon balm extract prevents formation of dangerous reactive chemicals formed by ionizing radiation exposure, ginkgo biloba extract protects DNA from cancer-causing damage, and spirulina extract supports bone marrow and its production of vital blood cells.

If you need to undergo a CT scan or other potent source of ionizing radiation, start supplementing with these <u>three</u> protective ingredients one to three times daily for <u>five</u> days prior to the scan and for a minimum of <u>five</u> days afterwards. They are now available in a combination formula for consumer convenience.

In addition, regular consumption of **blueberries** or **blueberry extract** markedly <u>enhances</u> **DNA repair**, which can help protect damaged cells from undergoing deadly mutations. •



If you have any questions on the scientific content of this article, please call a Life Extension®

Health Advisor at 1-866-864-3027.

References

- . Storrs C. Do CT scans cause cancer? Sci Am. 2013 Jul;309(1):30-2.
- Canadian Agency for Drugs and Technologies in Health. Radiation Emissions from Computed Tomography: A Review of the Risk of Cancer and Guidelines. Ottawa ON: 2014.
- Westra SJ. The communication of the radiation risk from CT in relation to its clinical benefit in the era of personalized medicine: part 1: the radiation risk from CT. *Pediatr Radiol.* 2014 Oct;44 Suppl 3:515-8.
- Mathews JD, Forsythe AV, Brady Z, et al. Cancer risk in 680,000 people exposed to computed tomography scans in childhood or adolescence: data linkage study of 11 million Australians. *BMJ*. 2013:346:f2360.
- Yu H. Typical cell signaling response to ionizing radiation: DNA damage and extranuclear damage. *Chin J Cancer Res.* Jun 2012:24(2):83-9.
- 6. Broustas CG, Lieberman HB. DNA damage response genes and the development of cancer metastasis. *Radiat Res.* 2014 Feb;181(2):111-30.
- Emerit I, Oganesian N, Sarkisian T, et al. Clastogenic factors in the plasma of Chernobyl accident recovery workers: anticlastogenic effect of Ginkgo biloba extract. *Radiat Res.* 1995 Nov;144(2):198-205.
- Emerit I, Arutyunyan R, Oganesian N, et al. Radiation-induced clastogenic factors: anticlastogenic effect of Ginkgo biloba extract. Free Radic Biol Med. 1995 Jun;18(6):985-91.
- 9. Lindholm C, Acheva A, Salomaa S. Clastogenic plasma factors: a short overview. *Radiat Environ Bioph.* 2010 May;49(2):133-8.
- Emerit I. Clastogenic factors as potential biomarkers of increased superoxide production. *Biomark Insights*. 2007;2:429-38.
- Morgan WF. Is there a common mechanism underlying genomic instability, bystander effects and other nontargeted effects of exposure to ionizing radiation? *Oncogene*. 2003 Oct 13;22(45):7094-9.
- Riklis E, Emerit I, Setlow RB. New approaches to biochemical radioprotection: antioxidants and DNA repair enhancement. Adv Space Res. 1996;18(1-2):51-4.
- Kobashigawa S, Kashino G, Suzuki K, Yamashita S, Mori H. Ionizing radiation-induced cell death is partly caused by increase of mitochondrial reactive oxygen species in normal human fibroblast cells. *Radiat Res.* 2015 Apr:183(4):455-64.
- Agrawal A, Kale RK. Radiation induced peroxidative damage: mechanism and significance. *Indian J Exp Biol.* 2001 Apr:39(4):291-309.
- Luzhna L, Kovalchuk O. Low dose irradiation profoundly affects transcriptome and microRNAme in rat mammary gland tissues. *Oncoscience*. 2014;1(11):751-62.
- Green DE, Rubin CT. Consequences of irradiation on bone and marrow phenotypes, and its relation to disruption of hematopoietic precursors. *Bone.* 2014 Jun:63:87-94.
- Heylmann D, Rodel F, Kindler T, Kaina B. Radiation sensitivity of human and murine peripheral blood lymphocytes, stem and progenitor cells. *Biochim Biophys Acta*. 2014 Aug;1846(1):121-9.
- Lara MS, Gutierrez JI, Timon M, Andres AI. Evaluation of two natural extracts (Rosmarinus officinalis L. and Melissa officinalis L.) as antioxidants in cooked pork patties packed in MAP. *Meat Sci.* 2011 Jul;88(3):481-8.
- Zeraatpishe A, Oryan S, Bagheri MH, et al. Effects of Melissa officinalis L. on oxidative status and DNA damage in subjects exposed to long-term low-dose ionizing radiation. *Toxicol Ind Health*. 2011 Apr;27(3):205-12.
- Zhang HQ, Lin AP, Sun Y, Deng YM. Chemo- and radio-protective effects of polysaccharide of Spirulina platensis on hemopoietic system of mice and dogs. *Acta Pharmacol Sin*. 2001 Dec;22(12):1121-4.

- 21. Schulpis KH, Papassotiriou I, Tsakiris S. 8-hydroxy-2-desoxyguanosine serum concentrations as a marker of DNA damage in patients with classical galactosaemia. Acta Paediatrica. 2006 Feb;95(2):164-9.
- 22. Rong Y, Geng Z, Lau BH. Ginkgo biloba attenuates oxidative stress in macrophages and endothelial cells. Free Radic Biol Med. 1996:20(1):121-7.
- 23. Dardano A, Ballardin M, Ferdeghini M, et al. Anticlastogenic effect of Ginkgo biloba extract in Graves' disease patients receiving radioiodine therapy. J Clin Endocrinol Metab. 2007 Nov;92(11):4286-9.
- 24. Hayashi O, Ono, S., Ishii, K., Shi, Y., Hirahashi, T., Katoh, T. Enhancement of proliferation and differentiation in bone marrow meatopoietic cells by Sirulina (Arthrospira) platensis in mice. J Appl Phycol. 2005;2006(18):47-56.
- 25. Hayashi O, Katoh T, Okuwaki Y. Enhancement of antibody production in mice by dietary Spirulina platensis. J Nutr Sci Vitaminol (Tokyo). 1994 Oct;40(5):431-41.
- 26. Available at: http://www.ncbi.nlm.nih.gov/books/NBK26884/. Accessed September 1, 2015.
- Zhang C-W, Tseng C-T, Zhang YZ. The effect of polysaccharide and phycocyanin from Spirulina platensis var. on peripheral blood and hematopoietic system of bone marrow in mice. Paper presented at the 2nd Asia-Pacific Conference on Algal Biotechnology. Malaysia. 1994.

- 28. Ivanova KG, Stankova KG, Nikolov VN, et al. The biliprotein Cphycocyanin modulates the early radiation response: a pilot study. Mutat Res. 2010 Jan:695(1-2):40-5.
- 29. Loseva LP, Tkatschenko, LW. The corrective influence on the immune system of spirulina platensis as a daily food supplement in general and in case of environmental pollution demonstrated by a study of children who are permanently exposed to low doses of radiation ("Chernobyl Children"). Minsk, Belarus. Clinic for Nuclear Medicine;1999
- 30. Dincer Y, Sezgin Z. Medical radiation exposure and human carcinogenesis-genetic and epigenetic mechanisms. Biomed Environ Sci. 2014 Sep;27(9):718-28.
- 31. Guttikonda R, Herts BR, Dong F, Baker ME, Fenner KB, Pohlman B. Estimated radiation exposure and cancer risk from CT and PET/CT scans in patients with lymphoma. Eur J Radiol. 2014 Jun:83(6):1011-5.
- 32. Kubo T, Ohno Y, Kauczor HU, Hatabu H. Radiation dose reduction in chest CT--review of available options. Eur J Radiol. 2014 Oct:83(10):1953-61.
- 33. Compagnone G, Casadio Baleni M, Di Nicola E, et al. Optimisation of radiological protocols for chest imaging using computed radiography and flat-panel X-ray detectors. Radiol Med. 2013 Jun;118(4):540-54.

CT Scans: A New Public Health Threat

When they were first made available, CT scanners were used only in major, tertiary-care hospital centers, and their use was restricted to only the sickest and most challenging patients. In those years, the benefits of getting a scan clearly outweighed the (then unknown) risks of the radiation exposure.

But in the past two decades, the collective radiation dose from medical images has grown 6-fold, with concomitant growth in the rate of new, preventable cancers.30

CT scans amount to a series of individual X-ray images, organized by a powerful computer to produce a high-contrast 3-D image of body contents. Differences in the "transparency" of tissues to the passage of X-rays produce the patterns that doctors can interpret to highlight individual organs and other structures, readily revealing their inner workings, and also revealing abnormal structures such as tumors, abscesses, and other signs of trouble. But that means that a single CT scan exposes your body to an amount of radiation comparable to that from multiple individual X-rays, thereby raising your cancer risk.

How great is that risk? The best way to understand this is by comparing CT scan radiation exposure to the exposure we constantly receive from outer space (mainly the sun).

Radiation exposure is measured in units called Sieverts (Sv), named after an early radiation researcher. The best estimates are that in one year, a person is naturally exposed to about 3 milli-Sieverts (mSv) from background radiation in space. By contrast, a single

head CT scan produces 2 mSv of exposure, while a full abdominal CT scan may involve more than 30 mSv. or 10-fold the annual natural exposure.2

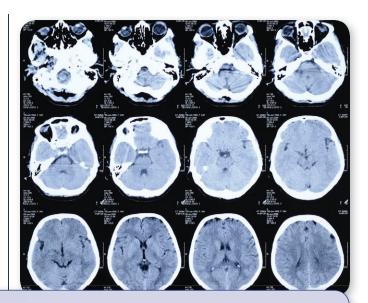
And that's a major concern.

Radiation doses between 5 and 125 mSv are associated with statistically significant increases in cancer risk, readily explaining the tremendous increase in cancer risk seen in the giant British Medical Journal study.2

People who already have a known malignancy are often exposed to even higher radiation levels than healthy individuals because of the frequency of scans for diagnostic and follow-up purposes. One study showed that patients with lymphomas underwent an average of 3.5 CT scans during their treatment periods. a rate that nearly doubled during their post-treatment surveillance period to 6.3 per person.31 The radiation intensity also increased, from an average 39 mSv during treatment to **53 mSv** during surveillance.

The value of frequent CT scans and more importantly, PET/CT scans, in patients undergoing cancer treatment outweighs the radiation risk. In this instance, insurance company cost-containment mandates are denying cancer patients optimal access to PET/CT scanning. The reason that properly read PET/CT scans are so important is that they can detect tiny malignant lesions before symptoms of recurring cancer manifest. This enables the treatment protocol to be adjusted with the objective of eradicating the small tumors before they grow too large.

- 34. Fuchs TA, Stehli J, Bull S, et al. Coronary computed tomography angiography with model-based iterative reconstruction using a radiation exposure similar to chest X-ray examination. Eur Heart J. 2014 May; 35(17):1131-6.
- 35. Rayo MF, Patterson ES, Liston BW, White S, Kowalczyk N. Determining the rate of change in exposure to ionizing radiation from CT Scans: a database analysis from one hospital. J Am Coll Radiol. 2014 Jul;11(7):703-8.
- 36. Yang CC, Liu SH, Mok GS, Wu TH. Evaluation of radiation dose and image quality of CT scan for whole-body pediatric PET/CT: a phantom study. Med Phys. 2014 Sep;41(9):092505.
- 37. Yamao Y, Yamakado K, Takaki H, et al. CT-fluoroscopy in chest interventional radiology: sliding scale of imaging parameters based on radiation exposure dose and factors increasing radiation exposure dose. Clin Radiol. 2013 Feb;68(2):162-6.
- 38. Ramanathan S. Ryan J. Radiation awareness among radiology residents, technologists, fellows and staff: where do we stand? Insights Imaging. 2015 Feb;6(1):133-9.
- 39. Available at: http://www.consumerreports.org/cro/magazine/2015/01/the-surprising-dangers-of-ct-sans-and-x-rays/index. htm. Accessed September 1, 2015.



The Overuse Of CT Scans

With the mounting evidence of the dangers caused by excessive radiation exposure, why isn't the medical community taking major steps to reduce the numbers of scans, or at the very least, to limit the dose of radiation when a scan can't be avoided? There are a few answers, and none of them are encouraging.

First, CT scans are extremely lucrative. Indeed, there are those who question whether physician ownership of radiology operations is to blame for its overuse. By some estimates, 26 to 44% of CT scans ordered are considered inappropriate and unnecessary.2

Technical improvements in CT scanners themselves have in fact permitted a reduction in radiation exposure per scan, but the ease and relatively low cost of such scans has led to a concomitant increase in the number of scans carried out every year, overshadowing the gains made by technology and leaving patients vulnerable.32

There is also simply old-fashioned resistance to change. Most hospitals now use digital radiographic equipment, which makes images with similar technology to the digital camera in your smart phone. The newer equipment can make images with much lower X-ray doses, but few radiology departments have bothered to take advantage of that. 16 Instead, they have simply continued to use the outdated protocols and high radiation levels needed to expose old-fashioned X-ray film. Studies show that substantial reductions in X-ray intensity are possible, which not only don't reduce image quality, but in fact enhance it.33

Radiologists also put pressure on the poorly paid technologists who perform the actual tests, wanting the highest-quality images the first time a patient goes through the scanner. This results in routine use of higher-than-necessary doses of radiation to achieve a sharp image-even though studies show that many CT scans can achieve good-quality studies with ultra-low radiation doses equivalent to a single chest X-ray. 32,34-37

Sadly, and frighteningly, very recent studies reveal that radiology workers (physicians, including those in training, technologists, and others) have poor knowledge about radiation risks, with technologists (those who work daily directly with the equipment that generates X-rays) having the lowest knowledge level.38 These knowledge gaps led directly to a significant underestimation of radiation doses and cancer risks from the kinds of X-rays done on a routine basis.38

Consumers, at least, are beginning to push back, as seen by an article highlighting all these risks in Consumer Reports in 2015.39 But despite the considerable power of consumer groups, the medical establishment is always slow to change-and your options are limited in the meantime.

You can and should ask that the lowest possible radiation dose be used when you must undergo a CT scan or other ionizing radiation-producing test, but that can be a difficult task for most of us not versed in medical terminology, and not understanding the multiple layers of decision-making in a large hospital or medical center.

Rather than skipping a potentially lifesaving test, and in addition to standing up for your own safety at the radiology suite, you can be nutritionally prepared ahead of time by loading your body with three specific nutrients known to be capable of mitigating the damage from ionizing radiation: lemon balm, ginkgo biloba, and spirulina.

JRINE

BOOSTS BRAIN CELL REGENERATION & SUPPORTS WHOLE-BODY HEALTH

Taurine, a free amino acid, has been described by scientists as "one of the most essential substances in the body." But as we age, taurine levels decline.

Cognitive Function and Brain Cell Regeneration

The benefits of taurine on brain cell growth are especially evident in those with a taurine deficiency, which includes aging individuals.

Promising research has found that taurine can promote **new brain cell formation** in the area of the brain associated with **learning** and **memory**. It does so by activating hibernating stem cells that are capable of growing into several different kinds of cells.2

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

Whole-Body Health

Past research has also shown the ability of taurine to maintain and support: 3-9

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

One capsule taken one to three times daily on an empty stomach, or as recommended by a health care practitioner, supports optimal cardiovascular levels.

- Mol Vis. 2012;18;2673-86. Stem Cell Res. 2015 May;14(3):369-79. 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.

- 6. J Biomed Sci. 2010 Aug 24;17 Suppl1:S1. 7. Amino Acids. 2008 Aug;35(2):469-73. 8. Amino Acids. 2012 Nov;43(5):1979-93. 9. Neurosci Lett. 2006 May 15;399(1-2):23-6.

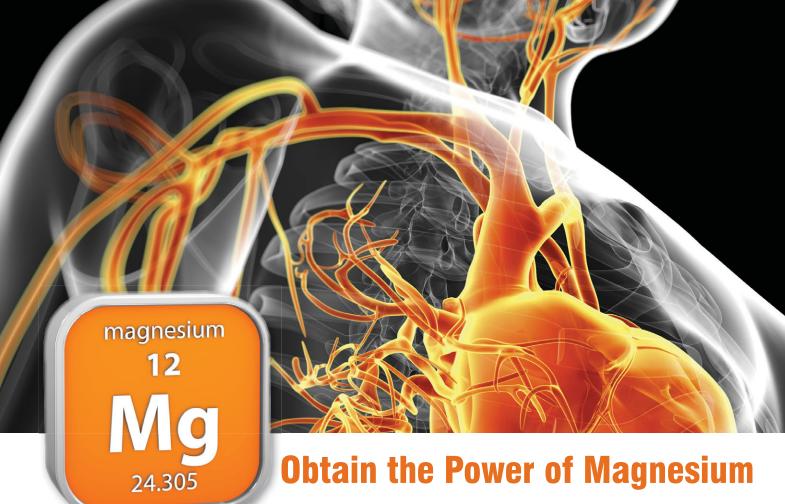


Item #01827 • 90 vegetarian capsules

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

Non-GMO

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



Are You Deficient in The Body's #1 Mineral?

Magnesium is the **most important** mineral in the body, yet most Americans do not obtain sufficient magnesium from their diet.

Magnesium is required for more than **300 biochemical reactions** and many of the body's critical functions are dependent upon it. Magnesium helps:^{1,2}

- Maintain normal muscle and nerve function.
- Keep **heart rhythm** steady.
- Support a healthy immune system.
- Keep bones strong.
- Maintain **blood sugar levels** already within normal range.
- Promote normal **blood pressure**. Magnesium is also...
- Involved in energy metabolism and protein synthesis.

The recommended intake of magnesium to maintain vascular health is **500 mg** or more a day. With **Life Extension® Magnesium Caps**, you can easily obtain **500 mg** of elemental magnesium for less than **7 cents a day!**

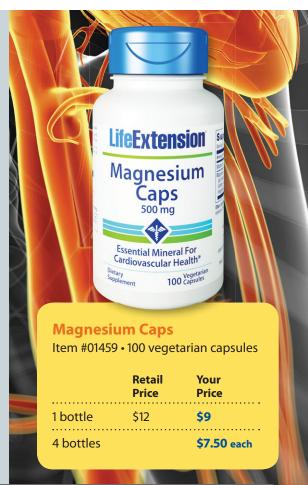
Non-GMO

Caution: If taken in high doses, magnesium may have a laxative effect. If this occurs, divide dosing, reduce intake, or discontinue use.

References

1. Am J Clin Nutr. 1987;45:1305-12. 2. Clinica Chimica Acta. 2000;294:1-26.

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



X-R Shield Maintain Cellular Integrity



From solar *ultraviolet* to *cosmic radiation* to *X-rays* from medical diagnostic imaging procedures like CT scans, *ionizing radiation* consists of waves and particles of sufficient energy to disturb atoms, "knocking out" electrons to create *ions*.

Also known as *free radicals*, these ions have the potential to interact with living tissues and disturb cell structures, including fragile DNA.

In addition to limiting and/or avoiding excessive ionizing radiation exposure from various sources (e.g. recurrent transcontinental plane travel and frequent medical diagnostic imaging procedures like CT and/or PET scans) whenever reasonable, additional cellular support during *ionizing radiation* exposure and *free radical* generation is helpful for optimal health.

A review of the published scientific literature reveals that certain nutrients may help support and nourish the body's cellular integrity in the context of **ionizing** radiation.

NEW X-R SHIELD FORMULA

The new **X-R Shield** provides <u>three</u> nutrients that help support and maintain cellular integrity¹⁻⁵:

LEMON BALM supports production of *superoxide dismutase* (SOD), *glutathione peroxidase*, and *catalase*—your body's natural defense against oxidative damage.¹They provide critical support by inhibiting the formation of dangerous free radicals¹²—supporting your body's natural defense of critical DNA.

GINKGO BILOBA supports the body's capacity to scavenge reactive oxygen species, which can interact with DNA.³ This further helps support your body's natural defense system in the context of DNA.⁴

ORGANIC SPIRULINA supports immune system production of white blood cells in the bone marrow.⁵ These immune cells are critical for a healthy immune response.

Prior to *ionizing radiation* exposure (e.g. transcontinental plane travel or medical diagnostic imaging procedures consider taking three capsules of **X-R Shield** one to three times daily beginning at least <u>five days</u> beforehand and continue taking until the bottle is finished

X-R Shield

Item #01919 • 90 vegetarian capsules

| | Retail Price | Your Price |
|-----------|-----------------|--------------------|
| 1 bottle | \$15 | \$11.25 |
| 4 bottles | | \$9.75 each |



References

- 1. Toxicol Ind Health. 2011 Apr;27(3):205-12.
- 2. Meat Sci. 2011 Jul;88(3):481-8.
- 3. Free Radic Biol Med. 1996;20(1):121-7.
- Free Radic Biol Med. 1995 Jun;18(6):985-91.
 Acta Pharmacol Sin. 2001 Dec;22(12):1121-4.

To order X-R Shield, call 1-800-544-4440 or visit www.LifeExtension.com



CHLOROPHYLLIN Protects Against Environmental Toxins

Every day, we are surrounded by environmental toxins. Studies show they can contribute to diabetes, heart disease, and cancer through induction of **mutations** to our DNA.¹⁻⁵

Fortunately, an inexpensive supplement—chlorophyllin—has been shown in multiple studies to detoxify these deadly chemicals, and protect DNA.

Proper use of **chlorophyllin** affords considerable defense against the natural and man-made toxins that permeate our food and environment.⁶⁻⁸



Environmental Toxins: An Unfortunate Fact Of Life

In the last 70 years, more than 80,000 new chemicals have been synthesized, and every year, over 4 **billion** pounds of these chemicals, many known carcinogens, are released into the environment.9 Even more disturbing is that the vast majority of these chemicals have never been adequately tested by any government agency, including the **EPA** and **FDA**, in regards to their effects on human health.¹⁰

On a daily basis, nearly everyone is exposed to this barrage of compounds through industrial and manufacturing facilities, agricultural runoff that includes pesticides and herbicides laced in foods, and emissions from trucks, cars, and planes. Even our brave military men and women face exposure from a multitude of chemicals used on bases and facilities around the world.

A study from the Centers for Disease Control and **Prevention** (CDC) published in 2009 confirmed these findings by examining the burden of 212 industrial chemicals in the bodies of US citizens.¹¹ In particular, the CDC report showed widespread exposure to commonly used industrial chemicals, including polybrominated diphenyl ethers (PBDEs) and bisphenol A (BPA), which is a chemical known for its toxic effects on reproduction.¹² BPA is reported to be present in at least 90% of the US population. Perfluoroalkyl chemicals are present in 98% of people sampled. 11-13 Perfluorooctanoic acid (PFOA), a type of perfluoroalkyl chemical, is best known for being part of the substance TeflonTM and is linked with multiple diseases, including ulcerative colitis, 14 kidney disease, 15 thyroid disease, 16 and cancer.17

Chlorophyllin: Protection Against Toxins

Chlorophyllin is a water-soluble derivative of the green plant substance *chlorophyll*.

In multiple studies, chlorophyllin exhibits powerful anticarcinogenic effects in regards to a variety of environmental toxins. 6-8 The way chlorophyllin accomplishes this is by targeting a number of molecules and pathways involved in cancer development, such as protecting against *mutations* of the p53 tumor sup**pressor gene**. Cell proliferation is partially controlled by the **p53 gene**, so protecting healthy expression of p53 is a critical factor that helps guard against cancerous changes.18

Chlorophyllin disables potent carcinogens such as polycyclic aromatic hydrocarbons and heterocyclic *amines* by forming complexes with these chemicals that limit the ability of these toxins to bind to normal cells to inflict malignant changes. 19-20

Chlorophyllin binds to a number of other common carcinogens including dibenzanthracene, dibenzopyrene and benzophenanthrene and limits their ability to form **DNA adducts**, which are strands of DNA bonded to cancer-causing toxins. Formation of DNA adducts is an early step in the formation of cancer-causing DNA mutations. **Chlorophyllin** limits DNA adduct formation caused by a variety of known carcinogens.²¹⁻²³

Chlorophyllin also functions as an antimutagenic agent against a number of prevalent chemicals in the environment such as **PhlP** (2-amino-1-methyl-6phenylimidazo[4,5-b]pyridine), known to be involved in colon, prostate, pancreatic, and breast cancer.²³⁻²⁴

Chlorophyllin Protects Against Multiple Cancers

The ways in which **chlorophyllin** provides protection against cancer—and the number of cancers it's effective against have been shown in multiple studies including:

Oral Cancer: A 2012 study in hamsters showed that chlorophyllin can both prevent and reverse genetic mutations linked to oral cancer caused by the chemical 7,12 dimethylbenzanthracene (DMBA).²⁵ Scientists in this study conclusively showed that chlorophyllin modulated and even reversed changes to 104 genes that were deleteriously affected by DMBA.

Pancreatic Cancer: A study published in 2014 examined the effects of a number of dietary supplements on pancreatic cell lines, with the results showing that chlorophyllin has antiproliferative effects on pancreatic cells.²⁶

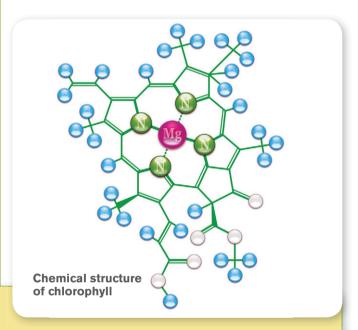
Colon Cancer: In a study on human colon cancer cells, researchers showed that chlorophyllin-treated colon cancer cells had a statistically decreased expression of human ribonucleotide reductase (RR), an enzyme used by these cancer cells for DNA synthesis and repair. By inhibiting RR, chlorophyllin made these cancer cells less able to grow and more receptive to mainstream cancer chemotherapeutic agents.²⁷

What You Need To Know

Chlorophyllin **Protects Against Environmental Toxins**

- Chlorophyllin provides protection against environmental toxins.
- Chlorophyllin is a safe, inexpensive supplement derived from the plant substance chlorophyll. It helps stop the damage done to DNA by environmental chemicals and toxins that are present in nearly everyone.
- Laboratory studies have shown that chlorophyllin is able to bind to and remove cancercausing toxins in the body.
- Chlorophyllin targets multiple molecules and pathways involved in cancer development brought about by environmental toxin exposure, such as mutations in the p53 tumor suppressor gene so that these mutations do not occur and raise the risk of cancer.

Bladder Cancer: In a 2014, researchers examined the combined effects of chlorophyllin with photodynamic therapy (PDT), a minimally invasive cancer treatment in which cancer cells take up a chemical or substance and are then exposed to a certain wavelength of light. In the presence of the photosensitizing substance, the PDT generates a form of oxygen that kills the cancer cells. In this study the researchers, using a special form of chlorophyllin, showed that when combined with photodynamic therapy (PDT), it resulted in approximately 85% destruction of bladder cancer cells.28



- The ability of cancer-causing toxins to form DNA adducts is significantly decreased in the presence of chlorophyllin.
- Chlorophyllin has been shown in numerous studies to protect against multiple forms of cancer, including bladder, breast, pancreatic, and colon cancer.
- Aflatoxins, the cause of a significant number of liver and other cancers around the world, is significantly reduced in the presence of chlorophyllin.
- Chlorophyllin is a potent free radical scavenger and has been shown in studies to protect mitochondria against oxidative damage.
- Chlorophyllin offers optimal protection against cancer-causing DNA and mitochondrial damage in a world contaminated with toxic chemicals.

Breast Cancer: Researchers at the University of Kentucky showed that when human breast cells were exposed to dibenzopyrene (DBP), a known carcinogen, dangerous and unstable DNA adducts were formed. However, in the presence of chlorophyllin a decrease of more than 65% in the number of DNA adducts was observed.29

Stomach Cancer: A 2014 study examined the effects of chlorophyllin on stomach cancer in rats and more specifically, the effects on transforming growth factor (TGF) beta signaling, a pathway that plays a vital role in cancer formation.³⁰ Chlorophyllin showed strong effects in stopping cancerous growth and metastasis, leading the authors of the study to conclude: "*Dietary* chlorophyllin that simultaneously abrogates TGF beta signaling pathway and the key hallmark events of cancer appears to be an ideal candidate for cancer chemoprevention."

Chlorophyllin Protects Against Liver Cancer

Out of the 600,000 new cases of liver cancer diagnosed yearly around the world, conservative estimates are that up to 155,000 of them are caused by aflatoxin found in multiple improperly stored foods including peanuts, corn, pistachios, and rice.31

Fortunately, studies show that **chlorophyllin** has the potential to significantly reduce the risk of liver cancer induced by aflatoxin by binding to carcinogenic byproducts of aflatoxin metabolism and therefore, decreasing bioavailability of these cancer-causing chemicals.³²⁻³³

In a study in the journal *Cancer Prevention Research*, researchers showed that in volunteers, ingestion of



150 mg of chlorophyllin significantly decreased the absorption of aflatoxin B.32

An even more compelling study on the preventive power of chlorophyllin was done in Oidong, China, a province where the population is known to have a high exposure to aflatoxins and therefore, putting them at high risk of liver cancer.³³ In this randomized, double-blind, placebo-controlled study, 180 men and women were randomized to receive either 100 mg of chlorophyllin or a placebo three times daily over a four-month period.

The researchers showed that those men and women randomized to the chlorophyllin arm of the study had a 55% reduction in a biochemical marker of aflatoxin when compared to those people not taking chlorophyllin. This significant finding led the authors of this study to conclude "...interventions with chlorophyllin or supplementation of diets with foods rich in chlorophylls may represent practical means to prevent the development of hepatocellular carcinoma or other environmentally induced cancers."

Aflatoxin: Nature's Own Deadly Toxin

It's not just industrial chemicals that increase the risk for cancer. There are also toxic compounds found in nature. Aflatoxin, a biochemical in the class of substances called mycotoxin, is produced by molds and is one of the most common and deadly carcinogens. While aflatoxin is associated with being a cause of liver cancer³⁹⁻⁴¹ through the formation of carcinogenic by-products and its ability to cause DNA mutations, studies show that aflatoxin is also the cause of other cancers. 42-44

An article in **JAMA** presented documentation that aflatoxin is a cause of gallbladder cancer. 42 In this case-control study conducted in Chile, researchers presented strong evidence that aflatoxin, known to cause gallbladder cancer in primates, is a cause of gallbladder cancer-the leading cause of cancer deaths in women in Chile-stating "...the associations between aflatoxin exposure and gallbladder cancer were statistically significant." A more recent article published in 2015 in the journal Tumour Biology showed evidence that aflatoxin is both a potential cause of lung cancer,55 and stimulates lung cancer cell migration, or metastasis.43

Chlorophyllin Shields Mitochondria

Another way that chlorophyllin provides protection against environmental toxins is via its free radical scavenging activities, both in general and specifically in mitochondria. In a study published in 2000, researchers looked at the protective effects of chlorophyllin on various organs and mitochondrial membranes in mice and rats.34 The authors of this study showed that not only did chlorophyllin protect mice brains, liver, and testes from oxidative damage, but it also proved to be a potent free radical scavenger in rat liver mitochondria, leading the researchers to conclude "...our studies showed that CHL [chlorophyllin] is a highly effective antioxidant, capable of protecting mitochondria against oxidative damage...'

In another study using rat liver mitochondria, researchers showed that chlorophyllin provided significant protection against ionizing radiation, a potent generator of free radicals.35 Finally, with mitochondrial damage strongly implicated in the aging process, 36-38 chlorophyllin may very well be useful in preventing chronic diseases and premature senescence.

Summary

Life Extension® described the antimutagenic properties of chlorophyllin in the mid-1980s.

There was ample evidence decades ago that taking **100 mg** of **chlorophyllin** with the heaviest meal of the day made sense because larger meals tend to contain more dietary mutagens.

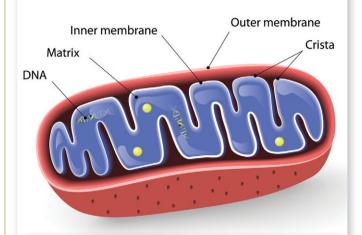
A popular formula used by health-conscious people today provides 100 mg of chlorophyllin along with other nutrients that are best taken with a heavy meal. If you ingest more than one large meal daily or are exposed to excessive dietary or environmental toxins, you may consider taking 100 mg of chlorophyllin with each meal.

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

References

- Fosslien E. Cancer morphogenesis: role of mitochondrial failure. Ann Clin Lab Sci. 2008 Autumn;38(4):307-29.
- Auger C, Alhasawi A, Contavadoo M, et al. Dysfunctional mitochondrial bioenergetics and the pathogenesis of hepatic disorders. Front Cell Dev Biol. 2015 Jun 25;3:40.
- Lim, S, Cho YM, Park KS, et al. Persistent organic pollutants, mitochondrial dysfunction, and metabolic syndrome. Ann NY Acad Sci. 2010 Jul;1201:166-76.
- Jia G, Aroor AR, Martinez-Lemus LA, et al. Mitochondrial functional impairement in reponse to environmental toxins in the cardiorenal metabolic syndrome. Arch Toxicol. 2015 Feb;89(2):147-53.

MITOCHONDRION



Environmental Toxins And Their Destructive Effects On Mitochondria

One way in which toxins may bring about the myriad of damage that they cause is through their destructive effects on mitochondria, microscopic organelles found in every human cell that convert the oxygen and food into energy. There are multiple studies that provide evidence that environmental toxins are poisons to these life-giving organelles and lead to multiple diseases including cancer,1-2 metabolic syndrome, insulin resistance,3-5 Alzheimer's disease,45 Parkinson's disease,46 chronic kidney disease,47 and cardiovascular disease.4

Environmental toxins damage mitochondria in multiple ways. 48-51 **Chlorophyllin** helps neutralize these poisons.



