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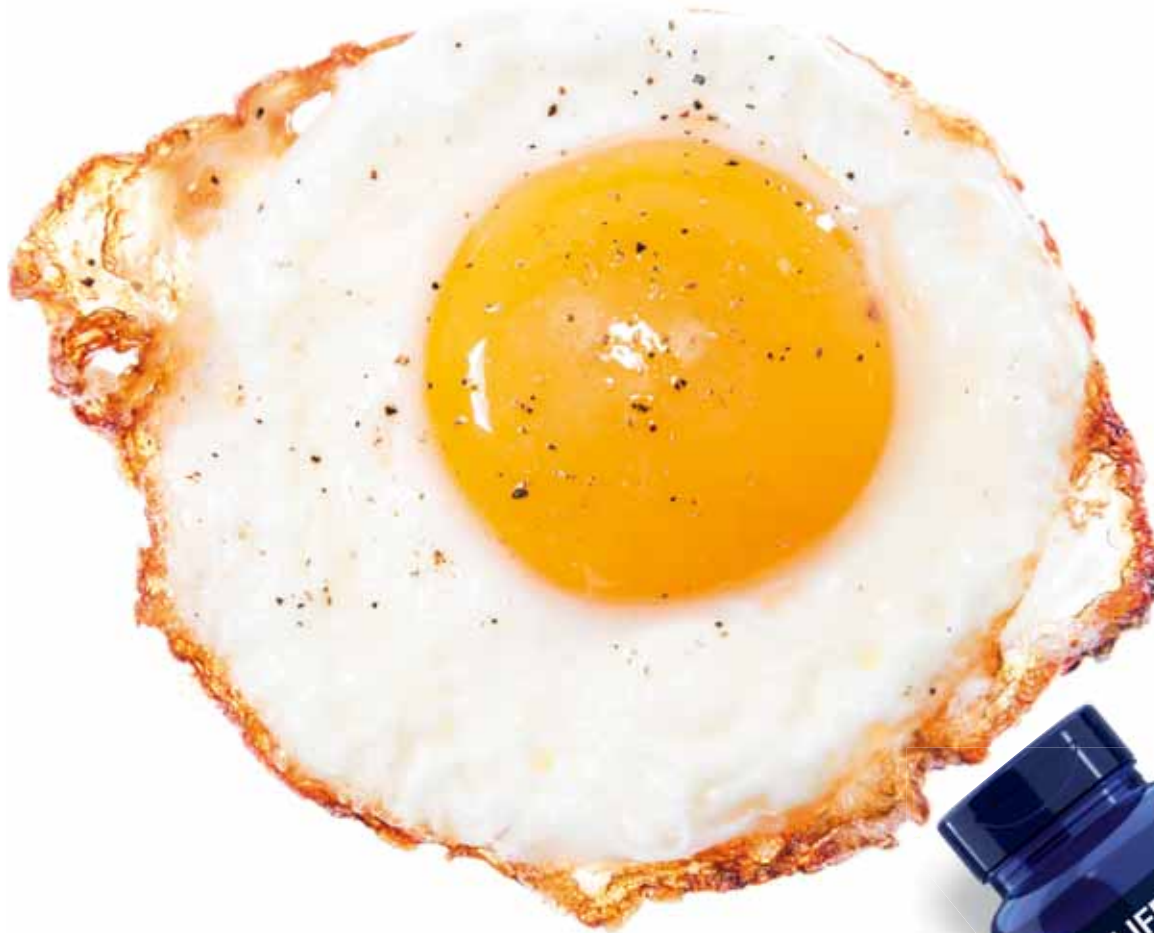
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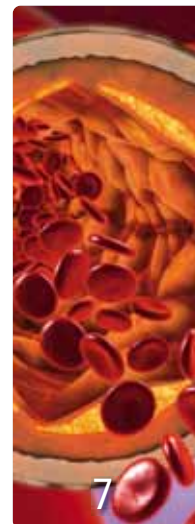