Environmental Toxins And Cancer

Despite the claims put out by national organizations like the American Cancer Society, we are not winning the war on cancer. Far from it. A report sent to President Obama in 2010 by the Presidents Cancer Panel, which over the period between September 2008 and January 2009 took testimony from 45 experts from academia, government, and industry regarding environmental toxins and cancer.⁵² In the panel's conclusion, they stated "...the true burden of environmentally induced cancer has been grossly underestimated. With nearly 80,000 chemicals on the market in the United States, many of which are...understudied and largely unregulated, exposure to potential environmental carcinogens is widespread." While research is on-going, recent studies on how environmental chemicals cause cancer include damaging changes to DNA and increasing cellular oxidative stress leading to unchecked free radical formation.53-54

- 5. Lee HK. Mitrochondrial dysfunction and insulin resistance: the contribution of dioxin-like substances. Diabetes Metab J. 2011 Jun:35(3):207-15.
- Ferguson LR, Philpott M, Karunasinghe N. Dietary cancer and prevention using antimutagens. Toxicology. 2004 May 20;198(1-
- 7. Williams DE. The rainbow trout liver cancer model: response to environmental chemicals and studies on promotion and chemoprevention. Comp Biochem Physiol C Toxicol Pharmacol. 2012 Jan;155(1):121-7.
- Nagini S, Palitti F, Natarajan AT. Chemopreventive potential of chlorophyllin: A review of the mechanisms of action and molecular targets. Nutr Cancer. 2015;67(2):203-11.
- 9. Available at: http://www.mountsinai.org/patient-care/service-areas/ children/areas-of-care/childrens-environmental-health-center/childrens-disease-and-the-environment/children-and-toxic-chemicals. Accessed September 11, 2015.
- 10. Thornton JW, McCally M, Houlihan J. Biomonitoring of industrial pollutants: health and policy implications of the chemical body burden. Public Health Rep. 2002 Jul-Aug;117(4):315-23.
- 11. Available at: http://www.cdc.gov/exposurereport/. Accessed September 11, 2015.
- 12. Available at: http://www.cdc.gov/biomonitoring/BisphenolA_Fact-Sheet.html. Accessed September 11, 2015.
- 13. Calafat AM, Wong LY, Kuklenyik Z, at al. Polyfluoroalkyl chemicals in the U.S. population: data from the National Health and Nutrition Examination Survey (NHANES) 2003-2004 and comparisons with NHANES 1999-2000. Environ Health Perspect. 2007 Nov;115(11):1596-602.
- 14. Steenland K1, Zhao L, Winquist A, et al. Ulcerative colitis and perfluorooctanoic acid (PFOA) in a highly exposed population of community residents and workers in the mid-Ohio valley. Environ Health Perspect. 2013 Aug;121(8):900-5.
- 15. Watkins DJ, Josson J, Elston B, et al. Exposure to perfluoroalkyl acids and markers of kidney function among children and adolescents living near a chemical plant. Environ Health Perspect. 2013 May;121(5):625-30.



- 16. Lopez-Espinosa MJ, Mondal D, Armstrong B, et al. Thyroid function and perfluoroalkyl acids in children living near a chemical plant. Environ Health Perspect. 2012 Jul;120(7):1036-41.
- 17. Barry V, Winquist A, Steenland K. Perfluorooctanoic acid (PFOA) exposures and incident cancers among adults living near a chemical plant. Environ Health Perspect. 2013 Nov-Dec:121(11-12):1313-8.
- 18. Gouda EM, Elbehairy AM, Ghoneim MA. Antimutagenic efficacy of some natural compounds on cyclophosphamide-induced p53 alterations. Z Naturforsch C. 2008 Nov-Dec;63(11-12):857-63.
- 19. Hayatsu H. Complex formation of heterocyclic amines with porphyrins: its use in detection and prevention. Princess Takamatsu Symp. 1995:23:172-80.
- 20. John K, Keshava C, Richardson DL, et al. Immune Response Signatures of Benzo() pyrene Exposure in Normal Human Mammary Epithelial Cells in the Absence or Presence of Chlorophyllin. Cancer Genomics Proteomics. 2009 Jan-Feb;6(1):1-11.
- 21. Lagerqvist A1, Håkansson D, Frank H, et al. Structural requirements for mutation formation from polycyclic aromatic hydrocarbon dihydrodiol epoxides in their interaction with food chemopreventive compounds. Food Chem Toxicol. 2011 Apr;49(4):879-86.
- 22. John K, Divi RL, Keshava C, et al. CYP1A1 and CYP1B1 gene expression and DNA adduct formation in normal human mammary epithelial cells exposed to benzo[a]pyrene in the absence or presence of chlorophyllin. Cancer Lett. 2010 Jun 28:292(2):254-60.
- 23. Mata JE, Yu Z, Gray JE, et al. Effects of chlorophyllin on transport of dibenzo(a, l)pyrene, 2-amino-1-methyl-6-phenylimidazo-[4,5-b] pyridine, and aflatoxin B(1) across Caco-2 cell monolayers. Toxicology. 2004 Mar 1;196(1-2):117-25.
- 24. Hirose M, Nishikawa A, Shibutani M, et al. Chemoprevention of heterocyclic amine-induced mammary carcinogenesis in rats. Environ Mol Mutagen. 2002;39(2-3):271-8.
- 25. Vidya Priyadarsini R, Kumar N, Khan I, et al. Gene expression signature of DMBA-induced hamster buccal pouch carcinomas: modulation by chlorophyllin and ellagic acid. PLoS One. 2012;7(4):e34628.
- 26. Konickova R, Vankova K, Vanikova J, et al. Anti-cancer effects of blue-green alga Spirulina platensis, a natural source of bilirubin-like tetrapyrrolic compounds. Ann Hepatol. 2014 Mar-Apr:13(2):273-83.
- 27. Chimploy K, Diaz GD, Li Q, et al. E2F4 and ribonucleotide reductase mediate S-phase arrest in colon cancer cells treated with chlorophyllin. International journal of cancer. Int J Cancer. 2009 Nov 1;125(9):2086-94.
- 28. Lihuan D, Jingcun Z, Ning J, et al. Photodynamic therapy with the novel photosensitizer chlorophyllin f induces apoptosis and autophagy in human bladder cancer cells. Lasers Surg Med. 2014 Apr:46(4):319-34.
- Smith WA, Freeman JW, Gupta RC. Effect of chemopreventive agents on DNA adduction induced by the potent mammary carcinogen dibenzo[a,l]pyrene in the human breast cells MCF-7. Mutat Res. 2001 Sep 1;480-481:97-108.
- 30. Thiyagarajan P, Kavitha K, Thautam A, et al. Dietary chlorophyllin abrogates TGFbeta signaling to modulate the hallmark capabilities of cancer in an animal model of forestomach carcinogenesis. Tumour Biol. 2014 Jul;35(7):6725-37.
- 31. Liu Y, Wu F. Global burden of aflatoxin-induced hepatocellular carcinoma: a risk assessment. Environmental health perspectives. Environ Health Perspect. 2010 Jun;118(6):818-24.
- 32. Jubert C, Mata J, Bench G, et al. Effects of chlorophyll and chlorophyllin on low-dose aflatoxin B(1) pharmacokinetics in human volunteers. Cancer Prev Res (Phila). 2009 Dec;2(12):1015-22.
- 33. Egner PA, Wang JB, Zhu YR, et al. Chlorophyllin intervention reduces aflatoxin-DNA adducts in individuals at high risk for liver cancer. Proc Natl Acad Sci U S A. 2001 Dec 4;98(25):14601-6.
- 34. Kamat JP, Boloor KK, Devasagayam TP. Chlorophyllin as an effective antioxidant against membrane damage in vitro and ex vivo. Biochim Biophys Acta. 2000 Sep 27;1487(2-3):113-27.
- 35. Boloor KK, Kamat JP, Devasagayam TP. Chlorophyllin as a protector of mitochondrial membranes against gamma-radiation and photosensitization. Toxicology. 2000 Nov 30;155(1-3):63-71.

- 36. Zapico SC, Ubelaker DH. Relationship Between Mitochondrial DNA Mutations and Aging. Estimation of Age-at-death. J Gerontol A Biol Sci Med Sci. 2015 Aug 18.
- 37. Dai DF, Chiao YA, Marcinek DJ, Szeto HH, Rabinovitch PS. Mitochondrial oxidative stress in aging and healthspan. Longev Healthspan. 2014 May 1;3:6.
- 38. Ross JM. Olson L. Coppotelli G. Mitochondrial and ubiquitin proteasome system dysfunction in ageing and disease: two sides of the same coin? Int J Mol Sci. 2015 Aug 17;16(8):19458-76.
- 39. Smith RJ. Nutrition and metabolism in hepatocellular carcinoma. Hepatobiliary Surg Nutr. 2013 Apr;2(2):89-96.
- 40. Kar P. Risk factors for hepatocellular carcinoma in India. J Clin Exp Hepatol. 2014 Aug;4(Suppl 3):S34-42.
- 41. Nault JC. Pathogenesis of hepatocellular carcinoma according to aetiology. Best Pract Res Clin Gastroenterol. 2014 Oct;28(5):937-47.
- 42. Nogueira L. Foerster C. Groopman J. et al. Association of aflatoxin with gallbladder cancer in Chile. JAMA. 2015 May 26;313(20):2075-7.
- 43. Cui A, Hua H, Shao T, et al. Aflatoxin B1 induces Src phosphorylation and stimulates lung cancer cell migration. Tumour Biol. 2015 Aug:36(8):6507-13.
- 44. Sieber SM, Correa P, Dalgard DW, Adamson RH. Induction of osteogenic sarcomas and tumors of the hepatobiliary system in nonhuman primates with aflatoxin B1. Cancer Res. 1979 Nov:39(11):4545-54.
- 45. Moulton PV, Yang W. Air pollution, oxidative stress, and Alzheimer's disease. J Environ Public Health. 2012;2012:472751.
- 46. Pan-Montojo F, Reichmann H. Considerations on the role of environmental toxins in idiopathic Parkinson's disease pathophysiology. Transl Neurodegener. 2014 May 9;3:10.
- 47. Kim NH, Hyun YY, Lee KB. Environmental heavy metal exposure and chronic kidney disease in the general population. J Korean Med Sci. 2015 Mar;30(3):272-7.
- 48. Ames BN, Shigenaga MK, Hagen TM. Oxidants, antioxidants, and the degenerative diseases of aging. Proc Natl Acad Sci U S A. 1993 Sep 1;90(17):7915-22.
- 49. Barja G. Free radicals and aging. Trends Neurosci. 2004 Oct;27(10):595-600.
- 50. Junqueira VB, Barros SB, Chan SS, et al. Aging and oxidative stress. Mol Aspects Med. 2004 Feb-Apr;25(1-2):5-16.
- 51. Ashok BT, Ali R. The aging paradox: free radical theory of aging. Exp Gerontol. 1999 Jun;34(3):293-303.
- 52. Available at: http://deainfo.nci.nih.gov/advisory/pcp/annualReports/pcp08-09rpt/PCP_Report_08-09_508.pdf. Accessed September 14, 2015.
- 53. Erkekoglu P, Kocer-Gumusel B. Genotoxicity of phthalates. Toxicol Mech Methods. 2014 Dec;24(9):616-26.
- 54. Kuppusamy SP, Kaiser JP, Wesselkamper SC. Epigenetic Regulation in Environmental Chemical Carcinogenesis and its Applicability in Human Health Risk Assessment. Int J Toxicol. 2015 Aug
- 55. Van Vleet TR, Watterson TL, Klein PJ, et al. Aflatoxin B1 alters the expression of p53 in cytochrome P450-expressing human lung cells. Toxicol Sci. 2006 Feb;89(2):399-407.



Throughout the body, omega-3 fatty acids support heart health, brain health, eye health, mood, the immune system, and much more.

Taking fish oil alone may not be enough to obtain optimal omega-3 benefits. Research indicates it is ideal to obtain omega-3s from multiple sources—such as fish oil and krill—since they target different cellular signaling pathways.

Life Extension®'s Super Omega with Krill, Astaxanthin, & Olive Fruit Extract is a unique formulation containing a potent concentration of 5 key ingredients that work together to maximize the protective benefits of omega-3s.

KRILL AND FISH OIL

Scientists suggest that combining **fish oil** and **krill oil** may provide enhanced benefits for the cardiovascular and nervous system more than either form by itself due to the different ways that fish oil and krill oil are taken up by our cells.1,2

OLIVE FRUIT EXTRACT

Olive polyphenols improve blood lipids and endothelial function that support arterial health along with other cardiovascular benefits.2,3

The daily dose of **Super Omega** provides the equivalent polyphenol content of 4 to 6 tablespoons of extra virgin olive oil.

ASTAXANTHIN

In Super Omega, the natural astaxanthin content of krill is fortified with additional astaxanthin for maximum stability and benefit.

Astaxanthin fights free radicals, 4 boosts mitochondrial function, 4 supports brain,^{5,6} cardiovascular,^{7,8} and DNA^{9,10} health, and supports healthy blood sugar levels for those already in the normal range.11,12

SESAME LIGNANS

The **Super Omega** formula contains standardized **sesame lignans** to enhance the overall benefits of EPA/DHA marine oils. 13,14

Sesame lignans help support a healthy inflammatory response and guard against lipid peroxidation, thereby extending the stability of DHA in the body.

Experts believe that taking both fish oil and krill oil together offers overlapping coverage to powerfully boost your health with benefits that add to and amplify those of fish oil alone.

Super Omega with Krill, **Astaxanthin, & Olive Fruit Extract**

Item #01988 • 120 softgels

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$45 | \$33.75 |
| 4 bottles | | \$31.50 each |



/DHA with Sesame Ligna Olive Extract, Krill & Astaxanthin

Non-GMO

References

- 1. Available at: http://www.jlr.org/ content/40/10/1867.long# xref-ref-23-1. Accessed March 24,
- 2. Lipids. 2011 Jan;46(1):37-46.
- J Photochem Photobiol B. 2006 Dec 1;85(3):205-15.
- J Nutr Biochem. 2010 May;21(5):381-9. 5. BMC Neurosci. 2012 Dec 29;13(1):156.
- 6. FASEB J. 2009 Jun;23(6):1958-68.
- 7. Arzneimittelforschung. 2011;61(4):239-46.
- 8. Nutr Res. 2011 Oct;31(10):784-9.
- 9. Nutr Metab (Lond), 2010;7:18. 10. Chem Biol Interact. 2011 Aug 15:193(1):79-87.
- 11. Biofactors. 2004;20(1):49-59.
- 12. Med Hypotheses. 2011 Oct;77(4):550-6.
- 13. Metabolism. 2006 Mar;55(3):381-90. 14. Biochem Biophys Acta. 2004 Jun
- 1;1682 (1-3):80-91.

To order Super Omega with Krill, Astaxanthin, & Olive Fruit Extract, call 1-800-544-4440 or visit www.LifeExtension.com

PROTECT D AGAINST CHEMICAL ASSAULT



On a daily basis, the **DNA** in each cell of your body is bombarded by an estimated 10,000 destructive oxidative hits. Environmental toxins, such as pesticides, cigarette smoke, coal dust, and diesel emission particles, challenge the integrity of your DNA and can profoundly modify outcomes of aging.^{2,3}

Supports Healthy DNA*

100 Vegetaria

Foods cooked at high temperatures also threaten cellular integrity. Deep-fried foods along with well-done beefsteak, hamburgers, and bacon trigger the formation of gene-threatening heterocyclic amines.^{4,5} Even "healthy" foods can contain small amounts of undesirable substances.6

To meet this challenge, **Chlorophyllin** provides a water-soluble form of **chlorophyll** shown to protect DNA.^{7,8} It binds to certain toxic chemicals, allowing the body to safely eliminate them before they can enter the bloodstream.

Chlorophyllin also helps the body neutralize all major oxygen species and acts to protect mitochondria.^{9,10}

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

Chlorophyllin Item #01571 • 100 vegetarian capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$24 | \$18 |
| 4 bottles | | \$15 each |

The suggested dose is to take one **Chlorophyllin** capsule with each meal that contains mutagenic byproducts. Each capsule provides:

100 mg Chlorophyllin

(from sodium magnesium chlorophyllin)

Non-GMO

- **References**1. *Proc Natl Acad Sci USA*. 1993 Sep 1;90(17):7915-22.

- Proc Nati Acad Sci USA. 1993 Sep 1;90(17):7915-22.
 Science. 1997 Oct 17;278(5337):407-11.
 Pharm Res. 2008 Sep;25(9):2097-116.
 Cancer Sci. 2004 Apr;95(4):290-9.
 Cancer Epidemiol Biomarkers Prev. 2001 May;10(5):559-62.
 Proc Natl Acad Sci USA. 1990 Oct;87(19):7777-81.
 Environ Mol Mutagen. 1997, 30(4): 468-74.
 Toxicology. 2000 Nov 30;155(1-3):63-71.
 Free Radic Res. 2001 Nov;35(5):563-74.
 B Rinchim Biophys Acta 2000 Sep 27:1487(2-3):113-27.

- 10.*Biochim Biophys Acta*. 2000 Sep 27;1487(2-3):113-27.

FOR WHOLE-BODY SUPPORT

Highly Absorbable CURCUMIN

Used medicinally for over 4,000 years, curcumin benefits almost every organ in the body.¹⁻³ The challenge in obtaining these benefits is that most supplements are poorly **absorbed** into the bloodstream and are not well retained in the body.

Life Extension e's curcumin supplements utilize a patented, bio-enhanced curcumin preparation that can reach up to 7 times higher concentrations in the blood than standard curcumin.4

Studies comparing standard curcumin to Super Bio-Curcumin® and Advanced Bio-Curcumin® with Ginger & Turmerones found:5,6

- Nearly 2 times the support for immune health,
- Nearly twice the support for healthy inflammatory response, and
- Approximately double the free-radical fighting support.

Life Extension® offers the choice of two super-absorbing curcumin formulas that require only one serving a day dosing:

- Super Bio-Curcumin[®] provides optimal potency of highly absorbable curcumin.
- **Advanced Bio-Curcumin® with Ginger & Turmerones** provides the following additional nutrients for those seeking more comprehensive support for prolonged functional inflammatory responses:
 - Ginger to complement health benefits,7
 - Turmerones to increase the amount of curcumin inside cells, and
 - Phospholipids to further enhance absorption.8



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

Non-GMO

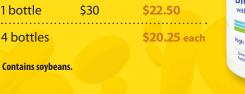


1. Br J Nutr. 2010 Jun; 103(11):1545-57. 2. Nat Sci Biol Med. Jan-Jun;4(1):3-7. 3. Biofactors. 2013 Jan-Feb;39(1):2-13.

Advanced Bio-Curcumin® with Ginger & Turmerones

Item #01808 • 30 softgels

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





- 4. Indian J Pharm Sci. 2008 Jul-Aug; 70(4):445-9.
- 5. Int J Pharmocol. 2009;5(6):333-45. 6. Food Nutr Res. 2009;48(3):148-52.
- 7. J Med Food. 2012 Mar;15(3):242-52. 8. Cancer Chemother Pharmacol. 2007:60:171-7.

trademarks of Dolcas-Biotech, LLC. U.S. Patent Nos. 7,883,728, 7,736,679 and 7,879,373.

Bio-Curcumin® and BCM-95® are registered

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

Dual-Action Formula Supports Healthy Cholesterol Levels

As you age, it becomes critically important to maintain an optimum balance between "good" and "bad" cholesterol, otherwise known as **HDL** and **LDL**, in order to maintain a healthy cholesterol profile.

For those who seek a safe and natural way to support healthy cholesterol levels, researchers have discovered *two compounds* that help promote healthy levels of both LDL and HDL already within normal range.¹⁻⁴

CHOL-Support™ is formulated with a triple standardized **artichoke leaf extract** and **pantethine**, the biologically active form of pantothenic acid³ to maintain healthy HDL, LDL, and total cholesterol levels already within the normal range.¹-⁴

Artichoke Leaf Extract

Clinical studies now demonstrate that artichoke leaf extract's polyphenolic compounds² support optimal HDL/LDL ratios already within normal range. ¹²

Pantethine

Clinical trials show pantethine safely maintains LDL and total cholesterol already within the normal range, without reducing protective CoQ10 levels.³

Two daily liquid vegetarian capsules of **CHOL-Support**™ offer comprehensive cardiovascular protection by providing unparalleled support for an optimal cholesterol profile.

Non-GMO

Pantesin° is a registered trademark of Daiichi Fine Chemical Co., Ltd. Pycrinil° is a registered trademark of Indena S.p.A., Italy.

References

- 1. Arzneimittelforschung. 2000 Mar;50(3):260-5.
- 2. Int J Food Sci Nutr. 2013 Feb;64(1):7-15.
- 3. Vasc Health Risk Manag. 2014;10:89-100.
- 4. Nutr Res. 2011 Aug;31(8):608-15.



Item #01910 • 60 liquid vegetarian capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$52 | \$39 |
| 4 bottles | | \$34.50 each |



To order CHOL-Support[™], call 1-800-544-4440 or visit www.LifeExtension.com



CONFERENCE REPORT

The 2015 American Society For Nutrition Conference

A good diet is essential for anyone wishing to achieve healthy longevity.

With conflicting stories appearing in the mass media, ascertaining what the best food choices are can be a daunting task.

There is disagreement among nutrition scientists, and there are often new studies or opinions that substantially alter dietary recommendations.

For example, saturated fats have been claimed to cause cardiovascular disease, 1 but this claim is debatable among nutrition scientists. 2-5 Interestingly, on a low-carbohydrate diet, high dietary saturated fats do not increase saturated fats in the bloodstream. 6

Discussion of the effects of dietary saturated fat rarely acknowledge the fact that there are about a dozen different saturated fats, each of which has unique properties. Ralmitate is the most abundant saturated fat found in foods. Palmitate constitutes nearly a third of butterfat and beef fat. Palmitate is more likely to cause inflammation than the other saturated fats (particularly in obese people). But palmitate is not inflammatory if consumed with olive oil or fish oil. A saturated fat called myristate strongly raises blood cholesterol, he but the saturated fat stearate does not. The saturated fat butyrate improves insulin sensitivity.

The human body derives energy either from glucose (from carbohydrate) or from fat (or from ketones from fat). ¹⁹ Carbohydrates only became a primary source of energy with the advent of agriculture. As a source of energy, glucose is much more damaging to blood vessels than fat (**saturated fat usually can't be oxidized**). Glucose is the preferred energy source for cancer cells. ²⁰ With the exception of dietary fiber, the fewer carbohydrates in the diet the better, unless there are compensating benefits, such as eating them with blueberries.

Protein is much more complicated. Protein contains essential amino acids, which are important for growth, maintenance, and metabolism of the human body. These functions are especially important for growing children, but are also important for the elderly and to prevent muscle loss in those who exercise. Adequate protein is also important for good immune system function.²¹

Population studies indicate that consumption of red meat and processed meat increases cancer and cardiovascular disease.²² Subjects randomly assigned to a vegetarian diet or an omnivorous diet had lower blood pressure when on the vegetarian diet.²³ Moderately low protein intake seems advisable for most adults (plant protein preferred), although the amount is hard to specify and is dependent on circumstances.

In seeking a healthy diet, I frequently make substantial changes to my eating patterns. In my search for greater clarity, I attended the *American Society for Nutrition* conference in Boston, March 28 to April 1, 2015.



My reports of scientific conferences normally only describe the presentations made by the scientists. But because everyone eats, because diet and dieting have such substantial influence on health, and because the discoveries of nutrition scientists so often seem contradictory or confusing, for this report I am following each presentation description with my interpretations and recommendations. These comments will be my opinions, which are not necessarily those of the Life Extension® Foundation.

My dietary opinions have been substantially influenced by scientists advocating a ketogenic diet, that is, a diet low in carbohydrates, and moderately low in protein. I have been on a vegetarian ketogenic diet for over a year. Insofar as food can only consist of carbohydrate, protein, and fat, my diet is high in fat and fiber. When dietary carbohydrate is low, the liver converts fat into ketones to provide energy for the brain. Although fiber is classified as a carbohydrate, when digested, fiber is converted to beneficial fats in the large intestine.

What do I eat? My "main course" is a vegetable salad (broccoli, green leafy vegetables, shredded cabbage) laden with healthy oil (like olive oil), to which I add walnuts, shredded coconut, and a scoop of whey protein. Among other things I eat are cream cheese with cauliflower, as well as avocado mixed with healthful flavorings such as cocoa powder and wheat grass powder.

Although this is a report on a scientific conference, because diet is such a personal matter, and because I am making recommendations, I need to disclose the basis for my opinions. I believe there is good science behind these opinions, and for that reason I am beginning by reporting the views of Dr. Eric Westman, one of the leading scientific advocates of a ketogenic diet.

Health Benefits Of A Ketogenic Diet

Eric Westman, MD, (Duke University Outpatient Clinic, Durham, North Carolina) is one of the pioneers in challenging the benefits of low-fat diets. He has conducted a number of studies showing that low-carbohydrate, high-fat diets (**ketogenic diets**) result in less hunger, better compliance, and greater **weight loss** than low-fat diets.²⁴⁻²⁶

Dr. Westman conducted a clinical trial in which type II diabetics were randomized to low-calorie diets or to ketogenic diets that did not restrict calories. The diabetics on the ketogenic diet showed greater weight loss, greater reduction in glycated hemoglobin (HbA1C), and greater reduction or elimination of required diabetic medications (95.2 versus 62%).²⁷ Subsequent studies by others have confirmed the benefit of a ketogenic



diet for treatment of type II diabetes. 28,29 Dr. Westman is opening a clinic entirely devoted to weight loss and diabetes treatment using a ketogenic diet (For more information, go to www.healclinics.com).

When carbohydrates are low, the liver converts fats into ketones, which can serve as a fuel source for many tissues, including the brain. 30 Dietary ketosis should not be confused with the ketoacidosis of diabetes, which involves acidity as well as much higher blood ketone concentrations.31,32 Unlike sugar, which stimulates appetite,33 ketones reduce appetite.34,35 As a source of energy for the brain, ketones make mitochondria more efficient, reduce free radical production, and protect against a variety of brain diseases, including epilepsy, stroke, and Alzheimer's disease. 36,37

A ketogenic diet has been shown to increase cognitive performance in aged rats.³⁸ Cancer cells thrive on the glucose resulting from dietary carbohydrates, but are generally unable to use ketones as an energy source.³⁹ Ketones have been shown to suppress tumors and prolong survival in mice that have metastatic cancer.40,41

Saturated fats have been claimed to cause cardiovascular disease,1 but this claim is debatable among nutrition scientists.²⁻⁵ A lot probably depends on the type of saturated fat and what other foods it is consumed with.

Insulin resistance results when excess fat accumulates in muscle, liver, and pancreas rather than in fat cells.42 But on a carbohydrate-restricted diet, fats are used for energy rather than stored. On a low-carbohydrate diet, high dietary saturated fats do not increase saturated fats in the bloodstream. 6 Cardiovascular risk factors may be reduced on a high-fat, low-carbohydrate (ketogenic) diet,43,44 and a low-carbohydrate diet has been shown to increase insulin sensitivity. 45,46

My interpretation/recommendation: A diet low in carbohydrates, moderately low in protein, and high in proper fats is recommended for persons trying to lose weight as well as for prevention and treatment of diabetes (and other age-related afflictions, notably cancer and cardiovascular disease).

Dietary Sugar

Deborah Sloboda, PhD, (Associate Professor, Biochemistry and Biomedical Sciences, McMaster University, Hamilton, Ontario, Canada) does research on the damaging effects of dietary sugar, particularly fructose.

Simple sugar constituted only about 2% of the prehistoric Paleolithic diet, but currently accounts for about 18 to 25% of total calories of the average modern Western diet. 47 Modern man is thus consuming about **10 times** more simple sugar than our early ancestors.

High fructose corn syrup, introduced in 1967, accounted for 42% of caloric sweeteners being used by the year 2000.⁴⁸ Fructose consumption increases visceral (abdominal) fat while reducing insulin sensitivity.⁴⁹ Dr. Sloboda is particularly concerned with fructose consumption during pregnancy, which can lead to gestational diabetes (temporary diabetes due to pregnancy), causing inflammatory metabolic disturbance for both the mother and the fetus. 50,51 The amino acid taurine can partially reduce the damaging effects of fructose.51,52

My interpretation/recommendation: Fructose consumption should be limited because fructose can elevate blood triglycerides while causing insulin resistance and the diseases of inflammation and diabetes. The soft drinks accounting for about 90% of the soda market (Coca-Cola®, Pepsi®, Dr. Pepper®, Mountain Dew[®], and Sprite[®]) all contain **60** to **65**% fructose.⁵³ Apple juice has more than twice the fructose as orange juice, yet apple juice is the most common sweetener in fruit drinks. Dried apples, dates, apricots, figs, and raisins are one-quarter to one-third fructose by weight, and should be eaten sparingly. Fructose can cause harm to both the mother and fetus during pregnancy and pregnant women should limit consumption of fructosecontaining beverages, and perhaps even supplement with taurine.

Dietary Starch

Diane Birt, PhD, (Distinguished Professor Emeritus, Iowa State University) is interested in using digestion-



resistant starch to improve human health. Starch, like sugar, is a form of carbohydrate, but starches that resist digestion in the stomach and small intestine, yet are digested in the large intestine, are called resistant starch or fiber.

Dr. Birt has described most foods in the typical Western diet as being highly digestible starches having a

high glycemic index (which raise blood glucose).⁵⁴ Such

foods result in cycles of high and low blood glucose and blood insulin, often causing insulin resistance and type II diabetes.⁵⁴ All starchy foods are composed of chains of sugar molecules, but not all starchy foods are as easily digested. Digestion-resistant starch eaten for breakfast reduces the surges of glucose and insulin at lunch.⁵⁵

Legumes (beans) contain more resistant starch than potatoes, rice, pasta, bread, or breakfast cereals. But Dr. Birt wishes to develop foods with higher resistant starch content than legumes.⁵⁴ Resistant starches have been shown to reduce obesity.⁵⁶ Starches (dietary fibers) that are more viscous reduce appetite more than those that are less viscous,⁵⁷ so the starches she is developing would be more viscous. Starches made of long unbranched chains of sugars are more *digestion-resistant* than starches having branched chains, a fact Dr. Birt also exploits when designing improved foods for health.⁵⁴

My interpretation/recommendation: Rapidly digested carbohydrate starches such as potatoes, pastry, white bread, and breakfast cereals should be avoided, but beans and waxy corn starch are good sources of dietary fiber that can satisfy appetite while reducing insulin surges after eating.^{54,58}

Dietary Fat

David Mutch, PhD, (Associate Professor, University of Guelph, Canada) is interested in how fats and inflammatory substances in the bloodstream relate to obesity.



Although obesity is often associated with insulin resistance and metabolic syndrome, he estimates that **13** to **29**% of obese persons (BMI greater than 30) are metabolically healthy, whereas **10** to **37**% of lean persons (BMI less than 25) are metabolically unhealthy, having insulin resistance.⁵⁹

Metabolically healthy obese people are less likely to have fat in heart cells

or in the insulin-producing cells of the pancreas.⁵⁹ Metabolically unhealthy people have higher blood levels of the saturated fat **palmitate**.^{60,61} Foods high in palminate include cocoa butter oil, palm oil, shortening, butter, lard, milk chocolate, fatty meats, pork and beef products, game meats, cashews, and eggs.⁶¹

Palmitate can induce inflammation and insulin resistance. 62-64 Elevated blood palmitate and glucose cause insulin-producing cells in the pancreas to die, but the **monounsaturated fat** oleic acid (high in **olive oil** and **avocado**) protects cells against glucose and palmitate toxicity. 13,14 *Omega-3* fatty acid supplementation of a mouse diet high in saturated fat also prevents inflammation from palmitate. 15



My interpretation/recommendation: Olive oil, fish oil, and avocado can protect against inflammation due to the saturated fat palmitate.

Kevin Fritsche, PhD, (Professor of Nutritional Immunology, University of Missouri) studies the effects of dietary fats on inflammation and the risk of car-



diovascular disease. He believes that the inflammatory blood component *C-reactive protein* is more strongly associated with cardiovascular disease than LDL cholesterol. ⁶⁵ Release of the inflammatory protein IL-18 from fat cells of obese people is triple of that seen in those who are not obese. ⁶⁶ Although he affirms that *trans-fatty acids* promote inflammation and

reduce function of the endothelial cells lining blood vessels,⁶⁷ he finds not much support for the claim that omega-6 fatty acids are pro-inflammatory.^{68,69}

The gut microbe *Bifidobacteria* has been shown to prevent the highly pro-inflammatory endotoxin lipopolysaccharide (or LPS, which constitutes **80**% of the cell wall of gram-negative bacteria) from leaking into the bloodstream from the intestine. ⁷⁰⁻⁷⁴ A high-fat, high-carbohydrate diet promotes inflammation by reducing the quantity of *Bifidobacteria*, ⁷³⁻⁷⁵ although some foods (including walnuts, olive oil, and wheat bran) can lessen this effect. ⁷⁶

The magnitude of inflammation resulting from a high-fat, high-carbohydrate meal is markedly greater in obese than in non-obese subjects. ^{77,78} Inflammation from a high-fat, high-carbohydrate meal may be independent of LPS concentration in the bloodstream. ⁷⁹

My interpretation/recommendation: LDL cholesterol is less of a cardiovascular disease concern than inflammation and insulin resistance. A high-fat diet should be avoided only if it is combined with a diet high in carbohydrates. Dietary carbohydrates cause insulin secretion, and insulin prevents fat from being used as an energy source, resulting in fat being stored in places other than fat cells, namely in muscle and pancreas, resulting in insulin resistance. Because this effect is largest in obesity, obese people should make a particular effort to reduce carbohydrates.

Weight-Loss Strategies

Catherine Hankey, PhD (Senior Lecturer in Human Nutrition, University of Glasgow, United Kingdom)



reviewed studies on intermittent fasting as a weight-loss strategy. At any given time, about 40% of women and 20% of men will be fasting for weight loss.80

Dr. Hankey began by acknowledging the bestselling book *The Fast* Diet by Dr. Michael Mosley and Mimi Spencer, which advocates reducing calories to 25% the usual daily intake

on two days each week (Monday and Thursday being the best choice for most people).

A high-protein diet can assist fasting because protein has been shown to reduce appetite. 81 High-protein diets (25% or more of calories) are associated with the greatest weight loss (and not much hunger) for a few months. By two years, however, there is little lasting weight loss.⁸¹ A three-month comparison of subjects restricting calories 25% either on a daily basis with high protein, or only on two days per week with carbohydrate restriction, showed greater body fat reduction along with improved insulin sensitivity for the intermittent-fasting, carbohydrate-restricting group.82

My interpretation/recommendation: Complete fasting for more than a day is intolerable for most people, but Dr. Mosley's book became a bestseller because so many people have succeeded in losing weight by eating only 25% of their usual calories twice weekly. I recommend this book to readers who have not succeeded with other weight-loss methods. The book contains recipes and strategies. Despite initial weight loss and reduction of hunger on a high-protein diet, high-protein diets are not recommended. Weight loss with a high-protein diet is only a temporary effect. Moreover, high dietary protein can result in chronic inflammation with subsequent increase in cancer and cardiovascular disease.83,84

Cynthia Kroeger and John Trepanowski, (PhD students, Department of Kinesiology and Nutrition, University of Illinois, Chicago) have conducted a number of studies together on the effects of alternate-day fasting (with 25% of usual calories consumed on the fast day).85-87 In one such study, the weight loss from endurance exercise and fasting in combination was compared with exercise and fasting alone. The combination resulted in a 13.2 pound weight loss, fasting alone resulted in a 6.6 pound weight loss, and exercise alone resulted in a 2.2 pound weight loss.88

Exercise alone is usually not very effective for weight loss,89,90 at least partially because of increased appetite. The most important role of exercise in dieting is to prevent loss of lean mass (and to boost AMPK enzyme activity).91 In another study, Kroeger and Trepanowski found that adding liquid meal replacement to intermittent fasting and calorie restriction resulted in an even greater reduction in weight, visceral fat, and LDL cholesterol. The liquid meals ensured greater portion control. Only the subjects receiving the liquid meal replacement showed a reduction in blood glucose, insulin, and homocysteine, as well as reduced LDL particle size.86

My interpretation/recommendation: Although exercise alone is not very effective for weight loss, exercise doubles the weight loss associated with a reduced-calorie diet, while preventing loss of muscle tissue. Combining exercise with calorie restriction is highly recommended for dieters.

Diana Thomas, PhD, (Professor of Mathematical Sciences, Montclair State University, Montclair, New Jersey) has created mathematical models to explain weight-loss effects associated with dieting. 92,93 Weight lost in the initial phases of dieting is typically the result of greater water loss associated with protein and carbohydrate loss.94 Unlike fat, proteins (like muscle) and carbohydrates (like glycogen) are stored with much water in the body, so loss of lean mass or carbohydrate results in loss of considerable amounts of water. 95,96 Calories required for weight loss increases with the duration of dieting because as dieting proceeds, weight loss increasingly means fat loss. According to Dr. Thomas, after four weeks of dieting, a reduction of about 2,000 dietary calories results in a loss of one pound. But by 24 weeks of dieting, a reduction of about 3,000 dietary calories is required to lose one pound.94

For an average person, resting metabolic rate accounts for 70% of calories consumed, physical activity accounts for 20%, and energy required to digest and metabolize food (the thermic effect of food) accounts for 10%.97

The thermic effect of protein is more than double the thermic effect of fat or carbohydrate. 98 The thermic effect of glucose is reduced with insulin resistance because insulin facilitates glucose storage (which consumes more energy than glucose used as an energy source).99

Resting metabolic rate typically declines 2 to 3% every decade after age 20, mostly due to loss of lean, fat-free mass.97 Physical activity also tends to decline with age. These factors make weight loss increasingly difficult with age. On average, body weight is highest for those in the 50-59 age group, but much of the reason for this may be due to higher death rates among the obese.⁹⁷

My interpretation/recommendation: Weight loss becomes more difficult with age largely because of the muscle loss associated with reduced activity. The more muscle (lean body mass) that a person has, the higher their resting metabolic rate will be. Resistance exercise is particularly important for building and maintaining muscle, as well as preventing the frailty too often associated with aging. Exercise should become a lifetime practice, not simply something done while dieting.

Muscle Loss In The Elderly

Ronenn Roubenoff, MD, (Internist, Tufts Medical Center, Boston) studies muscle loss in the elderly. He cited a three-year study of over 3,000 people in their



70s, which showed that loss of muscle quality was even greater than loss of muscle mass. 100 Deterioration of muscle in the elderly not only leads to functional impairment and falls, but to chronic heart failure and chronic obstructive pulmonary disease. 101-103 Endurance as well as resistance exercise, along with increased protein intake is recommended to prevent

muscular atrophy in the elderly, ¹⁰⁴ although those with kidney problems due to diabetes may need to limit their protein intake. ¹⁰⁵

My interpretation/recommendation: As stated above, endurance exercise, resistance exercise, and adequate protein can prevent muscle wasting and can reduce frailty in the elderly (for whom muscle-wasting can be life threatening).

Circadian Rhythm And Diet

Frank Scheer, PhD, (Assistant Professor of Sleep Medicine, Harvard Medical School) is an expert on the effects of circadian (day-night cycle) rhythm on dietary metabolism. The brain has a clock that normally causes **melatonin** secretion to increase in the evening (facilitating sleep), which causes secretion of growth hormone and prolactin at sleep onset, and which causes secretion of the stress hormone **cortisol** before awakening. The risk of a heart attack in the morning is elevated **40**% above the 24-hour average, the morning is elevated 40% above the 24-hour average, the stress partially because cortisol raises blood pressure. The Scheer's research has established that disruption of the circadian rhythm by jet lag or shift work leads to insulin resistance. The Melatonin supplementation can lessen these circadian stresses.

Dr. Scheer has also established that for those with a normal circadian rhythm, hunger is lowest at 8 a.m. and highest at 8 p.m. ¹¹⁰ There is also a circadian effect on appetite for certain foods. The 8 p.m. appetite for sweets, starches, and meat is increased, but there is no circadian effect for vegetables or dairy products. ¹¹⁰

Dr. Scheer conducted two studies showing that breakfast skippers and late eaters gain more weight (and have more trouble losing weight) than early eaters, despite consuming the same total number of calories. ^{111,112} But a randomized study focused only on those who skip breakfast showed no effect on weight gain or loss. ¹¹³ A review by other authors, while acknowledging that late-night eating contributes to metabolic syndrome, denied a similar effect for breakfast skipping, suggesting that breakfast skipping might be more successful than other strategies for calorie restriction. ¹¹⁴

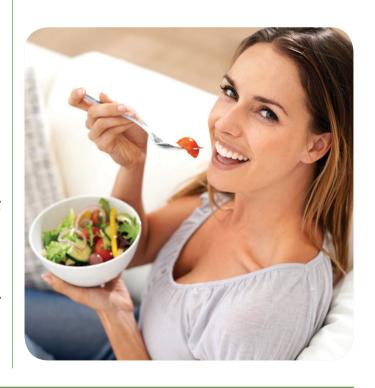
My interpretation/recommendation: Eating in the evening before bedtime can lead to greater weight gain than calorie counting would predict. Skipping breakfast is probably not a health hazard. Melatonin should be used to reduce the effects of jet lag or shift work.

Conclusions

There may be a revolution happening in the perception of the effects of dietary fat, especially in the context of a ketogenic diet. It would be hard to prove that the US Department of Agriculture dietary recommendations of high grain (carbohydrate, the "food pyramid") and low fat^{115,116} contributed significantly to the current obesity epidemic, but if true, the government will not readily admit blame.¹¹⁷

If you have any questions on the scientific content of this article, please call a Life Extension®

Health Advisor at 1-866-864-3027.



References

- Kris-Etherton PM, Innis S, American Dietetic Association, Dietitians of Canada. Position of the American Dietetic Association and Dietitians of Canada: dietary fatty acids. J Am Diet Assoc. 2007 Sep:107(9):1599-611.
- Siri-Tarino PW, Sun Q, Hu FB, Krauss RM. Meta-analysis of prospective cohort studies evaluating the association of saturated fat with cardiovascular disease. Am J Clin Nutr. 2010 Mar;91(3): 535-46
- Scarborough P, Rayner M, van Dis I, Norum K. Meta-analysis of effect of saturated fat intake on cardiovascular disease: over adjustment obscures true associations. Am J Clin Nutr. 2010 Aug;92(2):458-9; author reply 459.
- Chowdhury R, Warnakula S, Kunutsor S, et al. Association of dietary, circulating, and supplement fatty acids with coronary risk: a systematic review and meta-analysis. Ann Intern Med. 2014 Mar 18:160(6):398-406.
- Hooper L, Martin N, Abdelhamid A, Davey Smith G. Reduction in saturated fat intake for cardiovascular disease. Cochrane Database Syst Rev. 2015 Jun10;6:CD011737.
- Forsythe CE, Phinney SD, Feinman RD, et al. Limited effect of dietary saturated fat on plasma saturated fat in the context of a low carbohydrate diet. Lipids. 2010 Oct;45(10):947-62.
- Mensink RP, Zock PL, Kester AD, Katan MB. Effects of dietary fatty acids and carbohydrates on the ratio of serum total to HDL cholesterol and on serum lipids and apolipoproteins: a meta-analysis of 60 controlled trials. Am J Clin Nutr. 2003 May;77(5):
- Siri-Tarino PW, Sun Q, Hu FB, Krauss RM. Saturated fatty acids and risk of coronary heart disease: modulation by replacement nutrients. Curr Atheroscler Rep. 2010 Nov;12(6):384-90.
- Vock C, Gleissner M, Klapper M, Doring F. Identification of palmitate-regulated genes in HepG2 cells by applying microarray analysis. Biochim Biophys Acta. 2007 Sep;1770(9):1283-88.
- 10. French MA, Sundram K, Clandinin MT. Cholesterolaemic effect of palmitic acid in relation to other dietary fatty acids. Asia Pac J Clin Nutr. 2002;11Suppl7:S401-7
- 11. 11. Tian H, Liu C, Zou X, Wu W, Zhang C, Yuan D. MiRNA-194 regulates palmitic acid-induced toll-like receptor 4 inflammatory responses in THP-1 cells. Nutrients. 2015 May 13;7(5):3483-96.
- 12. Laugerette F, Furet JP, Debard C, et al. Oil composition of high-fat diet affects metabolic inflammation differently in connection with endotoxin receptors in mice. Am J Physiol Endocrinol Metab. 2012 Feb 1;302(3):E374-86.
- 13. Listenberger LL, Han X, Lewis SE, Cases et al. Triglyceride accumulation protects against fatty acid-induced lipotoxicity. Proc Natl Acad Sci U S A. 2003 Mar 18;100(6):3077-82.
- 14. Maedler K, Oberholzer J, Bucher P, Spinas GA, Donath MY. Monounsaturated fatty acids prevent the deleterious effects of palmitate and high glucose on human pancreatic beta-cell turnover and function. Diabetes. 2003 Mar;52(3):726-33.
- 15. Awada M, Meynier A, Soulage CO, et al. n-3 PUFA added to high-fat diets affect differently adiposity and inflammation when carried by phospholipids or triacylglycerols in mice. Nutr Metab (Lond). 2013 Feb 15;10(1):23.
- 16. Temme EH, Mensink RP, Hornstra G. Effects of medium chain fatty acids(MCFA), myristic acid, and oleic acid on serum lipoproteins in healthy subjects. J Lipid Res. 1997 Sep;38(9):1746-54.
- 17. Hunter JE, Zhang J, Kris-Etherton PM. Cardiovascular disease risk of dietary stearic acid compared with trans, other saturated, and unsaturated fatty acids: a systematic review. Am J Clin Nutr. 2010 Jan;91(1):46-63.
- 18. Gao Z, Yin J, Zhang J, Ward RE, Martin RJ, Lefevre M, Cefalu WT, Ye J. Butyrate improves insulin sensitivity and increases energy expenditure in mice. Diabetes. 2009 Jul;58(7):1509-17.
- Askew EW. Role of fat metabolism in exercise. Clin Sports Med. 1984 Jul;3(3):605-21.
- 20. Bozzetti F, Gavazzi C, Mariani L, Crippa F. Glucose-based total parenteral nutrition does not stimulate glucose uptake by humans tumours. Clin Nutr (Edinburgh). 2004 Jun; 23(3):417-21.

- 21. Rusu D, Drouin R, Pouliot Y, Gauthier S, Poubelle PE. A bovine whey protein extract can enhance innate immunity by priming normal human blood neutrophils. J Nutr. 2009 Feb;139(2):386-93.
- 22. Sinha R, Cross AJ, Graubard BI, Leitzmann MF, Schatzkin A. Meat intake and mortality: a prospective study of over half a million people. Arch Intern Med. 2009 Mar 23;169(6):562-71.
- 23. Rouse IL, Beilin LJ, Armstrong BK, Vandongen R. Blood-pressurelowering effect of a vegetarian diet: controlled trial in normotensive subjects. Lancet. 1983 Jan 1;1(8314-5):5-10.
- 24. Yancy WS Jr, Olsen MK, Guyton JR, Bakst RP, Westman EC. A low-carbohydrate, ketogenic diet versus a low-fat diet to treat obesity and hyperlipidemia: a randomized, controlled trial. Ann Intern Med. 2004 May 18;140(10):769-77.
- 25. Lecheminant JD, Gibson CA, Sullivan DK, et al. Comparison of a low carbohydrate and low fat diet for weight maintenance in overweight or obese adults enrolled in a clinical weight management program. Nutr J. 2007 Nov 1;6:36.
- 26. McClernon FJ, Yancy WS Jr, Eberstein JA, Atkins RC, Westman EC. The effects of a low-carbohydrate ketogenic diet and a low-fat diet on mood, hunger, and other self-reported symptoms. Obesity (Silver Spring). 2007 Jan;15(1):182-7.
- 27. Westman EC, Yancy WS Jr, Mavropoulos JC, Marquart M, McDuffie JR. The effect of a low-carbohydrate, ketogenic diet versus a low-glycemic index diet on glycemic control in type 2 diabetes mellitus. Nutr Metab (Lond). 2008 Dec19:5:36.
- 28. Hussain TA, Mathew TC, Dashti AA, Asfar S, Al-Zaid N, Dashti HM. Effect of low-calorie versus low-carbohydrate ketogenic diet in type 2 diabetes. Nutrition. 2012 Oct;28(10):1016-21.
- 29. Mobbs CV, Mastaitis J, Isoda F, Poplawski M. Treatment of diabetes and diabetic complications with a ketogenic diet. J Child Neurol. 2013 Aug;28(8):1009-14.
- 30. Greene AE, Todorova MT, Seyfried TN. Perspectives on the metabolic management of epilepsy through dietary reduction of glucose and elevation of ketone bodies. J Neurochem. 2003 Aug;86(3):529-37.
- 31. Manninen AH. Metabolic effects of the very-low-carbohydrate diets: misunderstood "villains" of human metabolism. J Int Soc Sports Nutr. 2004 Dec 31;1(2):7-11.
- 32. Paoli A, Bosco G, Camporesi EM, Mangar D. Ketosis, ketogenic diet and foodintake control: a complex relationship. Front Psychol. 2015 Feb 2;6:27.
- 33. Gaysinskaya VA, Karatayev O, Shuluk J, Leibowitz SF. Hyperphagia induced by sucrose: relation to circulating and CSF glucose and corticosterone and or exigenic peptides in the arcuate nucleus. Pharmacol Biochem Behav. 2011 Jan;97(3):521-30.
- 34. Gibson AA, Seimon RV, Lee CM, Ayre J, Franklin J, Markovic TP, Sainsbury A. Do ketogenic diets really suppress appetite? A systematic review and meta-analysis. Obes Rev. 2015 Jan;16(1):64-76.
- Veldhorst MA, Westerterp KR, van Vught AJ, Westerterp-Plantenga MS. Presence or absence of carbohydrates and the proportion of fat in a high-protein diet affect appetite suppression but not energy expenditure in normal-weight human subjects fed in energy balance. Br J Nutr. 2010 Nov;104(9):1395-405.
- 36. Maalouf M, Rho JM, Mattson MP. The neuro protective properties of calorie restriction, the ketogenic diet, and ketone bodies. Brain Res Rev. 2009 Mar;59(2):293-315.
- 37. Stafstrom CE, Rho JM. The ketogenic diet as a treatment paradigm for diverse neurological disorders. Front Pharmacol. 2012 Apr 9:3:59.
- 38. Xu K, Sun X, Eroku BO, Tsipis CP, Puchowicz MA, LaManna JC. Diet-induced ketosis improves cognitive performance in aged rats. Adv Exp Med Biol. 2010;662:71-5.
- 39. Seyfried TN, Flores RE, Poff AM, D'Agostino DP. Cancer as a metabolic disease: implications for novel therapeutics. Carcinogenesis. 2014 Mar;35(3):515-27.
- 40. Siri-Tarino PW, Sun Q, Hu FB, Krauss RM. Saturated fat, carbohydrate, and cardiovascular disease. Am J Clin Nutr. 2010 Mar:91(3):502-9.
- 41. Poff AM, Ari C, Arnold P, Seyfried TN, D'Agostino DP. Ketone supplementation decreases tumor cell viability and prolongs survival of mice with metastatic cancer. Int J Cancer. 2014 Oct 1;135(7):1711-20.

- 42. Borén J, Taskinen MR, Olofsson SO, Levin M. Ectopic lipid storage and insulin resistance: a harmful relationship. *J Intern Med*. 2013 Jul;274(1):25-40.
- 43. Sharman MJ, Kraemer WJ, Love DM, Avery NG, Gómez AL, Scheett TP, Volek JS. A ketogenic diet favorably affects serum biomarkers for cardiovascular disease in normal-weight men. *J Nutr*. 2002 Jul;132(7):1879-85
- 44. Wood RJ, Volek JS, Davis SR, Dell'Ova C, Fernandez ML. Effects of a carbohydrate-restricted diet on emerging plasma markers for cardio vascular disease. *Nutr Metab (Lond)*. 2006 May 4;3:19.
- 45. Gower BA, Goss AM. A lower-carbohydrate, higher-fat diet reduces abdominal and intermuscular fat and increases insulin sensitivity in adults at risk of type 2 diabetes. *J Nutr.* 2015 Jan;145(1): 177s-183s.
- Kirk E, Reeds DN, Finck BN, Mayurranjan SM, Patterson BW, Klein S. Dietary fat and carbohydrates differentially alter insulin sensitivity during caloric restriction. *Gastroenterology*. 2009 May;136(5):1552-60.
- Gray C, Long S, Green C, Gardiner SM, Craigon J, Gardner DS. Maternal fructose and/or salt intake and reproductive outcome in the rat: effects on growth, fertility, sex ratio, and birth order. *Biol Reprod*. 2013 Sep;89(3):51.
- 48. Bray GA, Nielsen SJ, Popkin BM. Consumption of high-fructose corn syrup in beverages may play a role in the epidemic of obesity. *Am J Clin Nutr*. 2004 Apr;79(4):537-43.
- Stanhope KL, Havel PJ. Fructose consumption: recent results and their potential implications. *Ann N Y Acad Sci.* 2010 Mar;1190: 15-24.
- Regnault TR, Gentili S, Sarr O, Toop CR, Sloboda DM. Fructose, pregnancy and later life impacts. *Clin Exp Pharmacol Physiol*. 2013 Nov;40(11):824-37.
- 51. Li M, Reynolds CM, Sloboda DM, Gray C, Vickers MH. Maternal taurine supplementation attenuates maternal fructose-induced metabolic and inflammatory dysregulation and partially reverses adverse metabolic programming in offspring. *J Nutr Biochem*. 2015 Mar;26(3):267-76.
- 52. Sloboda DM, Li M, Patel R, Clayton ZE, Yap C, Vickers MH. Early life exposure to fructose and offspring phenotype: implications for long term metabolic homeostasis. *J Obes*. 2014;2014:203474.
- Walker RW, Dumke KA, Goran MI. Fructose content in popular beverages made with and without high-fructose corn syrup. *Nutri*tion. 2014 Jul-Aug;30(7-8):928-35.
- 54. Birt DF, Boylston T, Hendrich S, et al. Resistant starch: promise for improving human health. *Adv Nutr.* 2013 Nov 6;4(6):587-601.
- 55. Brighenti F, Benini L, Del Rio D, et al. Colonic fermentation of indigestible carbohydrates contributes to the second-meal effect. *Am J Clin Nutr*. 2006 Apr;83(4):817-22.
- Belobrajdic DP, King RA, Christophersen CT, Bird AR. Dietary resistant starch dose-dependently reduces adiposity in obesityprone and obesity-resistant male rats. *Nutr Metab (Lond)*. 2012 Oct 25:9(1):93.
- 57. Wanders AJ, van den Borne JJ, de Graaf C, Hulshof T, Jonathan MC, Kristensen M, Mars M, Schols HA, Feskens EJ. Effects of dietary fibre on subjective appetite, energy intake and body weight: a systematic review of randomized controlled trials. *Obes Rev.* 2011 Sep;12(9):724-39.
- 58. Behall KM, Scholfield DJ, Yuhaniak I, Canary J. Diets containing high amylose vs amylopectin starch: effects on metabolic variables in human subjects. *Am J Clin Nutr*. 1989 Feb;49(2):337-44.
- 59. Badoud F, Perreault M, Zulyniak MA, Mutch DM. Molecular insights into the role of white adipose tissue in metabolically unhealthy normal weight and metabolically healthy obese individuals. *FASEB J.* 2015 Mar;29(3):748-58.
- Perreault M, Zulyniak MA, Badoud F, Stephenson S, Badawi A, Buchholz A, Mutch DM. A distinct fatty acid profile underlies the reduced inflammatory state of metabolically healthy obese individuals. *PLoS One*. 2014 Feb 10;9(2):e88539.
- 61. http://www.lifeextension.com/pdf/BloodTestSamples/Omega_ Check_Interpretation_Guide.pdf. Accessed September 29, 2015.

- 62. Nakamura S, Takamura T, Matsuzawa-Nagata N, et al. Palmitate induces insulin resistance in H4IIEC3 hepatocytes through reactive oxygen species produced by mitochondria. *J Biolog Chem.* 2009 May 29;284(22):14809-18.
- 63. Shi H, Kokoeva MV, Inouye K, Tzameli I, Yin H, Flier JS. TLR4 links innate immunity and fatty acid-induced insulin resistance. *J Clin Invest*. 2006 Nov;11;6(11):3015-25.
- Shaw B, Lambert S, Wong MH, Ralston JC, Stryjecki C, Mutch DM. Individual saturated and monounsaturated fatty acids trigger distinct transcriptional networks in differentiated 3T3-L1 preadipocytes. J Nutrigenet Nutrigenomics. 2013;6(1):1-15.
- 65. Emerging Risk Factors Collaboration, Kaptoge S, Di Angelantonio E, et al. C-reactive protein concentration and risk of coronary heart disease, stroke, and mortality: an individual participant meta-analysis. *Lancet*. 2010 Jan 9;375(9709):132-40.
- Packard RR, Libby P. Inflammation in atherosclerosis: from vascular biology to biomarker discovery and risk prediction. *Clin Chem*. 2008 Jan;54(1):24-38.
- 67. Lopez-Garcia E, Schulze MB, Meigs JB, Manson JE, Rifai N, Stampfer MJ, Willett WC, Hu FB. Consumption of trans fatty acids is related to plasma biomarkers of inflammation and endothelial dysfunction. *J Nutr.* 2005 Mar;135(3):562-6.
- 68. Johnson GH, Fritsche K. Effect of dietary linoleic acid on markers of inflammation in healthy persons: a systematic review of randomized controlled trials. *J Acad Nutr Diet*. 2012 Jul;112(7):1029-41, 1041.e1-15.
- Khandelwal S, Kelly L, Malik R, Prabhakaran D, Reddy S. Impact of omega-6fatty acids on cardiovascular outcomes: A review. *J Preventive Cardiol.* 2013 Feb;2(3):325-336.
- Park MS, Kim MJ, Ji GE. Assessment of lipopolysaccharide-binding activity of Bifidobacterium and its relationship with cell surface hydrophobicity, autoaggregation, and inhibition of interleukin-8 production. *J Microbiol Biotechnol*. 2007 Jul;17(7):1120-1126.
- 71. Silhavy TJ, Kahne D, Walker S. The bacterial cell envelope. *Cold Spring Harb Perspect Biol.* 2010 May;2(5):a000414.
- 72. Van Amersfoort ES, Van Berkel TJ, Kuiper J. Receptors, mediators, and mechanisms involved in bacterial sepsis and septic shock. *Clinical microbiology reviews*. 2003 Jul;16(3):379-414.
- Cani PD, Neyrinck AM, Fava F, et al. Selective increases of bifido bacteria in gut microflora improve high-fat-diet-induced diabetes in mice through a mechanism associated with endotoxaemia. *Diabetologia*. 2007 Nov;50(11):2374-83.
- 74. Cani PD, Bibiloni R, Knauf C, et al. Changes in gut microbiota control metabolic endotoxemia-induced inflammation in high-fat diet-induced obesity and diabetes in mice. *Diabetes*. 2008 Jun;57(6):1470-81.
- 75. Kim KA, Gu W, Lee IA, Joh EH, Kim DH. High fat diet-induced gut microbiota exacerbates inflammation and obesity in mice via the TLR4 signaling pathway. *PLoS One*. 2012;7(10):e47713.
- Kelly CJ, Colgan SP, Frank DN. Of microbes and meals: the health consequences of dietary endotoxemia. *Nutr Clin Pract*. 2012 Apr;27(2):215-25.
- 77. Patel C, Ghanim H, Ravishankar S, et al. Prolonged reactive oxygen species generation and nuclear factor-kappa Bactivation after a high-fat, high-carbohydrate meal in the obese. *J Clin Endocrinol Metab*. 2007 Nov;92(11):4476-9.
- 78. Ghanim H, Abuaysheh S, Sia CL, et al. Increase in plasma endotoxin concentrations and the expression of Toll-like receptors and suppressor of cytokine signaling-3 in mononuclear cells after a high-fat, high-carbohydrate meal: implications for insulin resistance. *Diabetes Care*. 2009 Dec;32(12):2281-7.
- Fogarty CL, Nieminen JK, Peräneva L, et al. High-fat meals induce systemic cytokine release without evidence of endotoxemia-mediated cytokine production from circulating monocytes or myeloid dendritic cells. *Acta Diabetol.* 2015 Apr;52(2):315-22.
- 80. Johnstone A. Fasting for weight loss: an effective strategy or latest dieting trend? *Int J Obes (Lond)*. 2015 May;39(5):727-33.
- 81. Weigle DS, Breen PA, Matthys CC, et al. A high-protein diet induces sustained reductions in appetite, ad libitum caloric intake, and body weight despite compensatory changes in diurnal plasma leptin and ghrelin concentrations. *Am J Clin Nutr.* 2005 Jul;82(1):41-8.

- 82. Harvie M, Wright C, Pegington M, et al. The effect of intermittent energy and carbohydrate restriction v. daily energy restriction on weight loss and metabolic disease risk markers in overweight women. Br J Nutr. 2013 Oct;110(8):1534-47.
- 83. Fontana L, Weiss EP, Villareal DT, Klein S, Holloszy JO. Longterm effects of calorie or protein restriction on serum IGF-1 and IGFBP-3 concentration in humans. Aging Cell. 2008 Oct;7(5):681-7.
- 84. Levine ME, Suarez JA, Brandhorst S, et al. Low protein intake is associated with a major reduction in IGF-1, cancer, and overall mortality in the 65 and younger but not older population. Cell Metab. 2014 Mar 4;19(3):407-17
- 85. Hoddy KK, Kroeger CM, Trepanowski JF, Barnosky A, Bhutani S, Varady KA. Meal timing during alternate day fasting: Impact on body weight and cardiovascular disease risk in obese adults. Obesity (Silver Spring). 2014 Dec;22(12):2524-31.
- 86. Klempel MC, Kroeger CM, Bhutani S, Trepanowski JF, Varady KA. Intermittent fasting combined with calorie restriction is effective for weight loss and cardio-protection in obese women. Nutr J. 2012 Nov 21:11:98.
- Kroeger CM, Klempel MC, Bhutani S, Trepanowski JF, Tangney CC, Varady KA. Improvement in coronary heart disease risk factors during an intermittent fasting/calorie restriction regimen: Relationship to adipokine modulations. Nutr Metab (Lond). 2012 Oct 31;9(1):98.
- 88. Bhutani S, Klempel MC, Kroeger CM, Trepanowski JF, Varady KA. Alternate day fasting and endurance exercise combine to reduce body weight and favorably alter plasma lipids in obese humans. Obesity (Silver Spring). 2013 Jul;21(7):1370-9.
- 89. Thomas DM, Bouchard C, Church T, et al. Why do individuals not lose moreweight from an exercise intervention at a defined dose? An energy balance analysis. Obes Rev. 2012 Oct;13(10):835-47.
- 90. Johns DJ, Hartmann-Boyce J, Jebb SA, Aveyard P, Behavioural Weight Management Review Group. Diet or exercise interventions vs combined behavioral weight management programs: a systematic review and meta-analysis of direct comparisons. J Acad Nutr Diet. 2014 Oct;114(10):1557-68.
- 91. Stiegler P, Cunliffe A. The role of diet and exercise for the maintenance of fat-free mass and resting metabolic rate during weight loss. Sports Med. 2006;36(3):239-62.
- 92. Thomas DM, Ciesla A, Levine JA, Stevens JG, Martin CK. A mathematical model of weight change with adaptation. Math Biosci Eng. 2009 Oct;6(4):873-87.
- 93. Heymsfield SB, Gonzalez MC, Shen W, Redman L, Thomas D. Weight loss composition is one-fourth fat-free mass: a critical review and critique of this widely cited rule. Obes Rev. 2014 Apr;15(4):310-21.
- 94. Thomas DM, Gonzalez MC, Pereira AZ, Redman LM, Heymsfield SB. Time to correctly predict the amount of weight loss with dieting. J Acad Nutr Diet. 2014 Jun;114(6):857-61.
- 95. Puolanne E, Halonen M. Theoretical aspects of water-holding in meat. Meat Sci. 2010 Sep;86(1):151-65.
- 96. Kreitzman SN, Coxon AY, Szaz KF. Glycogen storage: illusions of easy weight loss, excessive weight regain, and distortions in estimates of body composition. Am J Clin Nutr. 1992 Jul;56(1 Suppl):292S-293S.
- Villareal DT, Apovian CM, Kushner RF, Klein S; American Society for Nutrition; NAASO, The Obesity Society. Obesity in older adults: technical review and position statement of the American Society for Nutrition and NAASO, The Obesity Society. Am J Clin Nutr. 2005 Nov;82(5):923-34.
- 98. Welle S, Lilavivat U, Campbell RG. Thermic effect of feeding in man: increased plasma norepinephrine levels following glucose but not protein or fat consumption. Metabolism. 1981 Oct;30(10):953-8.
- Segal KR, Albu J, Chun A, Edano A, Legaspi B, Pi-Sunyer FX. Independent effects of obesity and insulin resistance on postprandial thermogenesis in men. J Clin Invest. 1992 Mar;89(3):824-33.
- 100. Goodpaster BH, Park SW, Harris TB, et al. The loss of skeletal muscle strength, mass, and quality in older adults: the health, aging and body composition study. J Gerontol A Biol Sci Med Sci. 2006 Oct;61(10):1059-64.

- 101. Ishigaki EY, Ramos LG, Carvalho ES, Lunardi AC. Effectiveness of muscle strengthening and description of protocols for preventing falls in the elderly: a systematic review. Braz J Phys Ther. 2014 Mar-Apr;18(2):111-8.
- 102. Gosker HR, Wouters EF, van der Vusse GJ, Schols AM. Skeletal muscle dysfunction in chronic obstructive pulmonary disease and chronic heart failure: underlying mechanisms and therapy perspectives. Am J Clin Nutr. 2000 May;71(5):1033-47.
- 103. Gouzi F, Maury J, Molinari N, et al. Reference values for vastuslateralis fiber size and type in healthy subjects over 40 years old: a systematic review and meta-analysis. J Appl Physiol (1985). 2013 Aug 1;115(3):346-54.
- 104. Deutz NE, Bauer JM, Barazzoni R, et al. Protein intake and exercise for optimal muscle function with aging: recommendations from the ESPEN Expert Group. Clin Nutr. 2014 Dec;33(6):929-36.
- 105. Bauer J. Biolo G. Cederholm T. et al. Evidence-based recommendations for optimal dietary protein intake in older people: a position paper from the PROT-AGE Study Group. J Am Med Dir Assoc. 2013 Aug;14(8):542-59.
- 106. Hastings MH, Reddy AB, Maywood ES. A clockwork web: circadian timing in brain and periphery, in health and disease. Nat Rev Neurosci. 2003 Aug;4(8):649-61.
- 107. Whitworth JA, Brown MA, Kelly JJ, Williamson PM. Mechanisms of cortisol-induced hypertension in humans. Steroids. 1995 Jan:60(1):76-80.
- 108. Scheer FA, Hilton MF, Mantzoros CS, Shea SA. Adverse metabolic and cardiovascular consequences of circadian misalignment. Proc Natl Acad Sci U S A. 2009 Mar 17:106(11):4453-8.
- 109. Arendt J, Skene DJ. Melatonin as a chronobiotic. Sleep Med Rev. 2005 Feb;9(1):25-39
- 110. Scheer FA, Morris CJ, Shea SA. The internal circadian clock increases hunger and appetite in the evening independent of food intake and other behaviors. Obesity (Silver Spring). 2013 Mar:21(3):421-3
- 111. Garaulet M, Gómez-Abellán P, Alburguerque-Béjar JJ, Lee YC, Ordovás JM, Scheer FA. Timing of food intake predicts weight loss effectiveness. Int J Obes (Lond). 2013 Apr;37(4):604-11.
- 112. Jakubowicz D, Barnea M, Wainstein J, Froy O. High caloric intake at breakfast vs. dinner differentially influences weight loss of overweight and obese women. Obesity (Silver Spring). 2013 Dec;21(12):2504-12.
- 113. Dhurandhar EJ, Dawson J, Alcorn A, et al. The effectiveness of breakfast recommendations on weight loss: a randomized controlled trial. Am J Clin Nutr. 2014 Aug;100(2):507-13.
- 114. Zilberter T, Zilberter EY. Breakfast: to skip or not to skip? Front Public Health, 2014 Jun 3:2:59
- 115. Willett WC. The dietary pyramid: does the foundation need repair? Am J Clin Nutr. 1998 Aug;68(2):218-9.
- 116. Nestle M. Toward more healthful dietary patterns--a matter of policy. Public Health Rep. 1998 Sep-Oct;113(5):420-3.
- 117. Johnston CS. Uncle Sam's diet sensation: MyPyramid--an overview and commentary. Med Gen Med. 2005 Aug 2;7(3):78.