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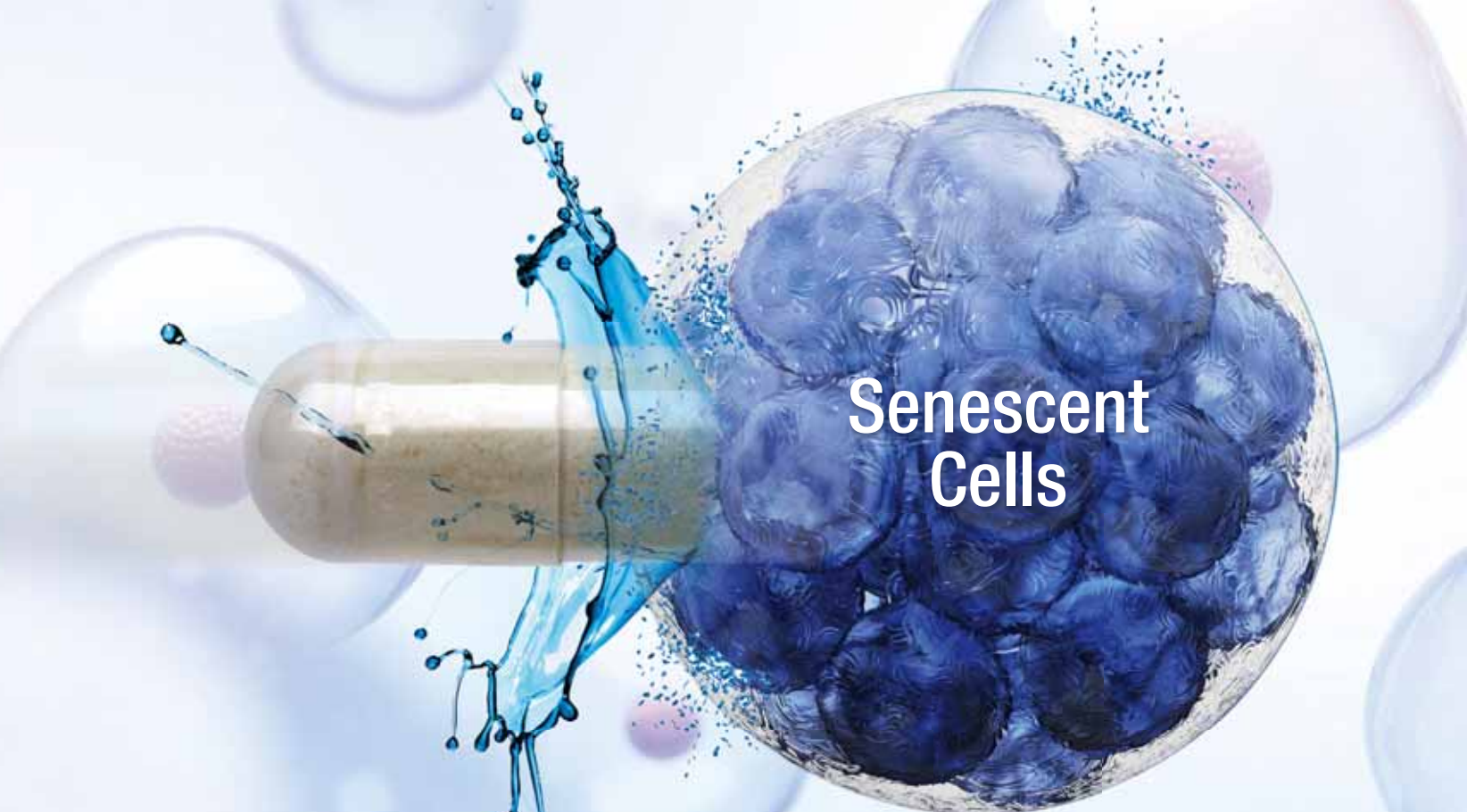
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Atherosclerosis in Sweden: Implications for Americans