NEW HIGHLY PURIFIED ALASKAN FISH OIL

The health benefits of eating **cold-water fish** are robust, yet concerns remain about **contaminants** found in wild and farm-raised fish.

This should <u>not</u> stop consumers from including fish in their diet, as the longevity advantages of consuming cold-water **fish** instead of foods like **beef** are substantial.

A recent study found that even **vegetarians** that include some **fish** in their diet fare better than strict vegetarians.¹

Eliminate Virtually All Fish-Derived Toxins

Consumers can exert significant control over their exposure to fish-borne toxins.

High-quality **fish oil** is distilled to remove synthetic and natural contaminants that existed in the **fish** itself. Enhanced molecular distillation techniques utilize redundant processes to virtually eliminate detectable environmental toxins.

The other safety concern about fish oil is that its delicate *omega-3 fats* are highly vulnerable to *rancidity*. No one wants to ingest oxidized (rancid) oils.

A new fish oil blend derived from pristine waters off the coast of **Alaska** utilizes a **multistep** process to remain exceptionally **fresh**. The result is that this **Alaskanderived** fish oil has a greater than **5-fold** <u>reduction</u> in the upper level threshold measurement for **oxidation**.

Current oxidation standards for quality fish oils ensure products free from rancidity. The <u>new Alaskan-derived</u> fish oil specification advances this premium standard **5-fold** better!

The chart below reveals the <u>reduction</u> in upper limit for **oxidation** of this <u>new</u> **Alaskan fish oil** blend over existing **quality** fish oils:

Higher Percentages Of EPA And DHA

An advantage to higher EPA and DHA fish oil concentrations is smaller sized omega-3 softgels.

The addition of this <u>new</u> **Alaskan-derived** fish oil to the **Super Omega-3** supplement group enables the same high-potency **EPA/DHA** to fit into slightly smaller softgels for easier swallowing.

International Fish Oil Association "Five-Star Rating"

The International Fish Oil Association (IFOS) is an independent organization that tests fish oils to determine their overall safety and quality. A Five-Star Rating indicates fish oils have been tested to meet very strict standards of quality as determined by EPA and DHA content, and for purity to rule out contamination with heavy metals, radiation, oxidation, and organic pollutants such as PCBs and dioxin.

The <u>new Alaskan-derived</u> fish oil enjoys the same **Five-Star Rating** mandated for all fish oils contained in the **Super Omega-3** family of supplements.

Sustainable Fishing

The Marine Stewardship Council is an independent nonprofit organization that sets a standard for **sustainable fishing** so that fishing can continue indefinitely with minimal environmental impact.

The <u>new</u> **Alaskan-derived** fish oil is the first refined omega-3 concentrate available worldwide that carries the prestigious **seafood sustainability** certification from the Marine Stewardship Council.

| Alaskan-Derived Specification | Current High-Quality Specification |
|----------------------------------|--|
| Maximum: 5 | Maximum: 26 |
| Max: 1.0 meq/kg | Max: 5.0 meq/kg |
| Maximum: 5 | Maximum: 20 |
| | Specification Maximum: 5 Max: 1.0 meq/kg |

Most Advanced Omega-3 Dietary Supplement

From supporting heart health and brain function to balancing the **inflammatory** response, there is no debating the broad-spectrum benefits of omega-3 fatty acids.²⁻⁴

There are hundreds of fish oil supplements on the market. Only one incorporates lifesaving findings to provide omega-3 and olive fruit extracts, along with sesame lignans, in a family of formulas called Super Omega-3.

Fish Oil + Olive Fruit Extract = **Greater Efficacy!**

Research findings indicate that a combination of fish oil and olive oil can support a healthy inflammatory response better than fish oil alone. Super Omega-3 incorporates the benefits of both fish oil and olive fruit extract into a single novel formula. A four softgel serving supplies the equivalent polyphenol content of 8 to 12 tablespoons of extra virgin olive oil.

Sesame Lignans Enhance Fish Oil Efficacy

Studies show that when sesame lignans are added to fish oil, there is a greater safeguard against oxidation along with the EPA/DHA fatty acids being directed toward pathways that help with inflammatory reactions.6

Benefits Of A Mediterranean Diet

The most popular **Super Omega-3** formula provides the following potencies of Mediterranean health benefits in just four smaller softgel capsules:

Four softgels contain:

| Alaskan Wild Fish Oil | Concentrate | 4,000 mg |
|-----------------------|-------------|-------------------|
| Omega-3s | | 2,400 mg |
| EPA | 1,400 mg | |
| DHA | 1,000 mg | |
| Polyphen-Oil™ Olive € | | af) 600 mg |
| Sesame seed lignan e | xtract | 20 mg |

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



Super Omega-3

Item #01982 • 120 softgels

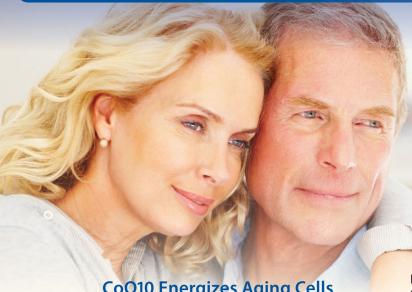
| | Retail Price | Your Price |
|------------|-----------------|---------------|
| 1 bottle | \$32 | \$24 |
| 4 bottles | | \$21 each |
| 10 bottles | | \$17.05 each |

Non-GMO

- 1. JAMA Intern Med. 2015;175(5):767-76. 2. Public Health Nutr. 2006 Dec;9(8A):1136-40. 3. Am J Prev Med. 2005 Nov;29(4):335-46. 4. J Am Diet Assoc. 2005 Mar;105(3):428-40. 5. Nutrition. 2005 Feb;21(2):131-6. 6. Biochem Biophys Acta. 2004 Jun 1;1682(1-3):80-91.

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

MAINTAIN YOUTHFUL MITOCHONDRIA FUNCTION



CoQ10 Energizes Aging Cells by Enhancing Mitochondria Activity

Published studies on CoQ10 absorption clearly show that **ubiquinol** CoQ10 is superior to the conventional ubiquinone form. In middle-age mice, **ubiquinol** proved **40%** more effective in slowing measurements of aging compared to ubiquinone.¹

A 2014 study further validates that **ubiquinol** activates mitochondrial functions to <u>slow</u> aging in mouse models.²

Life Extension® goes one step further and adds **shilajit** to its ubiquinol formula in a product called **Super Ubiquinol CoQ10**. Shilajit has been shown to promote mitochondrial metabolism, helping mitochondria convert fats and sugars into ATP—the main source of cellular energy.³⁻⁸ When combined with **ubiquinol** CoQ10, it has been shown to **double** levels of CoQ10 in the mitochondria.⁹

The latest studies reveal that when **shilajit** is combined with CoQ10, cellular energy substantially increases. Combining the two ingredients produced a **56%** increase in energy production in the brain—**40%** more than CoQ10 alone! In the muscles, there was a **144%** increase!¹⁰

References

- 1. Exp. Gerontol. 2006 Feb;41(2):130-40.
- 2. Antioxid Resox Signal. 2014 Jun 1;20(16):2606-20.
- Ghosal S. Shilajit in Perspective. Alpha Science International Limited; 2006.
- 4. Kaneka Corp. Unpublished study. 2007.
- 5. Sci Total Environ. 1987 Apr;62:347-54.
- 6. Environ Sci Technol. 2002 Jul 15;36(14):3170-5.
- 7. Environ Sci Technol. 2002 May 1;36(9):1939-46.
- 8. Environ Sci Technol. 2009 Feb 1;43(3):878-83.
- Systemic CoQ level in animals: Part II. Unpublished study. Natreon, Inc.; 2007.
- 10. Pharmacologyonline. 2009;1:817-25.

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

The Most Effective Form Of COQ10

Life Extension® combines these two energy-activating ingredients in an exclusive **ubiquinol-shilajit** formulation available in three different potencies:

Super Ubiquinol CoQ10

Item #01425 • 100 **50 mg** softgels **Non-GMO**

| | Retail Price | Your Price |
|------------|-----------------|---------------|
| 1 bottle | \$58 | \$43.50 |
| 4 bottles | | \$34.50 each |
| 10 bottles | | \$31.50 each |



Item #01426 • 60 **100 mg** softgels **Non-GMO**

| | Retail Price | Your Price |
|------------|-----------------|---------------|
| 1 bottle | \$62 | \$46.50 |
| 4 bottles | | \$39 each |
| 10 bottles | | \$36 each |



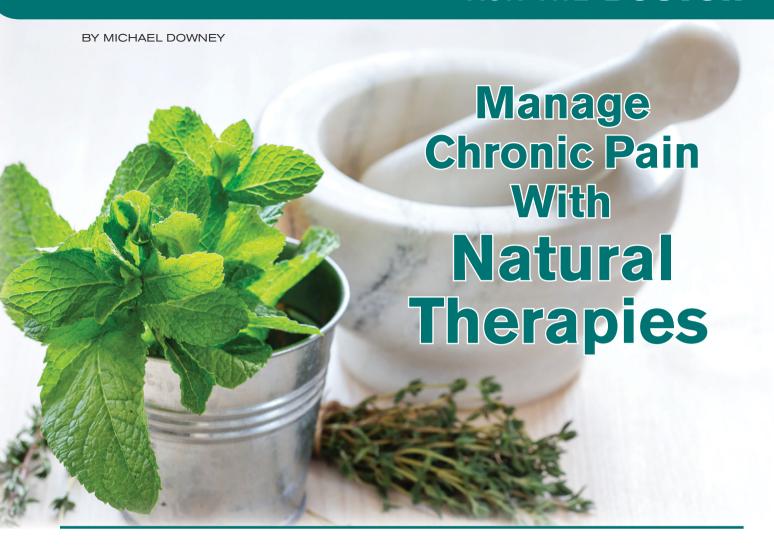
Item #01431 • 30 **200 mg** softgels **Non-GMO**

| | Retail Price | Your Price |
|------------|-----------------|---------------|
| 1 bottle | \$62 | \$46.50 |
| 4 bottles | | \$39 each |
| 10 bottles | | \$36 each |



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

ASK THE DOCTOR





Dr. Chris Kleronomos

A study published in the Journal of Pain reports that 19% of American adults-almost one in five-suffer from persistent pain. Among people over 60, that portion reaches almost 30%.1 Most doctors approach pain problems with a prescription pad, without ever considering other factors that can help mitigate and eventually end chronic pain. But Dr. Chris Kleronomos is an exception. He is a doctor of acupuncture and Oriental medicine, a nurse practitioner, and clinical director of the Fibromyalgia and Neuromuscular Pain Center of Oregon. In a thoroughly comprehensive approach to pain management, Dr. Kleronomos' clinic incorporates natural medicine, acupuncture, chiropractic, psychology, diet, biofeedback, injections, bee venom therapy, functional nutrition, and many other therapies. For those unable to visit his Salem, Oregon, clinic, he recently agreed to answer some questions for us about chronic pain and how it can be safely alleviated.

ASK THE DOCTOR

E: You started out treating pain as a Navy Corpsman and now operate a multimodal pain management clinic focused on therapies rooted in traditional and natural medicine. Can you explain how you first became interested in this field and the journey to where you are now?

CK: Yes, I was a Special Amphibious Reconnaissance Corpsman with the Marine Corp's elite Force Recon unit. So my interest [in pain management] was really born of the necessity to have field medical and survival options available—since we would often be alone and unsupported for extended periods. I began studying herbal therapies and natural techniques that indigenous people had used when fighting guerilla wars. When I got out of the service, I continued that journey with a Master's in Traditional Chinese Medicine. I then spent time in Southeast Asia working in animal conservation and learned directly from the indigenous people who worked for me. I eventually came back to the States and completed a doctoral program that focused on advanced natural therapies for oncology, chronic disease, and pain management. Due to my belief in integrated medicine, I went on to become an Advanced Practice Nurse Practitioner and eventually. completed an additional Master of Science in Functional Medicine and Clinical Nutrition.

Wany people suffer for years, even the rest of their lives, with chronic pain as a result of incurable conditions such as rheumatoid arthritis or fibromyalgia. Since pain medications seldom provide complete relief, are there other options for these patients?

CK: Lots of options are available for these people, and the research clearly shows that multimodal approaches produce the best outcomes. There is no "magic bullet." Diet therapy, in my opinion, is critical. Over the last several decades, we have seen a huge increase in allergies, asthma, and autoimmune disease, and it has been identified that chronic inflammation plays a critical role. Specific options such as acupuncture, chiropractic, psychology, hydrotherapy, supplements, and herbs all depend on the individual patient and the situation. Not every option works for everyone—just as not every medication works-and finding the right combination can be challenging. We approach every person as a unique case.

LE: Long-range side effects, or the risk of them, cause many patients to stop using, or cut back on, pain drugs. Clearly, patients are better off with an effective program of supplements and herbs, right?

CK: The majority of supplements and herbs are safe, but they still must be [used] appropriately and in context of the patient's condition and his or her other medications. Many supplements and herbs treat the contributing mechanisms of pain, such as inflammation or muscle spasms, help to reduce or prevent the escalation of pain medication dosing, or even increase the effect. Another strategy is to use supplements and herbs to address the associated side effects, such as anxiety or insomnia.

Many chronic pain sufferers find that drug painkillers dull, but don't eliminate, their pain. Surely, natural pain relievers cannot provide stronger pain suppression, can they?

CK: Generally speaking, the cumulative evidence for supplements and herbs is limited but excellent for what has been studied. My experience is that natural pain relievers can be as effective. and since they are applied differently and in combination, they can sometimes be stronger. It is really a matter of perspective. A narcotic medication has a strong, specifically targeted effect, but does not always address the root or underlying problem. For example, Corydalis, the classic "pain" herb is said to have a 1% equivalency to morphine—but it's also anti-inflammatory, decreases blood viscosity, and acts as a mild sedative and muscle relaxant. in addition to having some adaptogenic or balancing properties to the system. So, in many ways, it could be considered superior to prescription medications.

LE: Anti-inflammatory medicines such as aspirin, ibuprofen, or naproxen work by blocking the enzymes that trigger inflammation, swelling, and pain. Do nutritional and herbal supplements generally work by modulating the same mechanisms or by other, novel pathways?



Fibromyalgia: What Causes-And What Is-This Source Of Chronic Pain?

Although it is a common source of chronic pain, fibromyalgia is a disease that-unlike other painful conditions such as arthritis-does not show up on X-rays or in blood-work. Unfortunately, there is no lab test to confirm the diagnosis of fibromyalgia. It is primarily a diagnosis of exclusion, which means that other diseases and disorders must first be ruled out.

Although it is not well understood, fibromyalgia is identified as a neurosensory disorder characterized by disturbances in the way the central nervous system interprets and evaluates stimuli.2

Typically, it is associated with other regional pain syndromes, as well as mood and anxiety disorders. In fact, significant data supports the idea that fibromyalgia, chronic fatigue syndrome, regional chronic pain syndromes, and some emotional disorders all involve abnormal perturbations of the stress response system.3,4

Because many fibromyalgia patients appear well on physical examination, the diagnosis of fibromyalgia was historically considered controversial and unfortunately, written off by many conventional physicians as a psychosomatic condition.5,6

We asked Dr. Kleronomos to provide his assessment of the possible origins and mechanisms of fibromyalgia. Here is his analysis of the complexities of this source of chronic pain:

"Currently there is no definitive theory of fibromyalgia. Some critical aspects include increased central sensitization, altered pain signaling, sensory processing issues, hypothalamus-pituitary-adrenal dysfunction, and neurotransmitter imbalance.

"Emerging research performed by our psychologist, who is also my wife, has identified the relationship between 'dual-trauma' exposurehaving a trauma in both childhood and adulthood-and the development of fibromyalgia. [It] showed that the cascade of endocrine changes seen in severe trauma was similar in fibromyalgia patients, further supporting the concept of changes in brain neuroplasticity [your brain's ability to reorganize itself by forming new neural connections throughout life].

"My working theory is that there are four primary presentations of fibromyalgia, each with overlapping features.

- · Gut-mediated: This is associated with food sensitivities, dysbiosis [microbial imbalance on or inside the body], and leaky gut-which, admittedly, is a 'chicken-and-the-egg' pattern and tends to be the most predominant in long-standing cases.
- Structural: Typically, it has an onset of a triggering event, even a minor one, such as a car accident or a fall, and it may be the result of a longstanding, peripheral injury that became centralized or more widespread.
- · Psychogenic: This is related to neuroplasticity [and results] from sustained stress and trauma and shows increased activity in limbic structures [brain structures that govern emotions and behavior].
- Metabolic/nutrient: This is really 'pseudo-fibromyalgia syndrome,' resulting from deconditioning [a physiological decline in function], systemic inflammation, nutrient deficiencies, and other organic issues such as anemia, hypothyroidism, or chronic infections.

"The specific protocol we use at our clinic can ferret out what presentation it is-and address each in a variety of ways."

CK: [It depends.] Willow bark (Salix Spp.), from which modern aspirin is derived, has generally the same effects as the pharmaceutical. It has strong, graded evidence for use, but we know the analgesic actions are typically slow-acting, yet last longer than standard aspirin products. Boswellia spp., a resinous herb with a long history, has demonstrated in animal studies an analgesic effect comparable to a moderate dose of morphine, but it's known to also inhibit both of the major inflammatory pathways. Turmeric (Curcumin Spp.) is gaining lots of attention for its health and anti-inflammatory benefits. It does this by selectively inhibiting both major pathways of inflammation and the resulting downstream chemical triggers of inflammation. Interestingly, it also increases the potency of standard pharmaceutical nonsteroidal anti-inflammatory drugs, such as celecoxib. It has a variety of other health benefits, acting as a free radical scavenger, antineoplastic [tumor-inhibitor], antimicrobial, and it improves cholesterol synthesis.

LE: You provide a wide range of in-clinic therapies, such as injections. Can you give us an outline of these diverse therapies?

CK: Our clinic employs a variety of techniques. We use acupuncture, chiropractic, and physical modalities such as cupping, gua sha [instrument-assisted scraping of the skin], and Graston [a patented, instrument-assisted technique for breaking up scar tissue]. I also use several injection therapies, including trigger point muscle injections [injections into muscle knots], utilizing a combination of natural substances such as MSM—and anesthetic, as well as prolotherapy [injection

ASK THE DOCTOR

of irritating substances] for the neck, back, and joints. I'm probably most well-known, however, for the use of bee venom therapy. Bee venom, sometimes called apipuncture, has documented use going back thousands of years. Charlemagne is said to have used it to treat his gout during the Crusades, and Alexander the Great reportedly used it for chronic hip pain. Modern research is now able to validate the mechanisms of its action, which includes blocking peripheral pain-signaling, antiinflammation, central or brain modulation of pain, and immune regulation. Classically, it has been best known [to be] utilized for arthritic conditions, and in the US, has a following for use in multiple sclerosis. I have found it extremely effective for multiple pain conditions including fibromyalgia, rheumatoid arthritis, lupus, and complex regional pain syndrome.

LE: Aside from these in-clinic pain therapies, can you explain what supplement options are available that effectively treat chronic pain?

CK: A good, basic start is a whole-food multivitamin-some data suggests that you would need 27,000 calories to meet all of the recommended daily intake for micronutrients.7 This accounts for the most common nutrient deficiencies in the US, which are iron, calcium, magnesium, and D and B vitamins. Generally, I believe that most people need foundation support, depending on how good their diet is, where they live, and what underlying issues they have: an average dose of 4,000 IU daily of vitamin D3; 2,000 to 3,000 mg daily of omega-3; multi-strain live probiotic; and active B vitamins in

methylated forms. Also, a multimineral with calcium (600 mg twice daily), magnesium (1,000 to **2,000 mg** daily), and zinc (**20** mg daily), particularly if currently on an antacid medication or diuretic. I [have my patients] use iron with vitamin C if there is anemia, but I am cautious if there is a lot of inflammation. I always add CoO10 (100 to 300 mg), especially if they are on a statin drug. Other basics include glucosamine, chondroitin, and MSM, especially if there are degenerative changes. Selenium is a good option if [patients] have a thyroid issue—55 micrograms as basic, 200 micrograms if autoimmune. Using either NAC [N-acetyl cysteine] or glutathione can be useful. Green tea and garlic should be included in most people's regimen, either in food form or extract. These herbs have very broad effects that target several aspects of health. Green tea or EGCG can induce repair of cells at all stages of the cell cycle.

LE: You said that treatment choices depend on the individual patient. Do supplement recommendations also vary with each individual case?

CK: Yes. The individual's body effect on the supplement or herb—called pharmacokinetics is important. This can include sensitivities to fillers or the initial source from which it was derived. the amount of stomach acid available to break it down, and the integrity of the gastrointestinal tract, particularly the health of the microbiome. Individual genetics play an important role, especially in B vitamin synthesis, cellular energy production, detoxification, and in the CYP450 system, which is the liver's ability to process and detoxify.

LE: Is it safe to mix traditional drug pain relievers with supplement pain remedies?

CK: Yes, a review of the literature demonstrated that beliefs about herb-drug interactions are mainly theoretical considerations and not clinically observed facts. Drug interactions with herbs or nutrients do occur, but occur equally to common foods such as broccoli, grapefruit juice, and alcohol. This is also true of cigarette smoking. Certainly, it's far less of a problem than the over one million [Americans] injured or killed directly related to prescription medication error.

LE: Is exercising helpful and for what painful conditions?

CK: I often say that "motion is lotion." No question: Exercise is important for a lot of issues relating to chronic pain, including depression and anxiety. It's suitable for most chronic pain conditions in which furthering structural damage is not a risk—and for those risks, we recommend guidance from a physical therapist. Exercise helps strengthen the core, stabilize structures such as the spine and joints, and improve posture, and it's also important for reducing negative metabolic consequences such as obesity, high blood pressure, elevated blood sugar, and cholesterol. It even improves cellular energy production by the mitochondria...the recommendation is to maintain a moderate activity level overall, combining various activities such as walking, warm water swimming, and yoga.

E: Beyond supplements and exercise, do any other lifestyle changes improve pain?

CK: A healthy diet...addresses multiple systems simultaneously to

reestablish and support the body's ability to repair and heal. Many studies show that people who eat certain types of foods are less likely to have health issues. Although many fad diets are in the marketplace, there is a common theme of removing potentially problematic foods, increasing micronutrients, restoring healthy gut flora, and reestablishing an appropriate omega-6:omega-3 ratio to decrease inflammation. Carbohydrate restriction and avoidance of refined, processed, and GMO foods is a good start. Generally, anything that decreases depression, anxiety, and focuses on the health condition is helpful—including meditation, stress management, community activities, hobbies, recreational activities, time in nature, and relationships with family and friends. We [use] cognitive behavioral techniques, biofeedback, mindfulness, relaxation exercises, and guided imagery.

LE: For many patients, medical practitioners cannot identify a clear cause. Is there any way to track down the source of their pain?

CK: For complex conditions, there are multiple pieces to the pain "puzzle." I find a functional medicine approach [focusing on interactions between the environment and the gastrointestinal, endocrine, and immune systems] to be the most efficient in seeking antecedents, triggers, and mediators of the condition. I also look at blood work differently in an attempt to identify patterns. Specialized tests can be useful, such as salivary adrenal and hormone profiles, micronutrient assays, and stool analysis. I [emphasize] finding the underlying dysfunction and correction of systems-level concerns.

LE: Should people who are on blood-thinning drugs check with their doctor before using pain supplements?

CK: Yes, it's important to have a partnership with your health care provider before starting an herb or supplement regimen—research everything and ask questions. Although the evidence is not overwhelming for interactions, adverse effects have been documented. This is the one area where even food can have an impact.

LE: Very informative. Thank

CK: My pleasure.

For more information on Dr. Chris Kleronomos, or on the Fibromyalgia and Neuromuscular Pain Center of Oregon, visit http://pain-puzzle.com/ or call 1-844-724-6789.

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

References

- Kennedy J, Roll JM, Schraudner T, Murphy S, McPherson S. Prevalence of persistent pain in the U.S. Adult population: new data from the 2010 national health interview survey. J Pain. 2014 Oct:15(10):979-84.
- Gracely RH, Petzke F, Wolf JM, Clauw DJ. Functional magnetic resonance imaging evidence of augmented pain processing in fibromyalgia. Arthritis Rheum. 2002;46(5):1333-43.
- Buskila D, Sarzi-Puttini P. Biology and therapy of fibromyalgia. Genetics aspects of fibromyalgia syndrome. Arthritis Res Ther. 2006;8:218-22.
- Diatchenko L, Nackley AG, Slade GD, et al. Idiopathic pain disorders pathways of vulnerability. Pain. 2006 Aug;123(3):226-30.
- 5. Available at: http://www.uptodate.com/. Accessed May 1, 2015.
- Goldenberg DL. Fibromyalgia syndrome a decade later: what have we learned? Arch Intern Med. 1999:159(8):777-85.
- Calton JB. Prevalence of micronutrient deficiency in popular diet plans. J Int Soc Sports Nutr. 2010 Jun 10;7:24.



undreds of published studies validate the neurological properties of the <u>individual</u> ingredients contained in **Cognitex**®.

Scientists wondered what would happen if these cognitive-enhancing nutrients were combined into <u>one</u> formula. An open label study was performed to assess the effects of **Cognitex**® on the brain health of elderly human subjects.

Here is a summary from a study that was published in the *Journal of Dietary Supplements* in June 2011:

Study design: 26 elderly participants with memory complaints completed the study. Participants were given three capsules daily of **Cognitex**® for 12 weeks with assessments at two weeks and 12 weeks.

Memory at two weeks:

- Recall: +11.15 points vs. baseline—a 15% improvement
- Recognition: +8.68 points vs. baseline—a 11% improvement
- Spatial short-term memory: +19.85 points vs. baseline—a 42% improvement

Attention (sustained and focused) at two weeks:

- Sustained attention: +9.46 points vs. baseline a 12% improvement
- Focused attention: +3.77 points vs. baseline—
 a 4% improvement

Visuospatial learning at two weeks:

• +17.31 points vs. baseline—a 33% improvement

Activities of daily living (executive functions and mental flexibility) scores at two weeks:

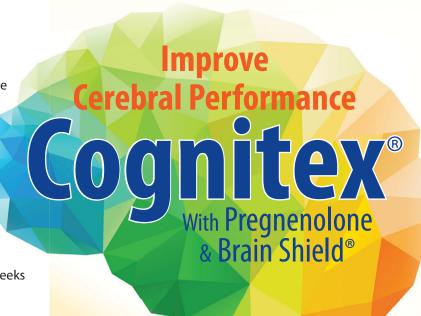
- Executive functions: +9.45 points vs. baseline a 14% improvement
- Mental flexibility: +9.92 points vs. baseline a 15% improvement

After 10 additional weeks of Cognitex® supplementation, further statistically significant improvements in activities of daily living (executive functions and mental flexibility) were observed:

- Executive function: +9.15 points vs. two-week scores—a 12% improvement
- Mental flexibility: +9.73 points vs. two-week scores—a 13% improvement

Cognitex® is designed to improve cerebral performance <u>and</u> protect against neurological problems over the long term. This study evaluated the effects of **Cognitex**® on boosting brain function over a short-term period.

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.



The suggested daily dose of three **Cognitex®** with **Pregnenolone & Brain Shield®** softgel caps provides:

| Alpha-Glyceryl Phosphoryl Choline (A-GPC) | 600 mg |
|---|--------|
| Phosphatidylserine (from Sharp-PS® Green) | 100 mg |
| Brain Shield® (Gastrodin) | 50 mg |
| Vinpocetine | 20 mg |
| Leucoselect [®] Grape Extract (seed) | 150 mg |
| Sensoril® Ashwagandha Extract | 125 mg |
| BlueActiv™ Wild Blueberry Extract | 150 mg |
| Uridine-5'-Monophosphate (disodium) | 50 mg |
| Proprietary NeuroProtection Complex Blend Perluxan Hops extract, Rosemary standardized extract | 125 mg |
| Pregnenolone | 50 ma |

Non-GMO

Cognitex® with Pregnenolone & Brain Shield® (Gastrodin)

Item #01897 • 90 softgels

| | Retail Price | Your Price |
|------------|-----------------|---------------|
| 1 bottle | \$62 | \$46.50 |
| 4 bottles | | \$39.75 each |
| 10 bottles | | \$37.50 each |

To order Cognitex® with Pregnenolone & Brain Shield®, call 1-800-544-4440 or visit www.LifeExtension.com



PROSTATE SUPPORT

In A Once-Daily Softgel

The majority of men find that their **prostate gland** undergoes significant changes as they age.¹ **ProstaPollen**™ supports healthy urination patterns and prostate function.

Triple Strength ProstaPollen[™] contains concentrated pollen extracts specifically selected for effective **prostate** support.²⁻⁴

Clinical studies have demonstrated that the flower pollen extracts in *Triple Strength* ProstaPollen[™] provide healthy support for aging prostate glands.²⁻⁴

Two fractions in *Triple Strength* ProstaPollenTM— $G60^{TM}$ (water-soluble) and NAX^{TM} (lipid-soluble)—support prostate health by helping to maintain smooth muscle tone in the prostate, bladder, and urethra.⁵

Life Extension® has long offered **pollen extracts** in the comprehensive **Ultra Natural Prostate** formula for maintaining healthy prostate function. For the first time, this <u>new</u> *Triple Strength* **ProstaPollen**™ European extract is available to Americans, providing a more potent dose⁶ in a convenient once-daily softgel!

For men using Life Extension®'s **Ultra Natural Prostate** formula, <u>additional</u> prostate support benefits can be found by adding just <u>one</u> *Triple Strength* **ProstaPollen™** softgel daily.*

To order **Triple Strength ProstaPollen**™, call **1-800-544-4440** or visit **www.LifeExtension.com**



The suggested dosage of <u>one</u> softgel of **Triple Strength ProstaPollen**™ provides:

Graminex® Flower Pollen Extract™**(from rye)

Graminex° **G60**[™] (water-soluble fraction) (360 mg) **Graminex**° **NAX**[™] (lipid-soluble fraction) (18 mg)

*Ultra Natural Prostate contains 252 mg of original Graminex extract providing 60 mg of G60™ water-soluble fraction and 3 mg of NAX™ lipid-soluble fraction in two softgels. Men completely satisfied with the effects of the Ultra Natural Prostate formula may not need this new Triple Strength ProstaPollen.

** Graminex® is a registered trademark of Graminex LLC.

References

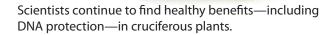
- 1. AJR. 1989 Jan.152:7-81.
- 2. Br J Urol. 1989 Nov;64(5):496-9.
- 3. Br J Urol. 1993 Apr;71(4):433-8.
- 4. Eur Urol. 2009 Sep;56(3):544-51.
- Prostate Support: Graminex flower pollen extract. 2003 Jan:4-9.
- 6. Clin Ther. 1995 Jan-Feb:17(1):82-7.

Not Eating Enough Veggies? No Problem!

Get All The Protective Benefits Of

Cruciferous Vegetables

In One Easy-To-Take Supplement



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

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

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

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

References

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

Triple Action Cruciferous Vegetable Extract

Item #01468 • 60 vegetarian capsules

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

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 |



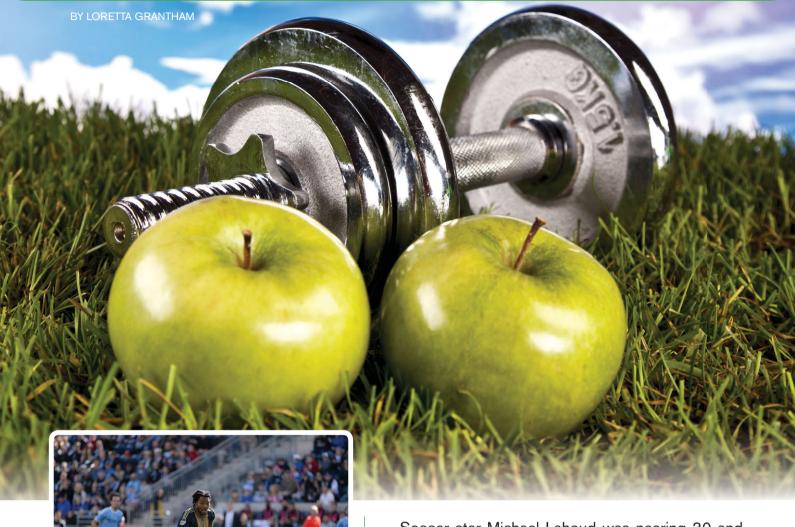
LifeExtension

Cruciferous Vegetable

or Cellular Protection &

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

WELLNESS PROFILE



SOCCER SUPERSTAR

MICHAEL LAHOUD

SAYS VITAMIN D IMPROVED HIS STAMINA AND HIS GAME

Soccer star Michael Lahoud was nearing 30 and wanted an edge.

"When I played in college, youth and talent got me through," says Philadelphia Union midfielder, 29, who graduated from Wake Forest University in 2008 and was a first-round draft pick for Chivas USA, a former Major League Soccer team based in Los Angeles, before joining Union three years ago.

"By the end of college, I'd stopped drinking soda and eating fast food. But that was pretty much it as far as taking care of myself. When you excel as an athlete, you're told most of your life how special and talented you are. But at the professional level, you realize that everyone's special and talented, and in order to consistently stay at the top of your game, you have to do something more."

Remarkable Results

What Lahoud did, upon the advice of Union strength-and-conditioning coach Kevin Miller, was to begin supplementing with vitamin D. And the results, the player says, have been remarkable.

"I noticed an immediate impact within about two weeks, especially more energy," he says. "But not in the sense of drinking an energy booster and getting a jolt—energy in a consistent, sustained way. I'm not a morning person by trade, but I get paid to be a morning person, and I started to feel less foggy when I got woke up. I could dedicate that extra energy to doing more training and the other work that a good pro needs to do."

Vitamin D has long been linked to strong bones by aiding in the metabolism of calcium. It also may help lower the risk of degenerative disease. By directly influencing over 200 human genes, vitamin D notably binds to many genes associated with autoimmune disease and cancer, which has researchers studying vitamin D's potential benefits beyond bone health. The

versatile vitamin is also involved in muscle function, the respiratory system, and brain development.

"I'm always reading about supplements because as a coach, you're constantly looking for ways to give your players an advantage," says Kevin Miller. "Obviously strong bones are important. But power, speed, and recovery from exertion are also key factors. Anything that can reduce the chance of injury or illness is critical when it comes to an athlete."

Team Credits Vitamin D For Its Comeback

Vitamin D started to gain lockerroom respect about five years ago
after the Chicago Blackhawks, following a 50-year championship
drought, won the Stanley Cup in
2010. The hockey team credited its
comeback, in part, to significantly
fewer injuries after most players
started taking a daily **5,000 IU**vitamin D supplement that season.
(The Blackhawks captured the Cup
again three years later.)



Study Shows Effects

Also in 2010, researchers measured the vitamin D levels of all 89 New York Giants football players in the spring, then compared the results to data on players who'd missed a game or practice during the previous season due to injury. The results, presented in 2011 at the annual meeting of the American Orthopaedic Society for Sports Medicine in San Diego, showed that the average vitamin D level of players who sustained injuries was about 20% lower than that of uninjured players regardless of age, height, weight, or body mass index (BMI).

"I was watching the Blackhawks while they were on their way to winning the Stanley Cup the second time, and they mentioned vitamin D again," Lahoud says, referring to the hockey team's 2012-2013 season. "It was funny to think that they turned themselves around with a supplement. I was really inspired by their story, and after we had a rough start to our season, [Coach Miller] brought up vitamin D, and he started the vitamin D challenge."

Miller asked Union players to take a blood test to measure their initial levels of vitamin D and then challenged them to increase their numbers.

"My number one goal for the players is to keep them healthy," says Miller. "Michael was great. He supplemented for about a year, and after we had the whole team tested again, he had the highest score."

Lahoud noted that dark skin makes it even harder to maintain adequate vitamin D levels because it inhibits ultraviolet B (UVB) radiation from sunlight, which is a precursor to the body making vitamin D. "After I began supplementing, my levels rose into the **80s (ng/dL)**," he says.



Recommended Dosages

Nutritional scientists recommend an optimal 25-hydroxy vitamin D blood level of between 50 ng/mL to 80 ng/mL.

The current daily Recommended Dietary Allowance (RDA) of vitamin D is only 600 IU, but widespread evidence of vitamin D deficiencies prompted researchers to increase that number to a daily intake of 1,000 IU. Life Extension® suggests that healthy adults supplement each day with 2,000-8,000 IU of vitamin D and have a blood test done to measure 25-hvdroxvvitamin D levels about 45 days later.

Miller, 43, a Union coach since 2010, is not just a trainer but also a triathlete who pursues peak fitness.

He has competed three times in the grueling Ironman Lake Placid in New York and says that he, too, has experienced the performance boost of vitamin D.

"I recall reading about vitamin D in Life Extension magazine, but I hadn't taken any myself at the time. The article recommended supplementing even if you spend time in the sun because of the weather, sunblock, and other things that affect your body's ability to produce an optimal level. I know Life Extension® is a very reputable company with safe products, so that's the brand I recommend."

Lahoud, for example, takes Life Extension®'s Vitamins D and K with Sea-IodineTM, which contains 5.000 IU of D to facilitate calcium absorption into the bloodstream and 2,100 mcg of vitamin K to transport calcium from the bloodstream into the bone. The midfielder, at Miller's recommendation, also takes L-carnitine, curcumin, zinc, and fish oil.

Feeling Fit On The Field

"I feel more prepared when I take my supplements every day," says Lahoud, an American citizen who plays internationally for his native country of Sierra Leone in high-profile tournaments such as the Africa Cup of Nations. "I some-

times have very long travel days to compete against the best players in the world, and I get an extra sense of being ready for the world stage if I take supplements on a regular

The defensive powerhouse, who says his position requires him to be one of the fittest on the field, emphasizes that you don't have to be a professional athlete to benefit from vitamin D.

"It's about doing your best consistently, whether you're a sports star, business leader, or just an everyday person. The way to do that is to create and stick to habits that involve taking care of yourself, including nutritional support. I've seen firsthand that vitamin D gives you the energy and focus to handle whatever life throws at you."

Benefits Of Vitamin D

Vitamin D is essential for calcium absorption as well as the maintenance of healthy bones and teeth. It also provides a protective effect against many diseases and conditions, such as type I diabetes and multiple sclerosis. Researchers also say that the steroid hormone:

- Boosts immunity,
- Supports the brain and nervous system,
- Regulates insulin levels.
- Strengthens lung function and heart health, and
- Influences the expression of genes involved in cancer growth.

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

Top Off Your
TESTOSTERONE
Naturally

Low Testosterone Levels May Lead to:

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

Maintaining healthy testosterone levels is one of the most important steps you can take to regain your health and improve your performance. With research showing that by the time a man is 70 years old, he may produce 60% less testosterone than he did at age 40, the time is now to add Life Extension®'s Super MiraForte with Standardized Lignans to your supplement regimen.

Super Miraforte with Standardized Lignans • Item #01698 • 120 capsules

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

Non-GMO

To order Super MiraForte with Standardized Lignans call 1-800-544-4440 or visit www.LifeExtension.com



Each daily dose of **Super MiraForte with Standardized Lignans** contains the following testosterone supporting ingredients:

| Chrysin | 1,500 mg |
|---|----------|
| Bioperine® | 15 mg |
| Muira puama | 850 mg |
| Nettle root | 282 mg |
| Chelated elemental zinc | 15 mg |
| Maca | 320 mg |
| HMRlignan™ Norway Spruce lignan extract | 33.4 mg |

Super Health. Super Libido. Super MiraForte.

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.



VITAMIN D3 SOFTGELS

For Superior Absorption

New research on the vital benefits of **vitamin D** emerges on a daily basis. Studies confirm that **optimal** levels of vitamin D are in the range of **50-80 ng/mL** of **25-hydroxy vitamin D**. **Life Extension**® has created a large selection of highly *absorbable* **vitamin D** supplements in softgels to help you to achieve your individual **vitamin D goals**. Keep in mind that you may already be getting **1,000-3,000 IU** of **vitamin D** in your current multi-nutrient formulas.



Vitamin D3 • 1,000 IU

250 softgels • Retail: \$12.50 Four bottles: **\$8.44** ea.

For most people, a **1,000 IU** potency is insufficient to attain optimal **vitamin D** blood levels. However, this potency may be suitable for smaller individuals who obtain **2,000-3,000 IU** in their multi-nutrient formulas (and children). **Item # 01751**Non-GMO.



Vitamin D3 • 5,000 IU With Sea-lodine™*

60 capsules (non-softgel) • Retail: \$14

Four bottles: \$9.38 ea.

Most people, especially those seeking to reduce their salt intake, do not ingest enough iodine. Combining **5,000 IU** of **vitamin D3** with **1,000 mcg** of **iodine** into one capsule makes taking these two nutrients economical and convenient. Due to the source of kelp, this product may contain fish and shellfish. Item # 01758



Vitamin D3 • 5,000 IU

60 softgels • Retail: \$11 Four bottles: **\$7.43** ea.

For those already obtaining **1,000-3,000 IU** of **vitamin D** in their multi-nutrient formulas, this **5,000 IU** potency is what may be needed to achieve optimal blood levels. **Item # 01713**



Vitamin D3 • 7,000 IU

60 softgels • Retail: \$14

Four bottles: \$9.45 ea.

Some people (such as those weighing more than 180 pounds) may require even more **vitamin D**. When combined with **1,000-3,000 IU** taken in a multi-nutrient formula, this **7,000 IU** softgel should enable these individuals to attain blood levels above **50 ng/mL**. Item # 01718



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

1 ounce • Retail: \$28 • Non-GMO Four bottles: **\$18.75** ea.

Great for travel and for those individuals who have difficulty absorbing enough vitamin D3 from softgels, this liquid vitamin D is ideal. Item # 01732

Also available without mint. (Item # 00864)

To order any of these high-potency vitamin D3 supplements, call 1-800-544-4440 or visit www.LifeExtension.com

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

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

THE MOST OPTIMAL FORM OF VITAMIN E

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

Prestigious scientific journals have highlighted gamma **tocopherol** as one of the most critically important forms of tocopherols, which includes d-alpha tocopherol (natural vitamin E) for those seeking optimal health benefits.

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

SESAME LIGNANS: THE NATURAL VITAMIN E BOOSTER

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

In a human study that combined gamma tocopherol with **sesame lignans**, gamma tocopherol/sesame was 25% more effective than gamma tocopherol/tocotrienols in suppressing tissue measurements for free-radical and inflammatory damage.4,5

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

WORLD'S MOST COMPREHENSIVE VITAMIN E FORMULA!

The Gamma E Tocopherol with Sesame Lignans formula provides potent doses of critically important gamma tocopherol along with sesame lignans to augment its antioxidant effects. Suggested dose is one softgel once or twice daily.

| 45 IU |
|--------|
| 359 mg |
| |

Contains soybeans.

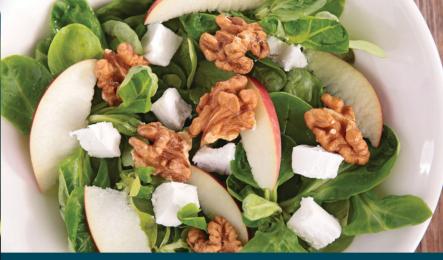


To order Gamma E Tocopherol with Sesame Lignans. call 1-800-544-4440 or visit www.LifeExtension.com

References

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



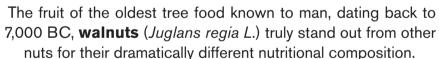


SUPER FOODS

BY MICHAEL DOWNEY

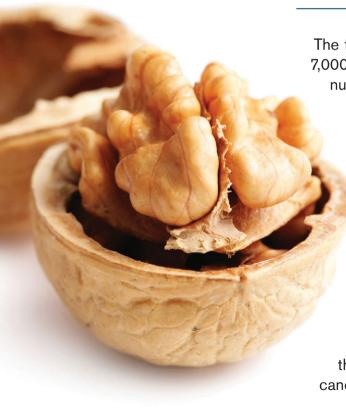
Walnuts

Abundant Disease-Prevention Benefits



The healthy fats in most nuts are primarily monounsaturated fatty acids. But walnuts are rich in polyunsaturated fatty acids¹-including omega-3 and omega-6 fatty acids-in addition to oleic acid, the powerful monounsaturate found in olive oil. In fact, the walnut is the only nut and one of the few foods that contains a significant amount of the lesser known omega-3 fat, alphalinolenic acid (ALA)-containing the highest ALA content of all edible plants² with a hefty **2.7 grams** per quarter-cup serving!³

And walnuts provide several rare and potent phytonutrients that are found in few commonly eaten foods. These unique compounds include the quinone *juglone*, the tannin *tellimagrandin*, and the flavonol *morin*⁴–all strong cancer inhibitors.



SUPER FOODS

Like other nuts, walnuts contain vitamin E—but instead of having most of their vitamin E present in the alpha-tocopherol form, walnuts provide an unusually high level of the particularly beneficial gamma-tocopherol form.⁴

Toss in walnuts' natural array of nutrients including melatonin, copper, manganese, molybdenum, biotin, folate, selenium, and magnesium—along with the rich supply of protein and fiber also found in other nuts—and you have a true super food.

"A Remarkable Radical-Scavenging Effect"

Underlying the walnut's compelling disease-inhibiting benefits is an abundance of polyphenols, as well as the rare, anticancer phytonutrients juglone, tellimagrandin, and morin.⁴ And these super-nuts are particularly rich in the phenolic compound ellagic acid, which has shown potent anti-inflammatory properties.⁶



An analysis by scientists who extracted the polyphenols from various raw and roasted nut types found that walnuts had the highest total polyphenols and the highest polyphenol efficacy.⁷

Researchers also investigated just some of walnut's tannins and ellagitannins and found that 14 different walnut polyphenols provided superoxide dismutase (SOD)-like activity and "a remarkable radical scavenging effect."8

Current evidence strongly suggests that polyphenols play a critical role in the prevention of atherosclerosis⁷ and other cardiovascular diseases, cancers, osteoporosis, neurodegenerative diseases, and diabetes mellitus.⁹ Nut supplementation has been shown to improve lipid profile, increase endothelial function, and reduce inflammation, all without causing weight gain.⁷ Walnut polyphenols in particular have been shown to inhibit, for example, chemically induced liver damage.¹⁰

Walnuts Cut The Risk Of Cancers

Walnuts have been shown to inhibit cancer development, slow its growth, and induce the death of cancer cells.

In one study, scientists found mice that ate the human equivalent of 2.4 ounces of whole walnuts for 18 weeks had substantially smaller and slower-growing prostate tumors compared to mice that consumed the same amount of fat but from other sources. Prostate cancer growth in the walnut group was 30 to 40% reduced, and these mice had lower blood levels of a protein (insulin-like growth factor or IGF-1) strongly linked prostate cancer. Also, there were beneficial effects on multiple genes related to controlling tumor growth and metabolism.11

In a study on breast cancer in mice, the human equivalent of two handfuls of walnuts daily cut breast cancer risk in half and slowed tumor growth by **50**%.¹²

Researchers then conducted a cell study to investigate whether walnut components have an effect on colon cancer stem cells. (Cancer stem cells are a subpopulation of cancer cells that can self-renew and undergo differentiation into multiple lineages, key elements of cancer progression.) Walnut extract significantly slowed the survival of the cancer stem cells and most striking, reduced the essential characteristics of their self-renewal capacity.¹³

Walnuts provide potent but very rare phytonutrients that block cancer. Scientists found that incubation of human liver cancer cells with the walnut quinone juglone caused obvious, destructive changes to the structure of the cancer cells—and induced their apoptosis.¹⁴

Why You Should Eat The Walnut Skin

Some food experts suggest removing the paperthin, outermost layer of a shelled walnut, citing the fact that they have a bitter taste. We're talking about the very fine, whitish outer part that is either flaky or sometimes, waxy. But you should definitely resist any advice to remove this skin from shelled walnuts. It's believed that up to 90% of walnuts' abundant phenols are found in this skin, making it the healthiest part to eat.

Look for walnuts that are raw and unpasteurized. Note that walnuts are highly perishable due to their high content of polyunsaturated fatty acids, which are easily damaged. If you buy shelled walnuts in bulk, avoid those that appear shriveled or rubbery, smell rancid, or that you cannot verify are fresh. Shelled walnuts should be stored in an airtight container and placed in the refrigerator, where they will keep for six months, or in the freezer, where they will last for a year. Unshelled walnuts should preferably be stored in the refrigerator, although as long as you keep them in a cool, dry, dark place they will stay fresh for up to six months.

Shelled walnuts are great as a quick snack just as they are. Or add them to your favorite salad, vegetable dish, fruit, or dessert-mix crushed walnuts into plain yogurt. But if you don't like the slightly bitter flavor, you can still get their rich therapeutic benefits by blending them into smoothies.



Other rare phytonutrients in wal-

nuts are known as tellimagrandin I and tellimagrandin II. Scientists identified these walnut members of the ellagitannins family as the predominant compounds responsible for the previously observed cytotoxic effect of walnuts on cancer cells. They also demonstrated that the mechanisms through which these walnut phytonutrients attack cancer cells include impairment of their mitochondrial function and induced apoptosis.15

Another rare and powerful phytonutrient found in walnuts is the flavonoid morin. A 2015 study published in the International Journal of Oncology reported that morin exerts strong anticancer activity against human colon cancer cells. It was determined to accomplish this by inducing generation of reactive oxygen species (ROS), suppressing anti-apoptotic proteins, and triggering apoptosis via several different pathways.¹⁶

Walnuts provide the amino acid L-arginine, shown to deliver vascular benefits to individuals who have heart disease or who have a higher risk for it due to multiple factors. 17

Effective

Cardiovascular

Protection

More critically, walnuts also contain the omega-3 fat alphalinolenic acid (ALA), which has anti-inflammatory effects and may prevent the formation of pathological blood clots. Those who eat an ALA-rich diet are less likely to have a fatal heart attack and have an almost 50% reduced risk of sudden cardiac death.¹⁸ It should be noted, however, that as people age, the *enzyme* required to convert ALA into beneficial EPA/ DHA declines, meaning that walnuts may not provide sufficient omega-3s. That's why fish oil supplementation is so important, even in those who consume ALA-rich foods like walnuts.

And it doesn't take a lot of walnuts to make a huge difference. A study found that blood levels of ALA were significantly raised by eating just four walnuts a day!19

Other research showed that eating just one ounce (seven walnuts) daily can decrease overall cardiovascular risk. 12 Also, increased frequency of nut consumption among people at high cardiovascular risk was found to result in a significantly lower risk of death.20

Walnut consumption supports healthful cholesterol levels. Scientists found that, when compared with control diets, walnutrich diets produced significantly greater decreases in total and LDL-cholesterol concentrations, with weighted mean decreases of 10.3 and 9.2 mg/dL, respectively.21

In the April 2015 issue of the FASEB Journal, scientists reported on an investigation into the effects of short-term walnut consumption on the reactive hyperemia index (RHI), a measure of human microvascular function—referring to the health of your body's smallest blood vessels such as capillaries. After just four weeks, the group of volunteers who ate 40 grams, or 1.5 ounces, of walnuts daily was found to have improved vascular function. The study concluded that this improvement was strongly associated with the effects of the walnut fatty acids, alpha-linolenic acid (ALA) and linoleic acid (LA).²²

Walnuts stand out because they contain an unusually high level of the gamma-tocopherol form of vitamin E instead of the alpha-tocopherol form. Gamma-tocopherol has been demonstrated—particularly in studies on the cardiovascular health of men—to provide significant protection from heart problems.⁴

Enhanced Protection Of Cognitive Function

Compounds in walnuts—gamma-tocopherol, folate, alpha-linolenic acid, omega-3 fatty acids, and antioxidant polyphenols—contribute to neuroprotection.

Animal research has indicated that consumption of foods such as blueberries and walnuts can decrease vulnerability to oxidative stress that occurs in aging and "may increase 'health span' and enhance cognitive and motor function in aging."²³

In a study on young and old rats, scientists noted the gene transcription effects—in the critical hippocampus region of the brain—of a diet with a walnut content equivalent to **1.5 ounces** in a human diet. Compared to controls, the walnut diet increased transcription of a particular immediate-early gene (IEG) that is essential to memory formation and synaptic plasticity. This prompted researchers to conclude "that dietary walnut may have protective effects on the aging brain."²⁴

Scientific attention was then shifted to humans. A crossover study published in the *British Journal of Nutrition* included 64 college students who were randomly assigned to follow an eight-week walnut diet and an eight-week placebo diet, with a sixweek washout period between diet

sequences. The walnut diet was demonstrated to boost inferential verbal reasoning by a significant **11.2%**. ²⁵

Further research has led scientists to suggest that walnuts may play a key role in the prevention of Alzheimer's disease.

First, a lab study found that walnut extract inhibited the formation of fibrillar amyloid beta-protein—a principal component of the amyloid plaques in the brains of Alzheimer's patients. Remarkably, walnut extract was also able to defibrillize already formed amyloid fibrils. The researchers concluded that "walnuts may reduce the risk, or delay the onset, of Alzheimer's disease by maintaining (fibrillar amyloid beta-protein) in the soluble form." ²⁶

Then, scientists conducted a study with transgenic mice that modeled Alzheimer's. Compared to the control diet, a diet containing the equivalent in human terms of **1.5 ounces** of walnuts daily significantly improved memory, learning ability, anxiety, and motor development. The study author suggested that "walnuts may have a beneficial effect in reducing the risk, delaying the onset, or slowing the progression of, or preventing [Alzheimer's disease]."²⁷

Finally in 2015, scientists investigated the cognitive and potentially anti-Alzheimer's effects of walnut consumption on humans aged 20 through 90. Controlling for all factors except walnut consumption, the researchers found that, compared to those who ate no nuts, those who ate a handful of walnuts daily performed substantially better on all six cognitive exams, taking considerably less time to finish questions and tasks. Crucially, those over age 60 reaped special benefits from walnut consumption, achieving recall



SUPER FOODS

Nutritional Content Of Walnuts

Just one ounce (28 grams) of shelled walnuts provides:35

| Calories | 185 |
|--|------------|
| Protein | 4.3 grams |
| Total fat | 18.4 grams |
| Cholesterol | 0.0 mg |
| Monounsaturated fat | 2,500 mg |
| Saturated fat | 1.7 grams |
| Polyunsaturated fat | 13.3 grams |
| Omega-3 fatty acids (precursor to EPA/DHA) | 2,565 mg |
| Omega-6 fatty acids | 10,761 mg |
| Trans fat | 0.0 grams |
| Thiamin | 0.1 mg |
| Vitamin B6 | 0.2 mg |
| Folate | 27.7 mcg |
| Manganese | 1.0 mg |
| Magnesium | 44.6 mg |
| Phosphorus | 97.8 mg |
| Copper | 0.4 mg |
| Phytosterols | 20.3 mg |
| Dietary fiber | 1.9 grams |

and other cognitive test scores that were more than seven percentile points higher than those of the same age group who did not eat walnuts.28

Other Effects

Various studies have suggested that walnuts may also provide other health benefits:

• Overweight adults with type II diabetes who ate one-quarter cup of walnuts daily reduced fasting insulin levels, compared to those who did not, within iust months.29

- An animal study found that a diet containing the human equivalent amount of two ounces of walnuts significantly altered the ratio of two major. gut bacteria communities, suggesting "a new mechanism...by which walnuts may exert their beneficial health effects."30
- Consuming walnuts has, after just three days, been associated with increased satiety and, consequently, potential weight loss.31
- Among men consuming a Western diet, adding 75 grams, or just over half a cup, of walnuts daily significantly improved sperm quality. including vitality, motility, and morphology.32

Summary

Unlike other nuts, walnuts are rich in omega-3 polyunsaturated fatty acids including alpha-linoleic acid (ALA)—a substantial **2.7 grams** per quarter cup—with an array of beneficial effects. They also contain oleic acid, the critical monounsaturate found in olive oil. Walnuts provide several very rare and highly potent, anticancer phytonutrients juglone, tellimagrandin, and morin. Studies show walnuts can inhibit cancer^{13,33,34} and promote brain,²³⁻²⁵ cardiovascular,^{19,20,22} microbiomic,30 and metabolic29 health, making them an undeniable super food. •

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

What You **Should Do**

Despite the benefits you've just read about walnuts, it may be unwise to add them to your diet. That's because they have a relatively high calorie content that could induce unwanted weight gain.

What you should do is replace unhealthy components of your diet with walnuts. Just imagine how much healthier the American population would be if they substituted walnuts for the dangerous snacks (such as potato chips and corn chips) that are endlessly advertised on TV?

References

- Flores-Mateo G, Rojas-Rueda D, Basora J, Ros E, Salas-Salvadó J. Nut intake and adiposity: meta-analysis of clinical trials. Am J Clin Nutr. 2013 Jun;97(6):1346-55.
- 2. Ros E. Health benefits of nut consumption. *Nutrients*. 2010 Jul;2(7):652-82.
- Available at: http://www.whfoods.com/ genpage.php?tname=nutrient&dbid=84. Accessed August 19, 2015
- Available at: http://www.whfoods. com/genpage.php?tname= foodspice &dbid=99. Accessed August 19, 2015
- S. Reiter RJ, Manchester LC, Tan DX. Melatonin in walnuts: influence on levels of melatonin and total antioxidant capacity of blood. *Nutrition*. 2005 Sep:21(9):920-4.
- Papoutsi Z, Kassi E, Chinou I, Halabalaki M, Skaltsounis LA, Moutsatsou P. Walnut extract (*Juglans regia* L.) and its component ellagic acid exhibit anti-inflammatory activity in human aorta endothelial cells and osteoblastic activity in the cell line KS483. *Br J Nutr*. 2008;99:715-22.
- 7. Vinson JA, Cai Y. Nuts, especially walnuts, have both antioxidant quantity and efficacy and exhibit significant potential health benefits. *Food Funct*. 2012 Feb;3(2):134-40.
- Fukuda T, Ito H, Yoshida T. Antioxidative polyphenols from walnuts (*Juglans regia* L.). *Phytochemistry*. 2003 Aug;63(7): 795-801.
- 9. Scalbert A, Johnson IT, Saltmarsh M. Polyphenols: antioxidants and beyond. *Am J Clin Nutr*. 2005 Jan;81(1 Suppl):215S-7S.
- 10. Shimoda H, Tanaka J, Kikuchi M, et al. Walnut polyphenols prevent liver damage induced by carbon tetrachloride and d-galactosamine: hepatoprotective hydrolyzable tannins in the kernel pellicles of walnut. *J Agric Food Chem*. 2008 Jun 25;56(12):4444-9.
- 11. Available at: http://www.sciencedaily. com/releases/2010/03/100322153953.htm. Accessed August 19, 2015
- 12. Available at: http://www.webmd.com/breast-cancer/news/20090421/walnuts-fight-breast-cancer. Accessed August 19, 2015
- 13. Kim YS, Choi SW, Min SJ, Lee J, Kim Y. Walnut phenol extracts inhibit stemness of colon cancer stem cells in vitro. *FASEB J.* 2015 April 29:752.10.
- 14. Chen L, Na-Shun BY, Zhang J, Yu J, Gu WW. Effect of juglone on the ultrastructure of human liver cancer BEL-7402 cells. Nan Fang Yi Ke Da Xue Xue Bao. 2009 Jun;29(6):1208-11.
- Le V, Esposito D, Grace MH, et al. Cytotoxic effects of ellagitannins isolated from walnuts in human cancer cells. *Nutr Cancer*. 2014;66(8):1304-14.

- 16. Hyun HB, Lee WS, Go SI, et al. The flavonoid morin from Moraceae induces apoptosis by modulation of Bcl-2 family members and Fas receptor in HCT 116 cells. *Int J Oncol*. 2015 Jun;46(6):2670-8.
- Available at: http://www.webmd.com/ heart/arginine-heart-benefits-and-sideeffects. Accessed August 19, 2015
- Available at: https://umm.edu/health/medical/altmed/supplement/alphalinolenicacid. Accessed August 19, 2015
- 19. Marangoni F, Colombo C, Martiello A, Poli A, Paoletti R, Galli C. Levels of the n-3 fatty acid eicosapentaenoic acid in addition to those of alpha linolenic acid are significantly raised in blood lipids by the intake of four walnuts a day in humans. Nutr Metab Cardiovasc Dis. 2007 Jul;17(6):457-61.
- Guasch-Ferré M1, Bulló M, Martínez-González MÁ, et al. Frequency of nut consumption and mortality risk in the PREDIMED nutrition intervention trial. BMC Med. 2013 Jul 16:11:164.
- 21. Banel DK, Hu FB. Effects of walnut consumption on blood lipids and other cardiovascular risk factors: a meta-analysis and systematic review. *Am J Clin Nutr*. 2009 Jul;90(1):56-63.
- 22. R, Yim SJ, Shearer G, Keen C, Djurica D, Newman J, Hackman R. The change in Holt human microvascular function and its relationship to plasma epoxide content after short-term walnut intake. FASEB J. April 2015 29:923.9.
- 23. Joseph JA, Shukitt-Hale B, Willis LM. Grape juice, berries, and walnuts affect brain aging and behavior. *J Nutr*. 2009 Sep;139(9):1813S-7S.
- 24. Poulose S, Bielinski D, Crott J, Roe A, Thangthaeng N, Shukitt-Hale B. Effects of aging and walnut-rich diet on dna methylation and expression of immediate-early genes in critical brain regions. *FASEB J*. 2015 April 29:749.7.
- Pribis P, Bailey RN, Russell AA, et al. Effects of walnut consumption on cognitive performance in young adults. *Br J Nutr*. 2012 May;107(9):1393-401.
- Chauhan N, Wang KC, Wegiel J, Malik MN. Walnut extract inhibits the fibrillization of amyloid beta-protein, and also defibrillizes its preformed fibrils. *Curr Alzheimer Res.* 2004 Aug;1(3):183-8.
- 27. Muthaiyah B, Essa MM, Lee M, Chauhan V, Kaur K, Chauhan A. Dietary supplementation of walnuts improves memory deficits and learning skills in transgenic mouse model of Alzheimer's disease. *J Alzheimers Dis.* 2014;42(4):1397-405.
- Arab L, Ang A. A cross sectional study of the association between walnut consumption and cognitive function among adult US populations represented in NHANES. J Nutr Health Aging. 2015 Mar;19(3):284-90.

- 29. Tapsell LC, Batterham MJ, Teuss G, et al. Long-term effects of increased dietary polyunsaturated fat from walnuts on metabolic parameters in type II diabetes. *Eur J Clin Nutr.* 2009 Aug;63(8):1008-15.
- 30. Byerley L, Ponder M, Lorenzo B, Banks S, Taylor C, Luo M, Blanchard E, Welsh D. Walnut consumption changes the relative abundance of bacteroidetes and firmicutes in the gut. *FASEB J*. April 2015 29:1006.1.
- 31. Brennan AM, Sweeney LL, Liu X, Mantzoros CS. Walnut consumption increases satiation but has no effect on insulin resistance or the metabolic profile over a 4-day period. *Obesity (Silver Spring)*. 2010 Jun;18(6):1176-82.
- 32. Robbins WA, Xun L, FitzGerald LZ, Esguerra S, Henning SM, Carpenter CL. Walnuts improve semen quality in men consuming a Western-style diet: randomized control dietary intervention trial. *Biol Reprod.* 2012 Oct 25;87(4):101.
- Le V, Esposito D, Grace MH, et al. Cytotoxic effects of ellagitannins isolated from walnuts in human cancer cells. *Nutr Cancer*. 2014;66(8):1304-14.
- 34. Fitschen PJ, Rolfhus KR, Winfrey MR, Allen BK, Manzy M, Maher MA. Cardiovascular effects of consumption of black versus English walnuts. *J Med Food*. 2011 Sep;14(9):890-8.
- 35. Available at: http://nutritiondata.self. com/facts/nut-and-seed-products/3138/2. Accessed August 19, 2015



Welcome LifeExtension Rx Welcome



Lowest Prices

Ask for LifeExtension Rx* Customer Discount Pricing 1-877-877-9700

Providing Trusted Prescription Compounding for Over 40 Years

- *Full Compounding Lab
- *Full Retail Pharmacy
- *Bio-identical Hormone Replacement Therapy
- *Free Standard Delivery/Shipping
- *Durable Medical Equipment
- *Trilingual (English, Spanish, French)
- *Liscensed to ship into 37 States



Proud members of PCCA, IA



PH:877-877-9700 FAX:877-877-9708

Renew Rx's online to receive \$1 off each Rx

(cash RX's only)

4401 Sheridan St. Hollywood, Fl 33021

www.POSTHASTEPHARMACY.com



Preserve Youthful CELLULAR

ENERGY

with Next-Generation

LIPOIC ACID

Published studies have shown the critical importance of **lipoic acid** in supporting healthy mitochondrial function.

Unlike other forms of lipoic acid, **Super R-Lipoic Acid** is more bioavailable, stable, and potent, achieving **10-30 times** higher peak blood levels than pure R-lipoic acid.¹ This unique **sodium-R-lipoate** can help you reach peak plasma concentrations within just 10-20 minutes² of supplementation.

Super R-Lipoic Acid provides more of the active "R" form of lipoic acid than any other supplement.

To order Super R-Lipoic Acid, call 1-800-544-4440 or visit www.LifeExtension.com

Super R-Lipoic Acid

Item# 01208 • 60 vegetarian capsules

| | Retail Price | Your Price |
|-----------|-----------------|---------------|
| 1 bottle | \$49 | \$36.75 |
| 4 bottles | | \$33.75 each |

Suggested dose is one to two capsules daily.

Non-GMO

References

- Carlson DA, Young KL, Fischer SJ, Ulrich H. In: Packer L. Patel M. eds. Lipoic Acid: Energy Production, Antioxidant Activity and Health Effects. London: Taylor & Francis Publishers; 2008:235-70.
- 2. Carlson DA, Smith AR, Fischer SJ, Young KL, Packer L. Altern Med Rev. 2007 Dec;12(4):342-51.

CAUTION: If you are taking glucose lowering medication, consult your healthcare provider before taking this product.

Bio-Enhanced® is a registered trademark of Geronova Research, Inc.

ULTIMATE HYDRATION

Maintaining eye support is essential for optimal eye health.

As we get older, our eyes become vulnerable to a variety of insults that can cause irritation and dry eye. With just a few drops of the proper eye lubricant, **eye irritation** stemming from **dryness** may be alleviated.

Brite Eyes III provides a powerful dose of <u>two</u> well-established lubricants in every drop, soothing eye discomfort without irritation.

Hydroxymethyl-cellulose and glycerin are FDA-approved for ophthalmic use and are uniquely preserved with potent antioxidants and anti-glycating agents.

The **Brite Eyes III** formula is buffered in a way to make it **soothing to the eye**. The suggested use of **Brite Eyes III** is to instill 1 to 2 drops in the affected eye as needed.

Brite Eyes III

Item #00893 • 2 vials (5 mL each)

| | Retail Price | Your Price |
|---------|-----------------|---------------|
| 1 box | \$34 | \$25.50 |
| 4 boxes | | \$24 each |

Brite Eyes III Sterile Lubricant Eve Drops

Manufactured for:
Quality Supplements and Vitamins Inc.
Ft. Lauderdale, Ft. 33309
info@lifeextension.com
www.lef.org
1.866-280-2852

2 Vials (5 mL each)

Each box of **Brite Eyes III** contains two individual vials that provide **5 mL** each. The reason for putting **Brite Eyes** into individual vials is to reduce the risk of bacterial contamination. Having small vials also makes it convenient for consumers to keep **Brite Eyes** readily accessible at home, the office, in one's purse or pocket, and other places where access to a soothing eye drop is needed.

To order Brite Eyes III, call 1-800-544-4440 or visit www.LifeExtension.com



Blood testing provides the ultimate information regarding correctable risk factors that may predispose you to disorders such as cancer, diabetes, cardiovascular disease, and more. Information about general health and nutritional status can also be gained through standard blood analysis. Standing behind the belief that blood testing is an essential component of any program designed to attain optimal health and longevity, *Life Extension*® offers this innovative and convenient service at a very affordable price. Not only is comprehensive blood testing an important step in safeguarding your health, it is a simple process from virtually anywhere in the United States.

Five Easy Steps:

- Call 1-800-208-3444 to discuss and place your order with one of our knowledgeable health advisors.
 (This order form can also be faxed to 1-866-728-1050 or mailed).
 Online orders can also be placed at www.lifeextension.com.
- After your order is placed, you will be mailed either a requisition form to take to your local LabCorp Patient Service Center or a Blood Draw Kit; whichever is applicable (Please note: If a blood draw kit is used, an additional local draw fee may be incurred.)
- 3. Have your blood drawn.
- 4. Your blood test results will be sent directly to you by Life Extension.
- 5. Take the opportunity to discuss the results with one of our knowledgeable health advisors 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.

Bloed Testing The Ultimate Information

MOST POPULAR PANELS

All of the blood test prices you see here are 25% off retail.

COMPREHENSIVE PANELS

MALE LIFE EXTENSION PANEL (LC322582) \$269

Chemistry Profile includes glucose, cholesterol, LDL, HDL, triglycerides, liver and-kidney function tests PLUS 20 additional tests. CBC includes immune (white) cell count, red blood cell count and platelet count. Also includes: C-Reactive Protein

DHEA-S Homocysteine
TSH for thyroid function Free Testosterone
Estradiol Total Testosterone

Vitamin D 25- hydroxy PSA (prostate-specific antigen)
Hemoglobin A1c

FEMALE LIFE EXTENSION PANEL (LC322535)

Chemistry Profile includes glucose, cholesterol, LDL, HDL, triglycerides, liver and-kidney function tests PLUS 20 additional tests. CBC includes immune (white) cell count, red blood cell count and platelet count. Also includes: C-Reactive Protein DHEA-S Homocysteine

TSH for thyroid function Free Testosterone
Estradiol Total Testosterone
Progesterone Vitamin D 25-hydroxy
Hemoglobin A1c

WEIGHT LOSS PANEL-COMPREHENSIVE (LC100028)

CBC/Chemistry profile (see description at right), DHEA-S, free and total Testosterone, Estradiol, Progesterone, Cortisol, TSH, Free T3, Free T4, Reverse T3, Insulin, Hemoglobin A1c, Vitamin D 25-hydroxy, C-Reactive Protein (high sensitivity), and Ferritin.

MALE ELITE PANEL* (LC100016)

Chem/CBC 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, Vitamin D 25-OH, hs-CRP, Ferritin, Homocysteine

FEMALE ELITE PANEL* (LC100017)

Chem/CBC 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, Vitamin D 25-OH, hs-CRP, Ferritin, Homocysteine

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 valuable information about a testosterone metabolite that can affect the prostate, and the mother hormone that acts as a precursor to all other hormones

To provide an even more in-depth analysis of a woman's hormone status, Life Extension has created this panel as an addition to the Female Life Extension Panel. This panel provides valuable information about total estrogen status, and the mother hormone that acts as a precursor to all other hormones.

LIFE EXTENSION THYROID PANEL (LC304131)
TSH, T4, Free T3, Free T4.

FEMALE COMPREHENSIVE HORMONE PANEL* \$2

(LC100011) CBC/Chemistry Profile (see description above right), DHEA-S, Estradiol, Total Estrogens, Progesterone, Pregnenolone, Total and Free Testosterone, SHBG, TSH, Free T3, Free T4, Cortisol.

MALE COMPREHENSIVE HORMONE PANEL* (LC100010) CBC/Chemistry Profile (see description above right), DHEA-S, Estradiol, DHT, PSA, Pregnenolone, Total and Free Testosterone, SHBG, TSH, Free T3, Free T4, Cortisol.

THE CBC/CHEMISTRY PROFILE (LC381822)

Note: This CBC/Chemistry Profile is included in many Life Extension panels. Please check panel descriptions. CARDIOVASCULAR RISK PROFILE

Total Cholesterol Cholesterol/HDL Ratio
HDL Cholesterol Estimated CHD Risk
LDL Cholesterol Glucose
Triglycerides Iron
LIVER FUNCTION PANEL

AST (SGOT)
ALT (SGPT)
Alkaline Phosphatase

KIDNEY FUNCTION PANEL

BUN BUN/Creatinine Ratio
Creatinine Uric Acid

BLOOD PROTEIN LEVELS

Total Protein Globulin

Albumin Albumin/Globulin Ratio BLOOD COUNT/RED AND WHITE BLOOD

CELL PROFILE

Red Blood Cell Count Monocytes White Blood Cell Count Lymphocytes Platelet Count Eosinophils Hemoglobin Basophils Polys (Absolute) Hematocrit Lymphs (Absolute) MCV Monocytes (Absolute) MCH Eos (Absolute) MCHC Baso (Absolute) Polynucleated Cells

RDW
BLOOD MINERAL PANEL

Calcium Sodium Potassium Chloride Phosphorus Iron

COMPREHENSIVE THYROID PANEL (LC100018)

\$575

\$120

TSH, T4, Free T4, Free T3, Reverse T3, TPO, ATA

\$199

\$198

\$125

\$249

\$149

\$90

\$330

FOOD SAFE ALLERGY TEST** (LCM73001)

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

STRESS MANAGEMENT PROFILE (LC100043)

Cortisol AM/PM, DHEA-S, Glucose, Insulin, Progesterone, Free T3, Lipid Panel

HEALTHY AGING PANEL-COMPREHENSIVE* (LC100026)

CBC/Chemistry profile (see description above), C-Reactive Protein (high sensitivity), Vitamin B12, Folate, Homocysteine, Vitamin D 25-hydroxy, Hemoglobin A1c, TSH, Free T3, Free T4, Ferritin, Urinalysis, Fibrinogen, and Insulin.

HEALTHY AGING PANEL-BASIC* (LC100025)

CBC/Chemistry profile (see description above), C-Reactive Protein (high sensitivity), Vitamin B12, Folate, Vitamin D 25-hydroxy, Hemoglobin A1c, TSH, Ferritin, and Insulin.

VAP™ TEST* (LC804500)

The VAPTM cholesterol test provides a more comprehensive coronary heart disease (CHD) risk assessment than the conventional lipid profile. Direct measurements, not estimations, are provided for total cholesterol, LDL, HDL, VLDL, and cholesterol subclasses.

VAP™ PLUS* (LC100009)

VAP, C-Reactive Protein (high sensitivity), Homocysteine, Fibrinogen, PLAC® Test (Lp-PLA2), Vitamin D 25-hydroxy.



Other Popular Tests and Panels

| Other Popular | | | | | | | | | |
|---|---------------------|--|--|--|--|--|--|--|--|
| NUTRIENT PANEL* (LC100024) Vitamin B12, Folate, Vitamin D 25-hydroxy, Vitamin C, Vitamin A, Selenium, Zinc, CoQ10, and RBC Magnesium. | \$349 | | | | | | | | |
| CHRONIC FATIGUE PROFILE (LC100005) CBC/Chemistry Profile (see description previous pa Epstein—Barr Virus antibodies (IgG and IgM), Cytomegalovirus Antibodies (IgG and IgM), Ferritin Total and Free Testosterone, DHEA-S, Free T3, Free Cortisol, C-Reactive Protein (high sensitivity), Vitamin B12, Folate, Insulin. | , | | | | | | | | |
| ANEMIA PANEL* (LC100006) CBC/Chemistry Profile (see description previous pa Ferritin, Total Iron Binding Capacity (TIBC), Vitamin B12, Folate | \$79 ge), | | | | | | | | |
| AUTOIMMUNE DISEASE SCREEN* (L100041) ANA screen, hs-CRP, TNF , Immunoglobulins, IgA, IgG, IgM | \$199 | | | | | | | | |
| DIABETES MANAGEMENT PROFILE – COMPREHENSIVE (LC100040) Hemoglobin A1C, Glucose, Insulin, Lipid Panel, Glycomark | \$129 | | | | | | | | |
| DIABETES MANAGEMENT PROFILE – BASIC (LC100039) Hemoglobin A1C, Glucose, Insulin | \$39 | | | | | | | | |
| ADVANCED CARDIAC BIOMARKERS ADVANCED OXIDIZED LDL PANEL* (LC100035) This panel looks at vascular inflammatory biomarks beginning with lifestyle choices to the development metabolic as well as cardiovascular disease and the formation of vulnerable plaque. The panel contain following tests: F2-Isoprostanes, Myeloperoxidase, Oxidized LDL. | t of ne s the | | | | | | | | |
| OXIDIZED LDL PANEL* (LC100034) This panel looks at vascular inflammatory biomarks beginning with the development of metabolic as we cardiovascular disease and the formation of vulner plaque. The panel contains the following tests: Myeloperoxidase and Oxidized LDL. | ell as | | | | | | | | |
| OXIDIZED LDL* (LC817472) OxLDL is a powerful initiator of inflammatory changes in the artery wall, which eventually lead to the formation of plaque. | \$75 | | | | | | | | |
| | | | | | | | | | |
| LE YOUR HEALTHY REWARDS | | | | | | | | | |
| With Your Healthy Rewards , you earn | 1 | | | | | | | | |

With **Your Healthy Rewards**, you earn **LE Dollars** back on every purchase you make — including blood tests!

See www.LifeExtension.com/Rewards for details.

HORMONES

DHEA-SULFATE (LC004020) This test shows if you are taking the proper amount of DHEA. This test normally costs \$100

MALE BASIC HORMONE PANEL (LC100012)
DHEA-S, Estradiol, Free and Total Testosterone, PSA

or more at commercial laboratories.

FEMALE BASIC HORMONE PANEL (LC100013)
DHEA-S, Estradiol, Free and Total Testosterone,
Progesterone

DIHYDROTESTOSTERONE (DHT)* (LC500142)
Measures serum concentrations of DHT.

\$50

\$33

\$29.90

\$55

\$33

\$47

\$28

\$39.68

ESTRADIOL (LC004515)
For men and women. Determines the proper amount in the body.

INSULIN FASTING (LC004333)
Can predict those at risk of diabetes, obesity, heart and other diseases.

PREGNENOLONE* (LC140707)
Used to determine ovarian failure, hirsutism, adrenal carcinoma, and Cushing's syndrome.

PROGESTERONE (LC004317)
Primarily for women. Determines the proper amount in the body.

SEX HORMONE BINDING GLOBULIN (SHBG)
(LC082016)
This test is used to monitor SHBG levels which are under the positive control of estrogens and

GENERAL HEALTH

VITAMIN D (250H) (LC081950)
This test is used to rule out vitamin D

thyroid hormones, and suppressed by androgens.

deficiency as a cause of bone disease.
It can also be used to identify hypercalcemia.

FERRITIN (LC004598)
Ferritin levels reflect your body's iron stores and is

also a biomarker for insulin resistance.

VITAMIN B12/FOLATE* (LC000810)

Measurements of B12 and Folate help evaluate your general health and nutritional status since the B vitamins are important for cardiac health as well as energy production.

PSA (PROSTATE SPECIFIC ANTIGEN) (LC010322)
Screening test for prostate disorders and possible cancer

Blood tests available in the continental United States only.
Restrictions apply in NY, NJ, PA, RI, and MA.
Not available in Maryland.

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

** This test is packaged as a kit, requiring a finger stick performed at home.

ORDER LIFE SAVING BLOOD TESTS FROM VIRTUALLY ANYWHERE IN THE US!

TERMS AND CONDITIONS

This blood test service is for informational purposes only and no specific medical advice will be provided. National Diagnostics, Inc., and Life Extension® contract with a physician who will order your test(s), but will not diagnose or treat you. Both the physician and the testing laboratory are independent contractors and neither National Diagnostics, Inc., nor Life Extension® will be liable for their acts or omissions. Always seek the advice of a trained health professional for medical advice, diagnosis, or treatment. When you purchase a blood test from Life Extension/National Diagnostics, Inc., you are doing so with the understanding that you are privately paying for these tests. There will be absolutely no billing to Medicare, Medicaid, or private insurance. I have read the above Terms and Conditions and understand and agree to them.

| Signature | | | |
|--------------------------|---|--------|--|
| Χ | | | |
| | | | |
| CUSTOMER NO. | | | |
| Male | | Female | |
| | | | |
| Name | | | |
| Date of Birth (required) | / | / | |
| Address | | | |
| City | | | |
| State | | Zip | |
| Phone | | | |
| Credit Card No. | | | |
| Expiration Date | | / | |

Mail your order form to:

LifeExtension

National Diagnostics, Inc

3600 West Commercial Boulevard Fort Lauderdale, FL 33309

Phone your order to: **1-800-208-3444**Fax your order to: **1-866-728-1050**

^{*} 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.

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
Endothelial Defense™ with Full-Spectrum
Pomegranate™ and CORDIART™
Endothelial Defense™ with GliSODin®
Full-Spectrum Pomegranate™
Natural BP Management
NitroVasc with CORDIART™
Pomegranate Extract Capsules

Bone Health

Bone Restore
Bone Restore with Vitamin K2
Bone Strength Formula with KoAct®
Bone-Up™
Calcium Citrate with Vitamin D
Dr. Strum's Intensive Bone Formula
Strontium Caps

Brain Health

Acetyl-L-Carnitine Acetyl-L-Carnitine Arginate **Blast** Brain Shield® Gastrodin Cognitex® Basics
Cognitex® with Brain Shield® Cognitex® with Pregnenolone & Brain Shield® Cognizin® CDP-Choline Caps DMAE Bitartrate (dimethylaminoethanol) Dopa-Mind[™] Ginkgo Biloba Certified Extract™ Huperzine A Lecithin Granules Migra-Eeze™ Migra-Mag with Brain Shield® Neuro-Mag® Magnesium L-Threonate Neuro-Mag® Magnesium L-Threonate with Calcium and Vitamin D3 Optimized Ashwagandha Extract . Prevagen™ PS (Phosphatidylserine) Caps Super Ginkgo Extract 28/7 Vinpocetine

Cholesterol Management

Advanced Lipid Control
Cho-Less™
CHOL-Support™
Policosanol
Red Yeast Rice
Theaflavins Standardized Extract
Vitamin B3 Niacin Capsules

Digestion Support

Artichoke Leaf Extract
Carnosoothe with PicroProtect™
Digest RC®
Effervescent Vitamin C - Magnesium Crystals
Enhanced Super Digestive Enzymes
Enhanced Super Digestive Enzymes
w/Probiotics
Esophageal Guardian
Extraordinary Enzymes

Fem Dophilus
Ginger Force®
Organic Golden Flax Seed
Pancreatin
Regimint
Tranquil Tract™
TruFiber™
WellBetX PGX plus Mulberry

Energy Management

Adrenal Energy Formula

Asian Energy Boost

D-Ribose Powder
D-Ribose Tablets
Forskolin
Mitochondrial Basics with BioPQQ®
Mitochondrial Energy Optimizer with BioPQQ®
NAD+ Cell Regenerator™
Peak ATP® with GlycoCarn®
PQQ Caps with BioPQQ®
Rhodiola Extract
RiboGen™ French Oak Wood Extract
Triple Action Thyroid

Eye Health

Mega EPA/DHA

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

Mega GLA with Sesame Lignans
OMEGA FOUNDATIONS™ Super Omega-3
EPA/DHA with Sesame Lignans &
Olive Extract
OMEGA FOUNDATIONS™ Super Omega-3
Plus EPA/DHA with Sesame Lignans,
Olive Extract, Krill & Astaxanthin
Organic Golden Flax Seed
Provinal® Purified Omega-7
Vegetarian Sourced DHA

Food

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® Cruciferous Vegetable Soup
Rich Rewards® Decaf Roast
Stevia Sweetener

Glucose Management

CinSulin® with InSea^{2®} and Crominex® 3+ CoffeeGenic® Green Coffee Extract Glycation Protection Formula Mega Benfotiamine Natural Glucose Absorption Control Tri Sugar Shield®

Heart Health

Aspirin (Enteric Coated)
Cardio Peak™ with Standardized Hawthorn and Arjuna
Fibrinogen Resist™ with Nattokinase
Folate & Vitamin B12 Caps
Optimized Carnitine with GlycoCarn®
Super Ubiquinol CoQ10
Super Ubiquinol CoQ10 with BioPQQ®
Super Ubiquinol CoQ10 with Enhanced
Mitochondrial Support™
Super-Absorbable CoQ10 Ubiquinone
with d-Limonene
TMG Powder
TMG Liquid Capsules

Hormone Balance

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

Immune Support

AHCC® Echinacea Extract Enhanced Zinc Lozenges i26 Hyperimmune Egg Immune Modulator with Tinofend® Immune Protect with PARACTIN® Immune Senescence Protection Formula™ Kinoko® Gold AHCC Kyolic® Garlic Formula 102 Kyolic® Garlic Formula 105 Kyolic® Reserve Lactoferrin (apolactoferrin) Caps NK Cell Activator™ Optimized Garlic Optimized Quercetin Peony Immune ProBoost Thymic Protein A Reishi Extract Mushroom Complex Standardized Cistanche Ten Mushroom Formula® Zinc Lozenges

Inflammation Management

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

Joint Support

Arthro-Immune Joint Support
ArthroMax® Advanced with UC-II® & AprèsFlex®
ArthroMax® with Theaflavins & AprèsFlex®
Bio-Collagen with Patented UC-II®
Fast-Acting Joint Formula
Glucosamine/Chondroitin Capsules
Krill Healthy Joint Formula
MSM (Methylsulfonylmethane)

Kidney & Bladder Support

Cran-Max[®] Cranberry Whole Fruit Concentrate Optimized Cran-Max[®] with Ellirose[™] Water-Soluble Pumpkin Seed Extract

Liver Health & Detoxification

Anti-Alcohol Antioxidants with
HepatoProtection Complex
Calcium D-Glucarate
Chlorella
Chlorophyllin
European Milk Thistle
Glutathione, Cysteine & C
HepatoPro
Liver Efficiency Formula
Mega L-Glutathione Capsules
N-Acetyl-L-Cysteine
PectaSol-C**
Silymarin
SODzyme* with GliSODin* & Wolfberry

Longevity & Wellness

AMPK Activator AppleWise Polyphenol Extract Berry Complete Blueberry Extract Blueberry Extract with Pomegranate CR Mimetic Longevity Formula DNA Protection Formula Enhanced Berry Complete with Acai Essential Daily Nutrients Grapeseed Extract with Resveratrol & Pterostilbene Mega Green Tea Extract (decaffeinated) Mega Green Tea Extract (lightly caffeinated) Optimized Fucoidan with Maritech® 926 Optimized Resveratrol Optimized Resveratrol with Nicotinomide . Riboside pTeroPure®

Pycnogenol® French Maritime
Pine Bark Extract Resveratrol with Pterostilbene RNA (Ribonucleic Acid)

Super Alpha-Lipoic Acid Super R-Lipoic Acid

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 Chromium Ultra Copper Iron Protein Plus Magnesium (Citrate) Magnesium Caps Only Trace Minerals Optimized Chromium with Crominex® 3+ Sea-Iodine™ Se-Methyl L-Selenocysteine Super Selenium Complex Vanadyl Sulfate Zinc Caps

Advanced Iodine Complete Solarshield® Sunglasses X-R Shield

Mood & Stress Management

5 HTP I-Theanine Natural Stress Relief SAMe (S-Adenosyl-Methionine)

Multivitamins

Children's Formula Life Extension Mix™ Comprehensive Nutrient Packs ADVANCED Comprenensive Nutrient Packs ADVANCED
Life Extension Mix™ Capsules without Copper
Life Extension Mix™ Capsules
Life Extension Mix™ Powder without Copper
Life Extension Mix™ Powder Life Extension Mix™ Tablets with Extra Niacin Life Extension Mix™ Tablets without Copper Life Extension Mix™ Tablets Once-Daily Health Booster One-Per-Day Tablets

Two-Per-Day Capsules Two-Per-Day Tablets

Personal Care

w/VERISOL®

Anti-Aging Rejuvenating Scalp Serum 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

Hair Suppress Formula Life Extension Toothpaste Sinus Cleanser Venotone Xyliwhite Mouthwash

Pet Care

Cat Mix Dog Mix

Probiotics

Bifido GI Balance BroccoMax® FlorAssist® Heart Health Probiotic FlorAssist® Oral Hygiene FlorAssist® Probiotic FlorAssist® Throat Health Jarro-Dophilus EPS® Theralac® Probiotics
TruFlora® Probiotics

Skin Care

Advanced Anti-Glycation Peptide Serum Advanced Lightening Cream
Advanced Peptide Hand Therapy Advanced Triple Peptide Serum Advanced Under Eye Serum with Stem Cells Amber Self MicroDermAbrasion Anti-Aging Face Oil Anti-Aging Mask Anti-Aging Rejuvenating Face Cream Anti-Glycation Serum with

Blueberry & Pomegranate Extracts Antioxidant Facial Mist Anti-Oxidant Rejuvenating Foot Cream Anti-Oxidant Rejuvenating Foot Scrub Anti-Oxidant Rejuvenating Hand Cream Anti-Redness & Adult Blemish Lotion Bioflavonoid Cream Broccoli Sprout Cream Collagen Boosting Peptide Serum Corrective Clearing Mask DNA Repair Cream Dual-Action MicroDermAbrasion Enhanced FernBlock® with Red Orange Complex Essential Plant Lipids Reparative Serum Face Rejuvenating Anti-Oxidant Cream Fine Line-Less Healing Formula Healing Mask Healing Vitamin K Cream Hyaluronic Facial Moisturizer Hyaluronic Oil-Free Facial Moisturizer Hydrating Anti-Oxidant Facial Mist Hydroderm Lifting & Tightening Complex Lycopene Cream Melatonin Cream Mild Facial Cleanser

Multi Stem Cell Skin Tightening Complex Neck Rejuvenating Anti-Oxidant Cream Pigment Correcting Cream Rejuvenating Serum
Rejuvenex® Body Lotion
RejuveneX® Factor Firming Serum Renewing Eye Cream Resveratrol Anti-Oxidant Serum Skin Lightening Serum Skin Restoring Phytoceramides with Lipowheat® Skin Stem Cell Serum Stem Cell Cream with Alpine Rose Tightening & Firming Neck Cream Ultra Lip Plumper Ultra Rejuvenex® Ultra RejuveNight® Ultra Wrinkle Relaxer Under Eye Refining Serum Under Eye Rescue Cream Vitamin C Serum

Vitamin D Lotion

Youth Serum

Vitamin E-ssential Cream

Sleep

Bioactive Milk Peptides Enhanced Natural Sleep® with Melatonin Enhanced Natural Sleep® without Melatonin Fast-Acting Liquid Melatonin Glycine L-Tryptophan Melatonin Optimized Tryptophan Plus

Sports Performance

Creatine Whey Glutamine Powder

Creatine Capsules

(Vanilla Flavor) DMG (N, N-dimethylglycine) New Zealand Whey Protein Concentrate, (Natural Chocolate and Vanilla Flavor) Pure Plant Protein Tart Cherry Extract Whey Protein Isolate (Chocolate and Vanilla Flavor)

Vitamins

Ascorbyl Palmitate Benfotiamine with Thiamine Beta-Carotene Biotin Buffered Vitamin C Powder Complete B-Complex Daily C+ Fast-C® with Dihydroquercetin Gamma E Tocopherol with Sesame Lignans Gamma E Tocopherol/Tocotrienols High Potency Optimized Folate Inositol Caps Liquid Emulsified Vitamin D3 Liquid Vitamin D3 Low-Dose Vitamin K2 Methylcobalamin MK-7 Natural Vitamin E No Flush Niacin Optimized Folate (L-Methylfolate)
Pantothenic Acid (Vitamin B-5) Pyridoxal 5'-Phosphate Caps Super Absorbable Tocotrienols Super Ascorbate C Capsules Super Ascorbate C Powder
Super K with Advanced K2 Complex Vitamin B12 Vitamin B6 Vitamin C with Dihydroquercetin Vitamin D3 with Sea-Iodine™ Vitamin D3

Vitamins D and K with Sea-Iodine™ **Weight Management**

7-Keto® DHEA Metabolite Advanced Anti-Adipocyte Formula Advanced Natural Appetite Suppress CalReduce Selective Fat Binder DHEA Complete Garcinia HCA
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 Guarana and Sesame Lignans Super CLA Blend with Sesame Lignans Waist-Line Control™

Women's Health

Advanced Natural Sex for Women® 50+ Breast Health Formula Femmenessence MacaPause® Natural Estrogen without Soy Isoflavones Natural Estrogen ProgestaCare® for Women Super-Absorbable Soy Isoflavones Ultra Soy Extract

| YOUR PRICE | | | | | | | | YO | UR PRIC | F |
|--|----------------------|--------------------------|-------------------|------------------------------|--------|--|----------------------|--------------------------|-------------------|------------------------------|
| ITEM No. 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 Total |
| A | Ÿ | Luon | Luon | Luon Q11 lotai | 01214 | BLUEBERRY EXTRACT • 60 veg. caps | | 16.88 | 15.00 | |
| 01524 ACETYL-L-CARNITINE • 500 mg, 100 veg. caps | 34.00 | 25.50 | 22.50 | | 01438 | BLUEBERRY EXTRACT W/ POMEGRANATE • 60 veg. caps | 30.00 | 22.50 | 20.25 | |
| 01525 ACETYL-L-CARNITINE ARGINATE • 100 veg. caps | 59.00 | 44.25 | 38.24 | | 01506 | BONE FORMULA (DR. STRUM'S INTENSIVE) • 300 caps | 56.00 | 42.00 | 37.50 | |
| 01628 ADRENAL ENERGY FORMULA • 60 veg. caps | 24.00 | 18.00 | 16.50 | | 01726 | BONE RESTORE • 120 caps | 22.00 | 16.50 | 14.25 | |
| 01630 ADRENAL ENERGY FORMULA • 120 veg. caps | 46.00 | 34.50 | 31.50 | | 01727 | BONE RESTORE W/VITAMIN K2 • 120 caps | 24.00 | 18.00 | 16.50 | |
| 01828 ADVANCED LIPID CONTROL • 60 veg. caps | 30.00 | 22.50 | 20.25 | | 01725 | BONE STRENGTH FORMULA W/KOACT® • 120 caps | 45.00 | 33.75 | 30.00 | |
| 00681 AHCC® • 500 mg, 30 caps | 59.98 | 44.99 | | | 00313 | BONE-UP® • 240 caps | 28.95 | 21.71 | 20.41 | |
| 29727 AHCC* (KINOKO* GOLD) • 500 mg, 60 veg. caps | 74.95 | 52.47 | | | 01661 | BORON • 3 mg, 100 veg. caps | 5.95 | 4.46 | 3.94 | |
| 00457 ALPHA-LIPOIC ACID W/BIOTIN (Super) • 250 mg, 60 caps | 37.00 | 27.75 | 24.00 | | 00202 | BOSWELLA • 100 caps | 38.00 | 28.50 | 22.50 | |
| 01907 AMPK ACTIVATOR • 90 veg. caps | 48.00 | 36.00 | 33.00 | | 01802 | BRAIN SHIELD® GASTRODIN • 300 mg, 60 veg. caps | 33.00 | 24.75 | 22.50 | |
| 01440 ANTI-ALCOHOL ANTIOXIDANTS W/HEPATOPRO • 100 caps | 26.00 | 19.50 | 17.25 | | 01253 | BRANCHED CHAIN AMINO ACIDS • 90 caps | 19.50 | 14.63 | 12.75 | |
| 01509 ANTI-ADIPOCYTE FORMULA W/MERATRIM® | 39.00 | 29.25 | 27.00 | | 01699 | BREAST HEALTH FORMULA • 60 caps | 34.00 | 25.50 | 22.50 | |
| & INTEGRA LEAN® (Advanced) • 60 veg. caps | | | | | 00893 | BRITE EYES III • 2 vials, 5 ml each | 34.00 | 25.50 | 24.00 | |
| 01625 APPLEWISE POLYPHENOL EXTRACT 600 mg, 30 veg. caps | 21.00 | 15.75 | 14.25 | | 26576 | BROCCO MAX® • 60 veg. caps | 26.95 | 20.21 | | |
| 01039 ARGININE/ORNITHINE • 500/250, 100 caps | 17.99 | 13.49 | | | 01203 | BROMELAIN (Specially-coated) | 21.00 | 15.75 | 14.25 | |
| 00038 ARGININE/ORNITHINE POWDER • 150 grams | | 17.21 | 14.25 | | | 500 mg, 60 enteric coated tablets | | | | |
| 01624 (L)-ARGININE CAPS • 700 mg, 200 veg. caps | 26.50 | 19.88 | 17.44 | | | С | | | | |
| 02004 ARTERIAL PROTECT • 30 veg. caps | | | 33.00 | | | CALCIUM CITRATE W/VITAMIN D • 300 caps | | | 15.94 | |
| 01617 ARTHROMAX® W/THEAFLAVINS & APRÈSFLEX® | | 33.00 | 30.00 | | 01651 | CALCIUM D-GLUCARATE • 200 mg, 60 veg. caps | | | 11.25 | |
| 120 veg. caps | | 00.00 | 00.00 | | ™01823 | CALREDUCE SELECTIVE FAT BINDER 120 mint chewable tablets | 45.00 | 33.75 | 28.50 | |
| 01618 ARTHROMAX® ADVANCED W/UC-II® & APRÈSFLEX® 60 caps | 36.00 | 27.00 | 24.00 | | 01700 | CARDIO PEAK™ w/STANDARDIZED HAWTHORN & ARJUNA 120 veg. caps | 36.00 | 27.00 | 24.00 | |
| 01404 ARTHRO-IMMUNE JOINT SUPPORT • 60 veg. caps | 32.00 | 24.00 | 21.00 | | 00916 | CARNITINE W/GLYCOCARN® (Optimized) • 60 veg. caps | 36.00 | 27.00 | 24.00 | |
| 00919 ARTICHOKE LEAF EXTRACT • 500 mg, 180 veg. caps | 30.00 | 22.50 | 21.00 | | 01532 | L-CARNITINE • 500 mg, 30 veg. caps | 15.00 | 11.25 | 9.90 | |
| 01533 ASCORBYL PALMITATE • 500 mg, 100 veg. caps | 22.50 | 16.88 | 15.00 | | 01258 | CARNOSOOTHE W/PICROPROTECT™ • 60 veg. caps | 29.95 | 22.46 | 20.25 | |
| 00888 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 | | 01687 | CARNOSINE (Super) • 500 mg, 90 veg. caps | 66.00 | 49.50 | 45.00 | |
| 01066 ASPIRIN • 81 mg, 300 enteric coated tablets | 6.00 | 4.50 | 4.00 | | | CAT MIX • 100 grams powder | 14.00 | 10.50 | 8.25 | |
| 01720 ASTAXANTHIN WITH PHOSPHOLIPIDS • 4 mg, 30 softgels | 16.00 | 12.00 | 10.50 | | 01891 | CHILDREN'S FORMULA LIFE EXTENSION MIX™ | 20.00 | 15.00 | 13.50 | |
| В | | | | | | 100 chewable tablets | | | | |
| 00920 BENFOTIAMINE W/ THIAMINE • 100 mg, 120 veg. caps | 19.95 | 14.96 | 13.95 | | 00550 | CHLORELLA • 500 mg, 200 tablets | 23.50 | 17.63 | | |
| 00925 BENFOTIAMINE (Mega) • 250 mg, 120 veg. caps | 30.00 | 22.50 | 20.25 | | 01571 | CHLOROPHYLLIN • 100 mg, 100 veg. caps | 24.00 | 18.00 | 15.00 | |
| 01206 BERRY COMPLETE • 30 veg. caps | 21.00 | 15.75 | 14.00 | | 01359 | CHO-LESS™ • 90 capsules | 35.00 | 26.25 | | |
| 01496 BERRY COMPLETE W/ACAI (Enhanced) • 60 veg. caps | 29.00 | 21.75 | 19.50 | | 01910 | CHOL-SUPPORT™ • 60 liquid veg. caps | 52.00 | 39.00 | 34.50 | |
| 00664 BETA-CAROTENE • 25,000 IU, 100 softgels | 11.25 | 8.44 | | | 01477 | CHROMIUM ULTRA • 100 veg. caps | 24.00 | 18.00 | 15.75 | |
| 01622 BIFIDO GI BALANCE • 60 veg. caps | 20.00 | 15.00 | 13.50 | | 01504 | CHROMIUM W/CROMINEX® 3+ (Optimized) | 9.00 | 6.75 | 6.00 | |
| 01073 BILBERRY EXTRACT • 100 mg, 100 veg. caps | 42.00 | 31.50 | 28.50 | | 04500 | 500 mcg, 60 veg. caps | 00.00 | 00.50 | 05.50 | |
| 01512 BIOACTIVE MILK PEPTIDES • 30 caps | 18.00 | 13.50 | 12.00 | | | CINSULIN® W/INSEA2® AND CROMINEX® 3+- 90 veg. caps | | | 25.50 | |
| 01631 BIO-COLLAGEN W/PATENTED UC-II® • 40 mg, 60 small caps | 36.00 | 27.00 | 24.00 | | | CISTANCHE (Standardized) • 30 veg. caps | | | 12.00 | |
| *01006 BIOSIL™ • 5 mg, 30 veg. caps | 18.95 | 15.16 | | | | CITRIMAX® (Super)- 180 veg. caps | | 30.00 | | 10.75 |
| *01007 BIOSIL™ • 1 fl oz | 31.99 | 25.59 | | | 00818 | CLA BLEND W/SESAME LIGNANS (Super) 1,000 mg, 120 softgels | 36.00 | 27.00 | 24.75 | 19./5 |
| 00102 BIOTIN • 600 mcg, 100 caps | 7.50 | 5.63 | 4.88 | | 00819 | CLA BLEND W/GUARANA & SESAME (Super) | 42.00 | 31.50 | 28.75 | |
| 01709 BLACK CUMIN SEED OIL • 60 softgels | 16.00 | 12.00 | 10.50 | | | 1,000 mg, 120 softgels | | | | |
| 01710 BLACK CUMIN SEED OIL W/BIO-CURCUMIN® • 60 softgels | 32.00 | 24.00 | 22.50 | | 01896 | COGNITEX® W/BRAIN SHIELD® • 90 softgels | 60.00 | 45.00 | 39.00 | 36.00 |
| 01008 BLAST™ • 600 grams of powder | 26.95 | 20.21 | | | 01897 | COGNITEX® W/PREGNENOLONE & BRAIN SHIELD® 90 softgels | 62.00 | 46.50 | 39.75 | 37.50 |
| 70000 BLOOD PRESSURE MONITOR (ACCUFIT™) • med/lg cuff | 79.99 | 49.99 | | | 01421 | COGNITEX® BASICS • 60 softgels | 38 00 | 28.50 | 26.25 | 24.00 |
| 70004 BLOOD PRESSURE MONITOR • Digital wrist cuff | 69.95 | 52.46 | | | | COGNIZIN® CDP CHOLINE CAPS • 250 mg, 60 veg. caps | | 27.00 | | |
| SUBTOTAL OF COLUMN 1 | | | | | | SUBTOTAL OF COLUMN 2 | | | | |
| TODIONE OF COLUMN 1 | | | | | | | | | | |

| | | | Y | UR PRIC | Œ | |
|--------|---|----------------------|-------------------|-------------------|--------------------|-----------|
| ITEM I | No. PRODUCT | Retail Each \$ | 1 Unit Each | 4 Unit Each | 10 Unit Each | QTY Total |
| 01835 | COMPLETE B-COMPLEX • 60 veg. caps | 10.00 | 7.50 | 6.75 | | |
| 01999 | COMPREHENSIVE NUTRIENT PACKS ADVANCED • 30 packs | 90.00 | 67.50 | 61.50 | | |
| 00119 | COPPER CAPSULES • 2 mg, 100 caps | 9.91 | 7.43 | | | |
| 00949 | COQ10 w/d-LIMONENE (Super-absorbable) 50 mg, 60 softgels | 25.00 | 18.75 | 16.50 | 15.00 | |
| 00950 | COQ10 w/d-LIMONENE (Super-absorbable) 100 mg, 100 softgels | 46.00 | 34.50 | 28.00 | 26.25 | |
| 01226 | COQ10 (Super ubiquinol) • 100 mg, 60 softgels | 56.00 | 42.00 | 36.00 | 33.00 | |
| 01733 | COQ10 w/BIOPQQ ® (Super ubiquinol) • 100 mg, 30 softgels | 54.00 | 40.50 | 33.00 | 30.00 | |
| 01426 | COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) 100 mg, 60 softgels | 62.00 | 46.50 | 39.00 | 36.00 | |
| 01425 | COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) - 50 mg, 100 softgels | 58.00 | 43.50 | 34.50 | 31.50 | |
| 01427 | COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) - 50 mg, 30 softgels | 20.00 | 15.00 | 12.00 | | |
| 01431 | COQ10 w/ENH MITOCHONDRIAL SUPPORT™ (Super ubiquinol) - 200 mg, 30 softgels | 62.00 | 46.50 | 39.00 | 36.00 | |
| 00862 | CRAN-MAX® • 500 mg, 60 veg. caps | 17.50 | 13.13 | 11.25 | | |
| 01424 | CRAN-MAX® WITH ELLIROSE™ (Optimized) • 60 veg. caps | 18.00 | 13.50 | 12.00 | | |
| 01529 | CREATINE CAPSULES • 120 veg. caps | 10.95 | 8.21 | 6.94 | | |
| 01746 | CREATINE WHEY GLUTAMINE POWDER • 454 grams (vanilla) | 30.00 | 22.50 | 20.25 | | |
| 01429 | CR MIMETIC LONGEVITY FORMULA • 60 veg. caps | 39.00 | 29.25 | 27.00 | | |
| 00407 | CURCUMIN® (Super bio) • 400 mg, 60 veg. caps | 38.00 | 28.50 | 26.25 | | |
| 01808 | CURCUMIN® W/GINGER & TURMERONES (Advanced bio) 30 softgels | 30.00 | 22.50 | 20.25 | | |
| 01804 | CYTOKINE SUPPRESS™ W/EGCG • 30 veg. caps | 30.00 | 22.50 | 20.25 | | |
| | COSMESIS | | | | | |
| 80157 | ADVANCED ANTI-GLYCATION PEPTIDE SERUM • 1 oz | 53.00 | 39.75 | 34.50 | | |
| 80154 | ADVANCED LIGHTENING CREAM • 1 oz | 65.00 | 48.75 | 42.75 | | |
| 80155 | ADVANCED PEPTIDE HAND THERAPY • 4 oz | 46.00 | 34.50 | 29.25 | | |
| 80152 | ADVANCED TRIPLE PEPTIDE SERUM • 1 oz | 65.00 | 48.75 | 42.75 | | |
| 80140 | ADVANCED UNDER EYE SERUM W/STEM CELLS • .33 oz | 49.00 | 36.75 | 31.50 | | |
| 80139 | AMBER SELF MICRODERMABRASION • 2 oz | 49.00 | 36.75 | 31.50 | | |
| 80158 | ANTI-AGING FACE OIL • 1 oz | 59.00 | 44.25 | 39.00 | | |
| 80118 | ANTI-AGING MASK • 2 oz | 72.00 | 54.00 | 47.52 | | |
| 80151 | ANTI-AGING REJUVENATING FACE CREAM • 2 oz | 65.00 | 48.75 | 42.75 | | |
| 80153 | ANTI-AGING REJUVENATING SCALP SERUM • 2 oz | 46.00 | 34.50 | 29.25 | | |
| 80134 | ANTI-GLYCATION SERUM W/BLUEBERRY & POMEGRANATE EXTRACTS • 1 oz | 33.00 | 24.75 | 23.51 | | |
| 80133 | ANTIOXIDANT FACIAL MIST • 2 oz | 32.00 | 24.00 | 22.80 | | |
| 80127 | ANTIOXIDANT REJUVENATING FOOT CREAM • 2 oz | 45.00 | 33.75 | 32.10 | | |
| 80128 | ANTIOXIDANT REJUVENATING FOOT SCRUB • 2 oz | 59.00 | 44.25 | 38.94 | | |
| 80117 | ANTIOXIDANT REJUVENATING HAND CREAM • 2 oz | 64.00 | 48.00 | 43.12 | | |
| 80105 | ANTI-REDNESS & ADULT BLEMISH LOTION • 1 oz | 74.50 | 55.88 | 49.17 | | |
| 80147 | BIOFLAVONOID CREAM • 1 oz | 46.00 | 34.50 | 29.25 | | |
| 80144 | BROCCOLI SPROUT CREAM • 1 oz | 46.00 | 34.50 | 29.25 | | |
| 80156 | COLLAGEN BOOSTING PEPTIDE SERUM • 1 oz | 59.00 | 44.25 | 39.00 | | |
| 80120 | CORRECTIVE CLEARING MASK • 2 oz | 64.50 | 48.38 | 42.57 | | |
| 80141 | DNA REPAIR CREAM • 1 oz | 49.00 | 36.75 | 31.50 | | |
| 80108 | ESSENTIAL PLANT LIPIDS REPARATIVE SERUM • 1 oz | 74.95 | 56.21 | 49.46 | | |
| | | | | | | |
| | SUBTOTAL OF COLUMN 3 | | | | | |

| | | | YO | UR PRIC | E | | |
|--------|--|----------------------|--------------------------|-------------------|--------------------|-----|-------|
| ITEM N | o. PRODUCT | Retail Each \$ | 1 Unit Each | 4 Unit Each | 10 Unit Each | OTY | Total |
| 80123 | FACE REJUVENATING ANTIOXIDANT CREAM • 2 oz | | 52.13 | 45.87 | | | |
| 80107 | FINE LINE-LESS • 1 oz | 74.50 | 55.88 | 49.17 | | | |
| 80131 | HAIR SUPPRESS FORMULA • 4 oz | 59.00 | 44.25 | 38.94 | | | |
| 80137 | HEALING FORMULA ALL-IN-ONE CREAM • 1 oz | 53.00 | 39.75 | 34.07 | | | |
| 80115 | HEALING MASK • 2 oz | 64.50 | 48.38 | 42.57 | | | |
| 80102 | HEALING VITAMIN K CREAM • 1 oz | 79.50 | 59.63 | 52.47 | | | |
| 80109 | HYALURONIC FACIAL MOISTURIZER • 1 oz | 58.00 | 43.50 | 38.28 | | | |
| 80110 | HYALURONIC OIL-FREE FACIAL MOISTURIZER • 1 oz | 58.00 | 43.50 | 38.28 | | | |
| 80138 | HYDRATING ANTIOXIDANT FACE MIST • 4 oz | 39.95 | 29.96 | 28.50 | | | |
| 80103 | LIFTING & TIGHTENING COMPLEX • 1 oz | 74.50 | 55.88 | 49.17 | | | |
| 80146 | LYCOPENE CREAM • 1 oz | 28.00 | 21.00 | 19.05 | | | |
| 80135 | MELATONIN CREAM • 1 oz | 33.00 | 24.75 | 20.33 | | | |
| 80114 | MILD FACIAL CLEANSER • 8 oz | 59.00 | 44.25 | 38.94 | | | |
| 80159 | MULTI STEM CELL SKIN TIGHTENING COMPLEX • 1 oz | 59.00 | 44.25 | 39.00 | | | |
| 80122 | NECK REJUVENATING ANTIOXIDANT CREAM • 2 oz | 64.00 | 48.00 | 42.24 | | | |
| 80111 | PIGMENT CORRECTING CREAM • 1/2 oz | 74.00 | 55.50 | 48.84 | | | |
| 80106 | REJUVENATING SERUM • 1 oz | 74.50 | 55.88 | 49.17 | | | |
| 80150 | RENEWING EYE CREAM • 1/2 oz | 65.00 | 48.75 | 42.75 | | | |
| 80142 | RESVERATROL ANTI-OXIDANT SERUM • 1 oz | 46.00 | 34.50 | 29.25 | | | |
| 80112 | SKIN LIGHTENING SERUM • 1/2 oz | 85.00 | 63.75 | 56.10 | | | |
| 80130 | SKIN STEM CELL SERUM • 1 oz | 74.00 | 55.50 | 51.75 | | | |
| 80143 | STEM CELL CREAM W/ALPINE ROSE • 1 oz | 66.00 | 49.50 | 43.50 | | | |
| 80148 | TIGHTENING & FIRMING NECK CREAM • 2 oz | 39.00 | 29.25 | 26.25 | | | |
| 80116 | ULTRA LIP PLUMPER • 1/3 oz | 64.00 | 48.00 | 42.24 | | | |
| 80101 | ULTRA WRINKLE RELAXER • 1 oz | 89.95 | 67.46 | 59.82 | | | |
| 80113 | UNDER EYE REFINING SERUM • 1/2 oz | 74.50 | 55.88 | 49.17 | | | |
| 80104 | UNDER EYE RESCUE CREAM • 1/2 oz | 74.50 | 55.88 | 49.17 | | | |
| 80129 | VITAMIN C SERUM • 1 oz | 85.00 | 63.75 | 56.10 | | | |
| 80136 | VITAMIN D LOTION • 4 oz | 36.00 | 27.00 | 25.25 | | | |
| 80145 | VITAMIN E-ESSENTIAL CREAM • 1 oz | 28.00 | 21.00 | 19.50 | | | |
| 80149 | YOUTH SERUM • 1 oz | | 48.75 | 42.75 | | | |
| | D | | | | | | |
| 01912 | DAILY C+ CITRUS FLAVOR • 30 stick packs | 21.00 | 15.75 | 14.25 | | | |
| 00658 | 7-KETO® DHEA METABOLITE • 25 mg, 100 caps | 28.00 | 21.00 | 18.00 | | | |
| 01479 | 7-KETO® DHEA METABOLITE • 100 mg, 60 veg. caps | 40.00 | 30.00 | 27.00 | | | |
| 01640 | DHA (Vegetarian sourced) • 30 veg. softgels | 20.00 | 15.00 | 13.50 | | | |
| 00607 | DHEA • 25 mg, 100 tablets (Dissolve in mouth) | 14.00 | 10.50 | 8.81 | | | |
| 01478 | DHEA COMPLETE • 60 veg. caps | 48.00 | 36.00 | 32.40 | | | |
| 00335 | DHEA • 25 mg, 100 caps | 18.00 | 13.50 | 11.25 | | | |
| 00454 | DHEA • 15 mg, 100 caps | 14.00 | 10.50 | 9.00 | | | |
| 00882 | DHEA • 50 mg, 60 caps | 19.00 | 14.25 | 12.75 | | Í | |
| 01689 | DHEA • 100 mg, 60 veg. caps | 24.00 | 18.00 | 16.50 | | Î | |
| 01358 | DIGEST RC® • 30 tablets | 19.95 | 14.96 | 12.75 | | Î | |
| 02021 | DIGESTIVE ENZYMES (Enhanced Super) • 60 veg. caps | 22.00 | 16.50 | 15.00 | | Í | |
| 02022 | DIGESTIVE ENZYMES w/PROBIOTICS (Enhanced Super) • 60 veg. caps | 28.00 | 21.00 | 18.00 | | Î | |
| | D,L-PHENYLALANINE • 500 mg, 100 veg. caps | 18.75 | 14.06 | 12.00 | | ĺ | |
| | SUBTOTAL OF COLUMN 4 | | | | | | |

| ITEM N | io. PRODUCT | Retail | Y0 | UR PRIC | 10 | | | ITEM | No. PRODUCT | Retail | Y0 | UR PRIO | 10 |
|---------|---|------------|--------------|--------------|----------------|-------|-------|---------|--|------------|--------------|--------------|----------------|
| | | Each \$ | Unit Each | Unit Each | Unit Each (| QTY : | Total | | | Each \$ | Unit Each | Unit Each | Unit Each (|
| 01540 | DMAE BITARTRATE • 150 mg, 200 veg. caps | 18.00 | 13.50 | 11.25 | | | | 00345 | 5 (L-) GLUTAMINE CAPSULES • 500 mg, 100 caps | 14.95 | 11.21 | 10.13 | |
| 00059 | DMG • 125 mg, 60 tablets | 24.80 | 18.60 | 17.02 | | | | 00141 | 1 (L-) GLUTAMINE POWDER • 100 grams | 22.00 | 16.50 | 15.00 | |
| 01570 | DNA PROTECTION FORMULA • 60 veg. caps | 34.00 | 25.50 | 24.00 | | | | 00522 | 2 GLUCOSAMINE/CHONDROITIN CAPSULES • 100 caps | 38.00 | 28.50 | 24.00 | |
| 01831 | DOG MIX • 100 grams powder | 18.00 | 13.50 | 11.25 | | | | 01541 | 1 GLUTATHIONE, CYSTEINE & C • 100 veg. caps | 20.00 | 15.00 | 13.50 | |
| 02006 | DOPA-MIND™ • 60 veg. tabs | 48.00 | 36.00 | 32.00 | | | | 00314 | 4 L-GLUTATHIONE (Mega) • 250 mg, 60 caps | 39.64 | 29.73 | | |
| 00321 | DR. PROCTOR'S ADVANCED HAIR FORMULA • 2 oz | 39.95 | 29.96 | 24.00 | | | | 01987 | 7 GLYCATION PROTECTION FORMULA • 60 veg. caps | 44.00 | 33.00 | 29.25 | |
| 00320 | DR. PROCTOR'S HAIR SHAMPOO • 8 oz | 24.95 | 18.71 | 16.50 | | | | 01669 | 9 GLYCINE • 1,000 mg, 100 veg. caps | 12.00 | 9.00 | 8.10 | |
| 00899 | DUAL-ACTION MICRODERMABRASION ADV. EXFOLIATE • 2.4 oz | 39.95 | 29.96 | 29.21 | | | | 01411 | The state of the s | 36.00 | 27.00 | 25.50 | |
| | E | | | | | | | 01604 | 100 mg, 60 veg. caps 4 GREEN COFFEE EXTRACT COFFEEGENIC® | 22.00 | 16.50 | 15.00 | |
| 01528 | ECHINACEA EXTRACT • 250 mg, 60 veg. caps | 14.35 | 10.76 | 9.38 | | | | 01004 | 200 mg, 90 veg. caps | 22.00 | 10.00 | 10.00 | |
| 01997 | ENDOTHELIAL DEFENSE™ w/FULL-SPECTRUM POMEGRANATE™ AND CORDIART™ • 60 softgels | 68.00 | 51.00 | 46.50 | | | | 01620 | O GREEN COFFEE EXTRACT COFFEEGENIC® 400 mg, 90 veg. caps | 32.00 | 24.00 | 21.00 | |
| 00997 | ENDOTHELIAL DEFENSE™ w/GLISODIN® • 60 veg. caps | 54.00 | 40.50 | 36.00 | | | | 00953 | 3 GREEN TEA EXTRACT (Mega) • lightly caffeinated, 100 veg. caps | 30.00 | 22.50 | 18.00 | |
| 00625 | EPA/DHA (Mega) • 120 softgels | 19.95 | 14.96 | 13.50 | | | | 00954 | 4 GREEN TEA EXTRACT (Mega) • decaffeinated, 100 veg. caps | 30.00 | 22.50 | 18.00 | |
| 01737 | ESOPHAGEAL GUARDIAN (Berry flavor) • 60 chewable tablets | 36.00 | 27.00 | 24.00 | | | | | н | | | | |
| 01042 | EUROPEAN LEG SOLUTION DIOSMIN 95 | 20.00 | 15.00 | 13.50 | | | | 01074 | 4 5 HTP • 100 mg, 60 caps | 27.95 | 20.96 | | |
| 01706 | 600 mg, 30 veg. tabs EXTRAORDINARY ENZYMES • 60 caps | 26.00 | 19.50 | 18.00 | | | | **02002 | HAIR, SKIN & NAIL REJUVENATION FORM W/VERISON® 90 tabs | 32.00 | 24.00 | 22.00 | |
| 01514 | EYE PRESSURE SUPPORT W/MIRTOGENOL $^{\circ}$ • 30 veg. caps | 38.00 | 28.50 | 25.50 | | | | 01738 | B HCA (Garnicia) • 90 veg. caps | 17.00 | 12.75 | 11.25 | |
| | F | | | | | | | 29754 | 4 HCACTIVE™ GARCINIA CAMBOGIA EXTRACT • 90 caps | 30.00 | 22.50 | | |
| 01054 | FACE MASTER® PLATINUM • Facial Toning System | 199.00 | 199.00 | | | | | 01393 | 3 HEPATOPRO • 900 mg, 60 softgels | 50.00 | 37.50 | 34.50 | |
| 00965 | FAST-ACTING JOINT FORMULA • 30 caps | 39.00 | 29.25 | 27.00 | | | | 01527 | 7 HUPERZINE A • 200 mcg, 60 veg. caps | 40.00 | 30.00 | 27.00 | |
| 01717 | FAST-C® W/DIHYDROQUERCETIN • 120 veg. tabs | 26.00 | 19.50 | 18.00 | | | | 00661 | 1 HYDRODERM® • 1 oz | 79.95 | 59.96 | 49.00 | |
| 20053 | FEM DOPHILUS® • 30 caps | 25.95 | 19.46 | | | | | | T. Control of the Con | | | | |
| 20055 | FEM DOPHILUS® • 60 caps | 39.95 | 29.96 | | | | | *01060 | 0 I26 HYPERIMMUNE EGG • 140 grams powder | 54.99 | 46.75 | | |
| 01064 | FEMMENESSENCE MACAPAUSE® • 120 veg. caps | 34.99 | 26.24 | | | | | 01704 | 4 IMMUNE MODULATOR W/TINOFEND® • 60 veg. caps | 17.00 | 12.75 | 11.25 | |
| 01728 | FERNBLOCK® W/RED ORANGE COMPLEX (Enhanced) | 42.00 | 31.50 | 28.50 | | | | 00955 | 5 IMMUNE PROTECT W/PARACTIN® • 30 veg. caps | 29.50 | 22.13 | 19.91 | |
| | 30 veg. caps | | | | | | | 02005 | 5 IMMUNE SENESCENCE PROTECTION FORMULA™ • 60 veg. tabs | 40.00 | 30.00 | 27.00 | |
| 00718 | FIBRINOGEN RESIST™ • 30 veg. caps | 49.00 | 36.75 | 33.00 | | | | 01049 | 9 INNERPOWER™ • 530 grams powder | 42.00 | 31.50 | | |
| | FLAX SEED (Organic golden) • 14 oz | 11.67 | 8.75 | | | | | 01674 | 4 INOSITOL CAPSULES • 1,000 mg, 360 veg. caps | 62.00 | 46.50 | 43.50 | |
| 01821 | FLORASSIST® HEART HEALTH PROBIOTIC • 60 veg. caps | 32.00 | 24.00 | 21.00 | | | | 01292 | 2 INTEGRA-LEAN® AFRICAN MANGO IRVINGIA | 28.00 | 21.00 | 18.00 | |
| 02011 | FLORASSIST® ORAL HYGIENE • 30 lozenges | 20.00 | 15.00 | 13.50 | | | | | 150 mg, 60 veg. caps | | | | |
| 01825 | FLORASSIST® PROBIOTIC • 30 liquid veg. caps | | 24.00 | | | | | 01248 | 8 IODINE COMPLETE (Advanced) • 12.5 mg, 180 tablets | 46.00 | 36.50 | | |
| | FLORASSIST® THROAT HEALTH • 30 lozenges | 20.00 | 15.00 | 13.50 | | | | 01677 | 7 IRON PROTEIN PLUS • 300 mg, 100 caps | 28.00 | 21.00 | 19.50 | |
| 01913 | FOLATE (Optimized) • 5,000 mcg, 30 veg. tablets | 25.00 | 18.75 | 16.50 | | | | 01492 | 2 IRVINGIA W/PHASE 3 TM CALORIE CONTROL COMPLEX (Ontimized African Manno) • 120 year cans | 56.00 | 42.00 | 36.00 | |
| 01939 | FOLATE (Optimized) • 1,000 mcg, 100 veg. tablets | 19.00 | 14.25 | 12.75 | | | | | (Optimized African Mango) • 120 veg. caps J, K, L | | | | |
| 01841 | FOLATE + VITAMIN B12 CAPS • 200 veg. caps | 10.50 | 7.88 | 7.13 | | | | 00056 | | 22.95 | 17.21 | | |
| 01544 | FORSKOLIN • 10 mg, 60 veg. caps | 16.00 | 12.00 | 10.50 | | | | | 9 JARRO-DOPHILUS EPS® • 30 caps | 39.95 | | | |
| 01513 | FUCOIDAN W/MARITECH® 926 (Optimized) • 60 veg. caps | 36.00 | 27.00 | 24.75 | | | | | 4 K W/ADVANCED K2 COMPLEX (Super) • 90 softgels | | 22.50 | 20.25 | |
| | G | | | | | | | 01600 | ` ` ` ` ` · · · | | 24.00 | | |
| 00559 | GAMMA E TOCOPHEROL/TOCOTRIENOLS • 60 softgels | | | 27.75 | | | | | O KRILL OIL • 60 softgels | 33.95 | | 21.70 | |
| 00759 | GAMMA E TOCOPHEROL W/SESAME LIGNANS • 60 softgels | 32.00 | | 21.75 | | | | | 6 KYOLIC® GARLIC FORMULA 102 • 200 veg. caps | 26.45 | | | |
| 01394 | GARLIC (Optimized) • 200 veg. caps | 24.95 | 18.71 | 15.75 | | | | 00310 | | 27.45 | | | |
| **01122 | GINGER FORCE® • 60 liquid caps | 34.95 | 26.21 | | | | | | 9 KYOLIC® RESERVE • 600 mg, 120 caps | | 20.96 | | |
| 01658 | GINKGO BILOBA CERTIFIED EXTRACT™ 120 mg, 365 veg. caps | 46.00 | 34.50 | 31.50 | | | | | 1 LACTOFERRIN • 60 caps | | | 36.00 | |
| 01648 | GINKGO EXTRACT 28/7 (Super) • 120 mg, 100 veg. caps | 29.00 | 21.75 | 19.88 | | | | 00020 | D LECITHIN • 16 oz granules | 18.00 | 13.50 | 12.00 | |
| 00756 | GLA WITH SESAME LIGNANS (Mega) • 60 softgels | 19.50 | 14.63 | 13.50 | | | | 02055 | 5 LIFE EXTENSION MIX™ • 315 tablets | 80.00 | 60.00 | 52.00 | 43.75 |
| | SUBTOTAL OF COLUMN 5 | | | | | | | | SUBTOTAL OF COLUMN 6 | | | | |

| | | | YO | UR PRI | Œ | |
|--------|--|----------------------|--------------------------|--------------------------|------------|-----------|
| ITEM N | o. PRODUCT | Retail Each \$ | 1 Unit Each | 4 Unit Each | 10 Unit | QTY Total |
| 02057 | LIFE EXTENSION MIX™ W/EXTRA NIACIN • 315 tablets | 80.00 | 60.00 | 52.00 | 43.75 | |
| 02054 | LIFE EXTENSION MIX™ • 490 caps | 90.00 | 67.50 | 58.00 | 47.50 | |
| 02056 | LIFE EXTENSION MIX™ POWDER • 14.81 oz | 80.00 | 60.00 | 52.00 | 43.75 | |
| 02065 | LIFE EXTENSION MIX™ • 315 tablets w/o copper | 80.00 | 60.00 | 52.00 | 43.75 | |
| 02064 | LIFE EXTENSION MIX™ • 490 caps w/o copper | 90.00 | 67.50 | 58.00 | 47.50 | |
| 02066 | LIFE EXTENSION MIX™ POWDER • 14.81 oz w/o copper | 80.00 | 60.00 | 52.00 | 43.75 | |
| 01608 | LIVER EFFICIENCY FORMULA • 30 veg. caps | 18.00 | 13.50 | 12.00 | | |
| 01639 | 5-LOX INHIBITOR W/APRÈSFLEX® • 100 mg, 60 veg. caps | 22.00 | 16.50 | 15.00 | | |
| 01678 | L-LYSINE • 620 mg, 100 veg. caps | 9.00 | 6.75 | 6.00 | | |
| 00455 | LYCOPENE (Mega) • 15 mg, 90 softgels | 35.00 | 26.25 | 22.50 | | |
| | M | | | | | |
| 01885 | MACUGUARD® OCULAR SUPPORT • 60 softgels | 22.00 | 16.50 | 14.85 | | |
| 01886 | MACUGUARD® OCULAR SUPPORT w/ASTAXANTHIN 60 softgels | 42.00 | 31.50 | 28.50 | | |
| 01459 | MAGNESIUM CAPS • 500 mg, 100 veg. caps | 12.00 | 9.00 | 7.50 | | |
| 01682 | MAGNESIUM (CITRATE) • 160 mg, 100 veg. caps | 9.00 | 6.75 | 5.63 | | |
| 01908 | MEDITERRANEAN TRIM WITH SINETROL™-XPUR 60 veg. caps | 18.00 | 13.50 | 12.00 | | |
| 01668 | MELATONIN • 300 mcg, 100 veg. caps | 5.75 | 4.31 | 3.75 | | |
| 01083 | MELATONIN • 500 mcg, 200 veg. caps | 18.00 | 13.50 | 12.00 | | |
| 00329 | MELATONIN • 1 mg, 60 caps | 5.00 | 3.75 | 3.47 | | |
| 00330 | MELATONIN • 3 mg, 60 caps | 8.00 | 6.00 | 5.16 | | |
| 00331 | MELATONIN • 10 mg, 60 caps | 28.00 | 21.00 | 18.00 | | |
| 00332 | MELATONIN • 3 mg, 60 veg. lozenges | 8.00 | 6.00 | 5.16 | | |
| 01734 | MELATONIN (Fast-Acting Liquid) • 2 oz (Citrus-Vanilla) | 12.00 | 9.00 | 8.25 | | |
| 01787 | MELATONIN TIMED RELEASE • 300 mcg, 100 veg. tabs | 12.00 | 9.00 | 8.25 | | |
| 01788 | MELATONIN TIMED RELEASE • 750 mcg, 60 veg. tablets | 8.00 | 6.00 | 5.25 | | |
| 01786 | MELATONIN TIMED RELEASE • 3 mg, 60 veg. tabs | 12.00 | 9.00 | 8.25 | | |
| 01536 | METHYLCOBALAMIN • 1 mg, 60 veg. lozenges (vanilla) | 9.95 | 7.46 | 6.00 | | |
| 01537 | METHYLCOBALAMIN • 5 mg, 60 veg. lozenges (vanilla) | 32.00 | 24.00 | 18.75 | 17.25 | |
| 00709 | MIGRA-EEZE™ (Butterbur) • 60 softgels | 29.50 | 22.13 | 19.75 | | |
| | MIGRA-MAG w/BRAIN SHIELD® • 90 veg. caps | 22.00 | 16.50 | 15.00 | | |
| 01522 | MILK THISTLE (European) • 60 veg. caps | 34.00 | 25.50 | 22.50 | | |
| 01822 | MILK THISTLE (European) • 60 softgels | 28.00 | 21.00 | 18.75 | | |
| 01817 | MILK THISTLE (European) • 120 softgels | | 33.00 | 30.00 | | |
| | MIRAFORTE w/STANDARDIZED LIGNANS (Super) • 120 caps | | | 42.00 | | |
| | MITOCHONDRIAL BASICS W/BIOPQQ® • 30 caps | | 33.00 | 30.00 | | |
| | · | | 54.00 | 48.00 | | |
| | MK-7 • 90 mcg, 60 softgels | | 21.00 | 18.75 | | |
| | MSM (Methylsulfonylmethane) • 1,000 mg, 100 caps | | 10.50 | 8.96 | | |
| | N | | | | | |
| 01534 | N-ACETYL-L-CYSTEINE • 600 mg, 60 veg. caps | 14.00 | 10.50 | 10.13 | | |
| | NAD+ CELL REGENERATOR™ • 100 mg, 30 veg. caps | 34.00 | 25.50 | 19.50 | | |
| | NATTOKINASE • 60 softgels | | 19.13 | | | |
| | NATURAL APPETITE SUPPRESS (Advanced) • 60 veg. caps | | 28.50 | 25.50 | | |
| | NATURAL BP MANAGEMENT • 60 tablets | | 33.00 | | | |
| | NATURAL ESTROGEN • 60 veg. tabs | | 28.50 | | | |
| | SUBTOTAL OF COLUMN 7 | | | | | |

| | ORDER ONLINE VISII: WWW.LIIEEXU | | | UR PRIC | E | | |
|--------|---|----------------|------------------|------------------|------------|-----|-------|
| ITEM N | o. PRODUCT | Retail Each | 1 Unit | 4 Unit | 10 Unit | | |
| 01902 | NATURAL ESTROGEN W/O SOY ISOFLAVONES • 30 veg. caps | \$ | Each | Each 21.00 | Each | QTY | Total |
| | NATURAL SEX FOR WOMEN® 50+ (Advanced) • 90 veg. caps | | 44.25 | 34.00 | | | |
| | NATURAL SLEEP® • 60 veg. caps | 13.00 | 9.75 | 7.50 | | | |
| | NATURAL SLEEP® w/ MELATONIN (Enhanced) • 30 caps | 22.00 | | 15.00 | | | |
| | NATURAL SLEEP® W/O MELATONIN (Enhanced) • 30 caps | | 15.00 | 13.50 | | | |
| | NATURAL SLEEP® MELATONIN • 5 mg, 60 veg. caps | 18.00 | 13.50 | 12.00 | | | |
| | NATURAL STRESS RELIEF • 30 veg. caps | | 21.00 | 18.00 | | | |
| | NERVIA® • 60 softgels | | 37.46 | 10.00 | | | |
| 01603 | NEURO-MAG® MAGNESIUM L-THREONATE • 90 veg. caps | | 30.00 | 27.00 | | | |
| 01602 | NEURO-MAG® L-THREONATE W/CALCIUM & VITAMIN D3 | 40.00 | 30.00 | 27.00 | | П | |
| | 225 grams • Lemon flavor | | | | | | |
| 01990 | NITROVASC w/CORDIART™ • 30 veg. caps | 18.00 | 13.50 | 12.00 | | | |
| 01903 | NK CELL ACTIVATOR™ • 30 veg. tablets | 45.00 | 33.75 | 31.50 | | | |
| 00373 | NO-FLUSH NIACIN • 800 mg, 100 caps | 19.00 | 14.25 | 12.75 | | | |
| | 0 | | | | | | |
| 01824 | OLIVE LEAF VASCULAR SUPPORT w/CELERY SEED EXTRACT (Advanced) • 60 veg. caps | 36.00 | 27.00 | 24.00 | | | |
| 01988 | OMEGA-3 PLUS EPA/DHA w/SESAME LIGNANS, OLIVE EXTRACT, KRILL & ASTAXANTHIN • 120 softgels | 45.00 | 33.75 | 31.50 | 24.75 | | |
| 01983 | OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 60 softgels | 18.00 | 13.50 | 12.00 | 9.38 | | |
| 01982 | OMEGA-3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 120 softgels | 32.00 | 24.00 | 21.00 | 17.05 | | |
| 01984 | OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 120 enteric coated softgels | 34.00 | 25.50 | 23.25 | 18.00 | | |
| 01985 | OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 60 enteric coated softgels | 20.00 | 15.00 | 13.50 | 10.50 | | |
| 01986 | OMEGA 3 EPA/DHA w/SESAME LIGNANS & OLIVE EXTRACT (Super) • 240 small softgels | 32.00 | 24.00 | 21.00 | 17.25 | | |
| 01981 | ONCE-DAILY HEALTH BOOSTER • 60 softgels | 52.00 | 39.00 | 36.00 | | | |
| 02001 | ONE-PER-DAY • 60 tablets | 22.00 | 16.50 | 15.00 | | | |
| 01328 | ONLY TRACE MINERALS • 90 veg. caps | 15.00 | 11.25 | 9.38 | | | |
| | P | | | | | | |
| 01789 | PALMETTOGUARD® SAW PALMETTO W/BETA-SITOSTEROL 30 softgels | 15.00 | 11.25 | 10.50 | 9.00 | | |
| 01790 | PALMETTOGUARD® SUPER SAW PALMETTO/ NETTLE ROOT W/BETA-SITOSTEROL • 60 softgels | 28.00 | 21.00 | 19.50 | 18.00 | | |
| 00073 | PANCREATIN • 50 caps | 13.22 | 9.92 | | | | |
| 01323 | PEAK ATP® WITH GLYCOCARN® • 60 veg. caps | 54.00 | 40.50 | 37.50 | | | |
| 00342 | PECTA SOL-C* MODIFIED CITRUS PECTIN • 454 grams powder | 109.95 | 82.46 | | | | |
| 01080 | PECTA SOL-C® MODIFIED CITRUS PECTIN • 270 veg. caps | 79.95 | 59.96 | | | | |
| 01811 | PEONY IMMUNE • 60 veg. caps | 36.00 | 27.00 | 24.00 | | | |
| 00673 | PGX® PLUS MULBERRY (WellBetX®) • 180 veg. caps | 34.95 | 26.21 | | | | |
| 01676 | PHOSPHATIDYLSERINE CAPS • 100 mg, 100 veg. caps | 54.00 | 40.50 | 36.00 | | | |
| 01436 | POLICOSANOL • 10 mg, 60 veg. caps | 20.00 | 15.00 | 11.25 | | | |
| 01423 | POMEGRANATE™ (Full-Spectrum) • 30 softgels | 24.00 | 18.00 | 15.75 | | | |
| 00956 | POMEGRANATE EXTRACT • 30 veg. caps | 19.50 | 14.63 | 13.16 | | | |
| 01797 | POMI-T® • 60 veg. caps | 33.33 | 25.00 | 22.50 | | | |
| 01500 | PQQ CAPS W/BIOPQQ® • 10 mg, 30 veg. caps | 24.00 | 18.00 | 13.50 | 12.00 | | |
| 01647 | PQQ CAPS W/BIOPQQ® • 20 mg, 30 veg. caps | 40.00 | 30.00 | 24.00 | 21.00 | | |
| 00302 | PREGNENOLONE • 50 mg, 100 caps | 26.00 | 19.50 | 16.50 | | | |
| | SUBTOTAL OF COLUMN 8 | | | | | | |

| ITEM N | lo. PRODUCT | Retail | 1 Y0 | UR PRIO | 10 | |
|----------|--|------------|--------------|--------------|-------|----------|
| IILIVIII | 0. 1100001 | Each \$ | Unit Each | Unit Each | Unit | QTY Tota |
| 00700 | PREGNENOLONE • 100 mg, 100 caps | 30.00 | 22.50 | 20.25 | | |
| *01373 | PRELOX® NATURAL SEX FOR MEN® • 60 tablets | 52.00 | 39.00 | 36.00 | | |
| 01576 | PREVAGEN® • 30 caps | 60.00 | 45.00 | | | |
| 01577 | PREVAGEN® ES • 30 caps | 70.00 | 60.00 | | | |
| 00525 | PROBOOST™ THYMIC PROTEIN A • 30 packets | 59.95 | 44.96 | | | |
| 01441 | PROGESTACARE® FOR WOMEN • 4 oz cream | 35.50 | 26.63 | 24.38 | | |
| 01898 | PROSTATE FORMULA (Ultra NAT) • 60 softgels | 38.00 | 28.50 | 26.25 | 24.00 | |
| 01909 | $\textbf{PROSTAPOLLEN}^{\intercal} \textbf{M} \text{ (Triple strength)} \bullet 30 \text{ softgels}$ | 28.00 | 21.00 | 18.75 | | |
| 01742 | PROTEIN-ISOLATE (Whey) Vanilla • 1 lb. powder | 30.00 | 22.50 | 20.25 | | |
| 01743 | PROTEIN-ISOLATE (Whey) Chocolate • 1 lb. powder | 30.00 | 22.50 | 20.25 | | |
| 01770 | PROTEIN CONCENTRATE (New Zealand Whey) Vanilla 520 grams | 30.00 | 22.50 | 19.95 | | |
| 01771 | PROTEIN CONCENTRATE (New Zealand Whey) Chocolate 660 grams | 30.00 | 22.50 | 19.95 | | |
| 01812 | PROVINAL® PURIFIED OMEGA-7 • 30 softgels | 27.00 | 20.25 | 18.00 | | |
| 01508 | PTEROPURE® • 50 mg Pterostilbene 60 veg. caps | 32.00 | 24.00 | 22.50 | | |
| 01209 | PUMPKIN SEED EXTRACT (Water-soluble) • 60 veg. caps | 20.00 | 15.00 | 13.50 | | |
| 01587 | PURE PLANT PROTEIN • Vanilla 540 grams powder | 38.00 | 28.50 | 26.25 | | |
| 01637 | PYCNOGENOL® FRENCH MARITIME PINE BARK EXTRACT 100 mg, 60 veg. caps | 64.00 | 48.00 | 45.00 | | |
| 01217 | PYRIDOXAL 5'-PHOSPHATE • 100 mg, 60 veg. caps | 22.00 | 16.50 | 14.85 | | |
| | Q, R | | | | | |
| 01309 | QUERCETIN (Optimized) • 250 mg, 60 veg. caps | 22.00 | 16.50 | 15.00 | | |
| 01030 | RED YEAST RICE (Bluebonnet) • 600 mg, 60 veg. caps | 16.95 | 13.56 | | | |
| 00605 | REGIMINT • 60 enteric-coated caps | 19.95 | 14.96 | 14.00 | | |
| 01708 | REISHI EXTRACT MUSHROOM COMPLEX • 60 veg. caps | 30.00 | 22.50 | 20.25 | | |
| 01448 | REJUVENEX® BODY LOTION • 6 oz | 24.00 | 18.00 | 14.85 | 12.75 | |
| 01621 | REJUVENEX® FACTOR FIRMING SERUM • 1.7 oz | 65.00 | 48.75 | 37.50 | | |
| 01220 | REJUVENEX® (Ultra) • 2 oz | 52.00 | 39.00 | 33.00 | 29.25 | |
| 00676 | REJUVENIGHT® (Ultra) • 2 oz | 39.95 | 29.96 | 27.00 | | |
| 01410 | RESVERATROL W/PTEROSTILBENE • 100 mg, 60 veg. caps | 36.00 | 27.00 | 24.00 | | |
| 02031 | RESVERATROL W/NICOTINAMIDE RIBOSIDE (Optimized) • 30 veg. caps | 42.00 | 31.50 | 27.00 | | L |
| 02030 | RESVERATROL (Optimized) • 60 veg. caps | 46.00 | 34.50 | 31.00 | | |
| 00889 | RHODIOLA EXTRACT • 250 mg, 60 veg. caps | 14.00 | 10.50 | 9.00 | | |
| 01900 | RIBOGEN™ FRENCH OAK WOOD EXTRACT 200 mg, 30 veg. caps | 36.00 | 27.00 | 24.75 | | |
| 00972 | (D) RIBOSE POWDER • 150 grams | 27.50 | 20.63 | 18.56 | | |
| 01473 | (D) RIBOSE TABLETS • 100 veg. tabs | 32.00 | 24.00 | 21.00 | | |
| 01609 | RICH REWARDS® BREAKFAST GROUND COFFEE • 12 oz. bag | 13.00 | 9.75 | | | |
| 01730 | RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Mocha • 12 oz. bag | 15.00 | 11.25 | 10.50 | | |
| | RICH REWARDS® BREAKFAST BLEND GROUND COFFEE Natural Vanilla • 12 oz. bag | | 11.25 | 10.50 | | |
| | RICH REWARDS® BREAKFAST BLEND WHOLE BEAN COFFEE 12 oz. bag | | | | | |
| | RICH REWARDST® DECAFFEINATED ROAST GROUND COFFEE 12 oz. bag | | | | | |
| 01530 | RICH REWARDS® CRUCIFEROUS VEGETABLE SOUP • 32 oz. | 11.95 | 8.96 | 8.44 | | |
| | SUBTOTAL OF COLUMN 9 | | | | | |

| | | | YO | UR PRIC | E | | |
|--------|---|----------------------|--------------------------|--------------------------|--------------------|-----|--------|
| ITEM N | o. PRODUCT | Retail Each \$ | 1 Unit Each | 4 Unit Each | 10 Unit Each | QTY | Total |
| 01208 | R-LIPOIC ACID (Super) • 240 mg, 60 veg. caps | 49.00 | 36.75 | 33.75 | | | |
| 00070 | RNA CAPSULES • 500 mg, 100 caps | 17.95 | 13.46 | 12.12 | | | |
| | S | | | | | | |
| 01432 | SAFFRON W/SATIEREAL® (Optimized) • 60 veg. caps | 36.00 | 27.00 | 24.00 | | | |
| 00358 | SAMe (S-ADENOSYL-METHIONINE) 200 mg, 20 enteric coated tablets | 16.00 | 12.00 | 10.50 | 9.75 | | |
| 00557 | SAMe (S-ADENOSYL-METHIONINE) 400 mg, 20 enteric coated tablets | 28.00 | 21.00 | 18.00 | | | |
| 01740 | SEA-IODINE™ • 1,000 mcg, 60 veg. caps | 8.00 | 6.00 | 5.40 | | | |
| 00046 | SELENIUM • 2 fl. oz dropper | 11.95 | 8.96 | | | | |
| 01679 | SE-METHYL L-SELENOCYSTEINE • 200 mcg, 100 veg. caps | 12.00 | 9.00 | 8.25 | | | |
| 00318 | SERRAFLAZYME • 100 tablets | 18.00 | 13.50 | 12.00 | | | |
| 01684 | SILYMARIN • 100 mg, 50 veg. caps | 9.25 | 6.94 | 6.19 | | | |
| 01249 | SINUS CLEANSER • 4 oz. bottle | 25.00 | 18.75 | | | | |
| 01596 | SKIN RESTORING PHYTOCERAMIDES w/LIPOWHEAT® 30 veg. liquid caps | 25.00 | 18.75 | 17.25 | | | |
| 00961 | SODZYME® w/GLISODIN® & WOLFBERRY • 90 veg. caps | 28.00 | 21.00 | 18.00 | | Ī | |
| 00657 | | 12.99 | 9.74 | 8.63 | | | \neg |
| 01097 | SOY EXTRACT (Ultra) • 150 veg. caps | 87.00 | 65.25 | 58.50 | | | |
| 00432 | , , , | 9.95 | 7.46 | | | | |
| | STEVIATM ORGANIC LIQUID SWEETENER (Better) • 2 oz | 11.00 | 8.25 | | | | |
| | STRONTIUM • 750 mg, 90 veg. caps | | | 13.50 | | | |
| | SUPER ABSORBABLE SOY ISOFLAVONES • 60 veg. caps | 28.00 | 21.00 | | | | |
| | SUPER SELENIUM COMPLEX • 200 mcg, 100 veg. caps | | 10.50 | 9.00 | 8.25 | | |
| 01770 | T | 14.00 | 10.50 | 3.00 | 0.23 | | |
| 01723 | TART CHERRY EXTRACT W/STANDARDIZED CHERRYPURE® 60 veg. caps | 22.00 | 16.50 | 15.00 | | | |
| 01827 | TAURINE • 1,000 mg, 90 veg. caps | 13.00 | 9.75 | 9.00 | | Ī | |
| 01918 | TEAR SUPPORT w/MAQUIBRIGHT® • 60 mg, 30 veg. caps | 18.00 | 13.50 | 12.00 | | Ī | |
| 00133 | L-TAURINE POWDER • 300 grams | 20.00 | 15.00 | 12.66 | | Ī | |
| | TEN MUSHROOM FORMULA® • 120 veg. caps | 39.95 | 29.96 | | | | \neg |
| | THEAFLAVIN STANDARDIZED EXTRACT • 30 veg. caps | 18.00 | 13.50 | 12.00 | | | |
| | (L) THEANINE • 100 mg, 60 veg. caps | | 18.00 | 15.38 | | | |
| | THERALAC® PROBIOTICS • 30 caps | | 35.96 | 10.00 | | | |
| | THYROID FORMULA (Metabolic Advantage™) • 100 caps | | 16.46 | | | | |
| | TMG POWDER • 50 grams | | 10.50 | 8.25 | | | |
| | TMG • 500 mg, 60 liquid veg. caps | 13.00 | 9.75 | 9.00 | | | |
| | TOCOTRIENOLS (Super-absorbable) • 60 softgels | | | 21.00 | | | |
| | TOOTHPASTE • 4 oz (Mint) tube | 9.50 | 7.13 | 6.50 | | | |
| | , , | | | 34.50 | | | |
| | TRANQUIL TRACT™ • 60 veg. caps TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT | | | | | | |
| | 60 veg. caps | | | 16.50 | | | |
| 01469 | TRIPLE ACTION CRUCIFEROUS VEGETABLE EXTRACT w/RESVERATROL • 60 veg. caps | 32.00 | | | | | |
| | TRIPLE ACTION THYROID • 60 veg. caps | | | 24.00 | | | _ |
| | TRI SUGAR SHIELD® • 60 veg. caps | | | 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 | | | |
| | SUBTOTAL OF COLUMN 10 | | | | | | |

| | 10 ORDER CALL: 1.954.766.64 | | | | | | . • |
|---------|---|------------|--------------|--------------|--------------|-----|-------|
| ITEM N | o. PRODUCT | Retail | 1 | UR PRIO | 10 | | |
| | | Each \$ | Unit Each | Unit Each | Unit Each | QTY | Total |
| 01721 | TRYPTOPHAN PLUS (Optimized) • 90 veg. caps | 32.00 | 24.00 | 21.75 | | | |
| 02016 | TWO-PER-DAY • 60 tablets | 10.50 | 7.88 | 7.13 | | | |
| 02015 | TWO-PER-DAY • 120 tablets | 20.00 | 15.00 | 13.50 | | | |
| 02014 | TWO-PER-DAY • 120 caps | 22.00 | 16.50 | 15.00 | | | |
| 00326 | L-TYROSINE • 500 mg, 100 tablets | 12.98 | 9.74 | | | | |
| | V | | | | | | |
| 00213 | VANADYL SULFATE • 7.5 mg, 100 veg. tablets | 15.00 | 11.25 | 9.38 | | | |
| 00408 | VENOTONE • 60 caps | 18.95 | 14.21 | 12.00 | | | |
| 01327 | VINPOCETINE • 10 mg, 100 veg. tablets | 18.00 | 13.50 | 10.50 | | | |
| 00372 | VITAMIN B3 NIACIN • 500 mg, 100 caps | 7.65 | 5.74 | 4.99 | | | |
| 00098 | VITAMIN B5 • 500 mg, 100 caps (Pantothenic Acid) | 10.50 | 7.88 | 7.04 | | | |
| 01535 | VITAMIN B6 • 250 mg, 100 veg. caps | 12.50 | 9.38 | 8.25 | | | |
| 00361 | VITAMIN B12 • 500 mcg, 100 lozenges | 8.75 | 6.56 | 5.44 | | | |
| 01634 | VITAMIN C w/DIHYDROQUERCETIN 1,000 mg, 60 veg. tablets | 10.00 | 7.50 | 6.75 | | | |
| 00927 | VITAMIN C w/DIHYDROQUERCETIN 1,000 mg, 250 veg. tablets | 25.50 | 19.13 | 17.44 | | | |
| 00084 | VITAMIN C POWDER (BUFFERED) • 454 grams | 23.95 | 17.96 | 16.50 | | | |
| 01736 | VITAMIN C-MAGNESIUM CRYSTALS (EFFERVESCENT) 180 grams | 20.00 | 15.00 | 13.50 | | | |
| 01732 | VITAMIN D3 • 2,000 IU, 1 fl oz, Mint flavor | 28.00 | 21.00 | 18.75 | | | |
| 01753 | VITAMIN D3 • 1,000 IU, 90 softgels | 7.00 | 5.25 | 4.50 | | | |
| 01751 | VITAMIN D3 • 1,000 IU, 250 softgels | 12.50 | 9.38 | 8.44 | | | |
| 01713 | VITAMIN D3 • 5,000 IU, 60 softgels | 11.00 | 8.25 | 7.43 | | | |
| 01718 | VITAMIN D3 • 7,000 IU, 60 softgels | 14.00 | 10.50 | 9.45 | | | |
| 01758 | VITAMIN D3 W/SEA-IODINE™ • 5,000 IU, 60 caps | 14.00 | 10.50 | 9.38 | | | |
| 00864 | VITAMIN D3 LIQUID EMULSION • 2,000 IU, 1 oz. | 28.00 | 21.00 | 18.75 | | | |
| 01840 | VITAMINS D AND K W/SEA-IODINE™ • 60 caps | 24.00 | 18.00 | 16.50 | | | |
| 01763 | VITAMIN E (Natural) • 400 IU, 100 softgels | 30.00 | 22.50 | 21.00 | 19.50 | | |
| 01225 | VITAMIN K2 (Low-dose) • 45 mcg, 90 softgels | 18.00 | 13.50 | 12.00 | | | |
| | w | | | | | | |
| 01902 | WAIST-LINE CONTROL™ • 120 veg. caps | 42.00 | 31.50 | 28.50 | | | |
| | X, Y | | | | | | |
| 01919 | X-R SHIELD • 90 veg. caps | 15.00 | 11.25 | 9.75 | | | |
| 00409 | XYLIWHITE™ MOUTHWASH • 16 oz | 10.00 | 7.50 | | | | |
| | Z | | | | | | |
| 01813 | ZINC HIGH POTENCY • 50 mg, 90 veg. caps | 7.95 | 5.96 | 5.25 | | | |
| | 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 softgels | 64.95 | 48.71 | | | | |
| 33880 | OUTSTANDING HEALTH: THE 6 ESSENTIAL KEYS | 24.95 | 18 71 | | | | |
| 33000 | TO MAXIMIZE YOUR ENERGY AND WELL BEING by Michael Galitzer, MD & Larry Trivieri Jr. • 2015 | 24.33 | 10.71 | | | | |
| 33878 | TESTOSTERONE REPLACEMENT THERAPY by Dr. John Crisler • 2015 | 19.99 | 14.99 | | | | |
| 33877 | THE TRUTH ABOUT MEN AND SEX by Abraham Morgentaler, MD, FACS • 2015 | 16.99 | 12.74 | | | | |
| 33876 | TOX-SICK • by Suzanne Somers • 2015 | 26.00 | 19.50 | | | | |
| 33875 | DOCTORED: THE DISILLUSIONMENT OF AN AMERICAN PHYSICIAN • by Sandeep Jauhar • 2015 | 26.00 | 19.50 | | | | |
| | SUBTOTAL OF COLUMN 11 | | | | | | |

| | | | Y0 | UR PRIC | E | _ | |
|--------|---|----------------------|-------------------|-------------------|--------------------|-----|-------|
| 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 | | | | |
| 33872 | GET SERIOUS • by Brett Osborn, MD • 2014 | 24.95 | 18.71 | | | | |
| 33868 | TOXIN TOXOUT: GETTING HARMFUL CHEMICAL OUT OF OUR BODIES AND OUR WORLD • by Bruce Lourie and Rick Smith • 2014 | 25.99 | 19.49 | | | | |
| 33867 | THE COMPLETE MEDITERRANEAN DIET by Michael Ozner, MD • 2014 | 19.95 | 14.96 | | | | |
| 33869 | UNLEASH THE POWER OF THE FEMALE BRAIN by Daniel Amen, MD • 2014 | 16.00 | 12.00 | | | | |
| 33870 | MAGNIFICENT MAGNESIUM by Dennis Goodman, MD • 2014 | 14.95 | 11.21 | | | | |
| 33864 | THE SUPPLEMENT PYRAMID by Michael A. Smith, MD • 2014 | 24.95 | 18.71 | | | | |
| 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 | | | |
| 33854 | THE GREAT CHOLESTEROL MYTH • by Jonny Bowden, PhD, CNS and Stephen Sinatra, MD • 2012 | 19.99 | 14.99 | | | | |
| 33958 | THE VITAMIN D SOLUTION by Michael F. Holick, PhD, MD (Paperback) • 2013 | 16.00 | 12.00 | | | | |
| 33838 | YOUR GUIDE TO HEALTHY SKIN THE NATURAL WAY by Gary Goldfaden, MD • 2012 | 26.00 | 15.00 | | | | |
| 33815 | KNOCKOUT • by Suzanne Somers • 2009 | 25.99 | 17.00 | | | | |
| 33809 | TESTOSTERONE FOR LIFE by Abraham Morgentaler, MD • 2008 | 16.95 | 11.87 | | | | |
| 33696 | LIFE EXTENSION REVOLUTION by Philip Lee Miller, MD (Paperback) | 16.00 | 12.00 | | | | |
| 33805 | MIAMI MEDITERRANEAN DIET WITH 300 RECIPES by Michael D. Ozner, MD, FACC, FAHA (Hardcover) • 2008 | 24.95 | 16.25 | | | | |
| 33906 | THE MIGRAINE CURE • by Sergey Dzugan, MD, PhD • 2006 | 24.00 | 15.60 | | | | |
| | 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.



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** \$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. SHIPPING ALL 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

INTRODUCING



Because You Deserve More

Introducing Your Healthy Rewards—the exciting new program exclusively for Life Extension customers. Your Healthy Rewards earns you 2% LE Dollars back on every purchase you make*...and the best part is, Your Healthy Rewards is FREE—no membership involved, no commitment required. The reason behind Your Healthy Rewards is simple: we believe that you deserve more. (Current Life Extension members earn 4%.)

And earn even more benefits when you upgrade to Your Healthy Rewards *Premier*!

For \$49.95, you get an immediate \$50 LE Dollar enrollment bonus, double LE Dollars (4%) back on purchases, complimentary CHOICE unlimited standard shipping service[†], and more. Your Healthy Rewards *Premier* is the ultimate way to earn LE Dollars and enjoy exclusive *Premier*-only perks! At the annual rate of just \$49.95 US/\$59.95 International, *Premier* pays for itself.

Learn more about YOUR HEALTHY REWARDS Call toll-free 1-888-224-8239 www.LifeExtension.com/Rewards

- * You earn LE Dollars on all your Life Extension purchases (except shipping fees, CHOICE and *Premier* program fees, *Life Extension* Magazine® subscriptions, or any purchases made with LE Dollars or gift card). Redeem LE Dollars for any purchase such as products, labs, sale items, and shipping fees at the rate of 1 LE Dollar being equal to \$1 U.S Dollar at checkout. LE Dollars may not be redeemed for *Premier* program fees, CHOICE program fees, *Life Extension* Magazine® subscriptions, or to purchase Gift Cards. LE Dollars have no cash value and are not redeemable for cash, transferable, or assignable for any reason.
- † CHOICE Standard pre-paid shipping offers unlimited shipping to any mailing address within the 50 U.S. states, excluding U.S. territories. CHOICE also gives you discounts on non-standard shipping, shipping outside of the United States, and expedited shipping costs. CHOICE pre-paid unlimited shipping excludes blood test products and gift cards. This offer is not available to international customers serviced by distributors of Life Extension products.

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 | |



You care about the quality of your family's supplements.

We do too.



Master Supplements Inc. is proud that we have been **posting online test results** for every single lot of our products **for over ten years.** These tests verify the purity, potency and high quality of our powerful product line. We rely on ingredients that are backed with scientific research, clinical studies and years of efficacy. We have protected our unique technology with **16 U.S. patents**, providing formulas that support digestive and immune health. Our probiotics, fiber and enzyme supplements all help **restore digestive comfort, regularity, and energy***.

Call your *Life Extension®* advisor to learn more.

Call Life Extension to place your order today.

1-800-544-4440

*These statements have not been evaluated by the Food and Drug Administration.

These products are not intended to diagnose, treat, cure or prevent any disease.







Bio-Cleansing Probiotic



Soluble Fiber with Enzymes

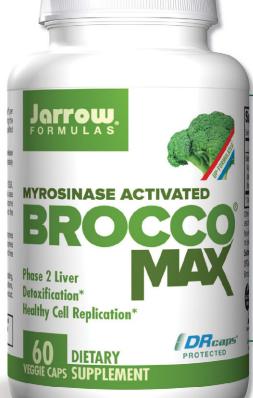
30 capsules Item# 01038 Retail: **\$47.95** Your Price: **\$35.96** 32 capsules Item# 01389 Retail: **\$42.95** Your Price: **\$32.21** 6.2 OZ Item# 01386 Retail: **\$32.95** Your Price: **\$24.71**





BROCCOMAX THE SULFORAPH GENERAI





BroccoMax[®] is protected by U.S. Patent # 9,017,666

- Patented, Ultra Clean, Super Critical Process
- 30 mg of SGS (Sulforaphane Glucosinolate, aka Glucoraphanin) Per Capsule
- Myrosinase Enzyme Activated
- DRcaps® Release in Small Intestine to Ensure the Conversion of SGS to Sulforaphane (and Not Sulforaphane Nitrile)
- Minimum 8 mg Sulforaphane Potential Per Capsule (in vitro test)

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

BroccoMax® Item # 26576 Retail Price \$26.95 Your Price \$20.21



trademark of Capsugel®



VISIT US AT WWW.JARROW.COM FOR MORE PRODUCT INFORMATION

*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.

Fortify Your Immune System with AHCC®

A Clinically Proven and Patented Medicinal Mushroom Extract

Every year, 23 million days of work are lost to feeling under the weather. While most people view immune challenges as part and parcel of the cold weather season, they are not, in fact, inevitable. After all, many people manage to stay well all year round, even though they are exposed to the same environments as those who have weaker resistance. The reason is because the environment is not responsible for whether you feel well or not. Your immune system is.

Innate Versus Adaptive Immunity

You have two basic types of immunity: innate and adaptive. Your innate immunity launches an immediate, general attack against a threat. Your adaptive immunity takes longer to kick in, but produces a targeted, specific response to a threat. Very few natural compounds have the ability to augment both innate and adaptive immunity. AHCC® (short for Active Hexose Correlated Compound) is one exception.*



877-937-2422 | www.qualityoflife.net

f Facebook.com/QualityofLifeLabs

☑ @QOLsupplements



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



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

How AHCC® Works

AHCC® is a patented, fermented, medicinal mushroom extract whose efficacy is supported by over 20 human clinical research studies. It has been shown to modulate immune response in several ways.

- AHCC® enhances the production of cytokines, the messengers of the immune system, so that your whole immune team can coordinate an organized response to outside threats.*
- AHCC® boosts populations of macrophages, the "street cleaners" of your substances and cellular debris.*
- AHCC® increases the activity of natural killer (NK) cells, your innate immune system's first line of defense against invasion.*
- And AHCC® raises levels of dendritic cells and T cells, key players in your adaptive immune system's highly specialized response to specific threats.*

Item# 29727 Retail \$74.95

Your Price \$52.47



Standardized TART CHERRY Extract

STRONG SUPPORT FOR **SORE MUSCLES**

After Exercise and Everyday Exertion

Tart cherries are chockfull of compounds found to block COX-1 and COX-2 inflammatory enzymes. Benefits of the fruit include: 2-5

- Rapid muscle recovery after exercise, and
- · Fast relief from the minor aches, discomfort, and stiffness following everyday activities.

Clinically Validated

Numerous studies have confirmed the muscle-supporting benefits of tart cherries. One clinical trial found that consuming tart cherry juice twice daily produced a substantial decrease in muscle symptoms related to exercise. In the same study, loss of strength due to exercise over four days was reduced from 22% to only 4%.4

Anthocyanins

Anthocyanins—the powerful flavonoids found in dark-pigmented fruit—have been studied for their many advantages, including heart, cellular, and cognitive health. 6-8 Tart cherries have a higher content of anthocyanins than many other fruits.1

Life Extension® offers 100% natural Tart Cherry Extract with Standardized CherryPURE®, which opens the door to the positive benefits of continuous exercise—at any age! This formulation provides all the muscle-supporting benefits of tart cherries and matches that anthocyanin dose used in successful clinical trials by providing a standardized **40 mg** dose of anthocyanins in each vegetarian capsule.2,4



CherryPURE® is a registered trademark of Shoreline Fruit, LLC.

References

- 1. Phytomedicine. 2001 Sep;8(5):362-9.
- 2. J Int Soc Sports Nutr. 2010 May 7;7:17.
- 3. Scand J Med Sci Sports. 2010 Dec;20(6):843-52.
- 4. Br J Sports Med. 2006 Aug;40(8):679-83.
- 5. Am J Vet Res. 2009 Jun;70(6):758-63.
- 6. Mol Nutr Food Res. 2007 Jun;51(6):675-83.
- 7. Biochemistry (Mosc). 2004 Jan;69(1):75-80.

8. Adv Nutr. 2011 Jan;2(1):1-7.



Tart Cherry Extract with Standardized CherryPURE®

Item #01723 • 60 vegetarian capsules

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



To order Tart Cherry Extract with Standardized CherryPURE®, call 1-800-544-4440 or visit www.LifeExtension.com

Dual-Action Support For Aging Joints!

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

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

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

Better Absorption For Optimum Benefit

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

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

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

ArthroMax® Advanced With UC-II® And AprèsFlex®

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

ArthroMax® Advanced with UC-II® & AprèsFlex®

Item #01618 • 60 capsules

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

Non-GMO

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

References

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





PO BOX 407198 FORT LAUDERDALE, FLORIDA 33340-7198

WHAT'S INSIDE

Visit us at www.LifeExtension.com

LifeExtension°

Magazine



7 CONSUMER REPORTS RECOGNIZES HAZARDS OF CT SCANS

Life Extension® long ago cautioned against unnecessary **X-ray** exposure, but these early warnings were largely ignored. **Consumer Reports** magazine has published their own expose on **radiation-emitting** imaging procedures and the staggering number of excess **cancers** being caused.



24 REDUCE RISK OF SORE THROAT

A novel **probiotic** lozenge has been found to markedly lower **strep** and other **throat infections**. Controlled studies show significant **62%-84%** reductions in strep throat incidences in humans.



34 MITIGATE IMPACT OF POST-MEAL GLUCOSE

Blood sugar levels surge after most meals. A fat-soluble form of vitamin B1 called **benfotiamine** can impede *glycation* reactions, helping to guard against this destructive mechanism of elevated blood sugar.



58 PROTECT AGAINST ENVIRONMENTAL TOXINS

Every year, over **4 billion pounds** of toxic chemicals are released into the environment, many of which are known carcinogens. **Chlorophyllin**, a watersoluble form of **chlorophyll** has been shown to neutralize toxic compounds and protect cellular DNA against mutations that can lead to cancer.



70 2015 AMERICAN SOCIETY FOR NUTRITION CONFERENCE

Exclusive report on a conference where experts describe the effects of the ketogenic diet, dietary sugar, dietary starch, dietary fat, weight-loss strategies, muscle loss in the elderly, and more.



97 BENEFITS OF WALNUTS

Walnuts are rich in polyunsaturated fats and oleic acid. Research shows regular ingestion of just small quantities of walnuts can protect against cancers, cardiovascular disease, cognitive decline, and metabolic disorders.