WILLIAM FALOON

The **American Heart Association** published a report on a study that detected **atherosclerosis** in **42%** of **Swedish** study subjects without known **heart disease**.¹

Compared to **Americans**, most **Swedish** citizens have lower vascular risks.²

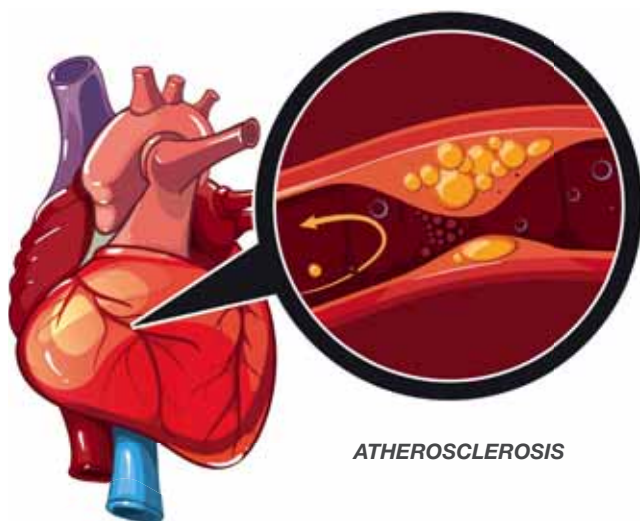
In 2019, the percentage of **obese** people in Sweden was **13.1%** compared to **40%** in the United States.³

Coronary **heart disease** deaths are **22.5% higher** in the **United States** than in **Sweden**.⁴

These data about artery disease in **Sweden** likely indicate *greater* **atherosclerosis** prevalence in the **United States**.

Autopsy studies on young Americans consistently show evidence of **atherosclerosis**.^{5,6}

Recognized risk factors of severe **artery disease** include hypertension,⁷⁻⁹ elevated lipids^{7,10-12} diabetes,^{7,13} and/or obesity.^{7,14}



A common misconception is that **atherosclerosis** is a **modern-day** phenomenon.

Studies using high-resolution images of **mummified bodies** from different world regions reveal the opposite. **Atherosclerosis** has plagued mankind for thousands of years.¹⁵

Atherosclerosis is the term used to describe damage and obstruction that occurs in the linings of our arteries.¹⁶

When **arterial blood flow** is impeded or blocked, the result can be death or disability from a **stroke** or **heart attack**.¹⁶

A misconception about **atherosclerosis** is that it did not exist in prior historic eras. I've heard so-called experts claim that people didn't suffer coronary artery disease in the past because their diets were so "natural."

These "experts" do not have sufficient data and have no basis to claim that atherosclerosis is a recent phenomenon. All they can cite is that few people developed **heart disease** in the past compared to today.

The reason more people weren't having **heart attacks** back then is that most died before the age of **40-60** from diseases that are now curable.

The advantage we have today is being able to delay **atherosclerotic** progression and reduce our risk of coronary **heart attack** and ischemic **stroke**.

Atherosclerosis in Ancient Cultures

A study published in **2011** found definitive or probable **atherosclerosis** present in **mummified** bodies that lived during every era of **ancient Egypt** studied.¹⁷

In **2013**, *The Lancet* showed that **mummified humans** dating back 4,000 years and spanning several geographic locations/cultures all suffered from **clogged arteries**.¹⁵

According to the lead investigator of *The Lancet* study:

“The fact that we found similar levels of atherosclerosis in all of the different cultures we studied, all of whom had very different lifestyles and diets, suggests that atherosclerosis may have been far more common in the ancient world than previously thought.”¹⁵

—Randall Thompson, MD

Modern Era Atherosclerosis

A **2021 Swedish** study found a **1.8-times** greater prevalence of **atherosclerosis** in people aged **60-64** compared to those **50-54** years old.¹



As we at *Life Extension* have long argued, **atherosclerosis** is a **disease of aging** that has long been prevalent.¹⁸

It took modern computed tomography **imaging** to document probable signs of **heart disease** in **34%** of mummified humans whose estimated mean age at death was **43** years. Those who died at an estimated mean age of **32** years were less likely to show signs of **atherosclerosis**.¹⁵

These findings suggest that **arterial disease** has always been widespread. The encouraging news is that methods to prevent and even reverse it have been validated in modern clinical studies.

What Caused Atherosclerosis In The Mummified Bodies?

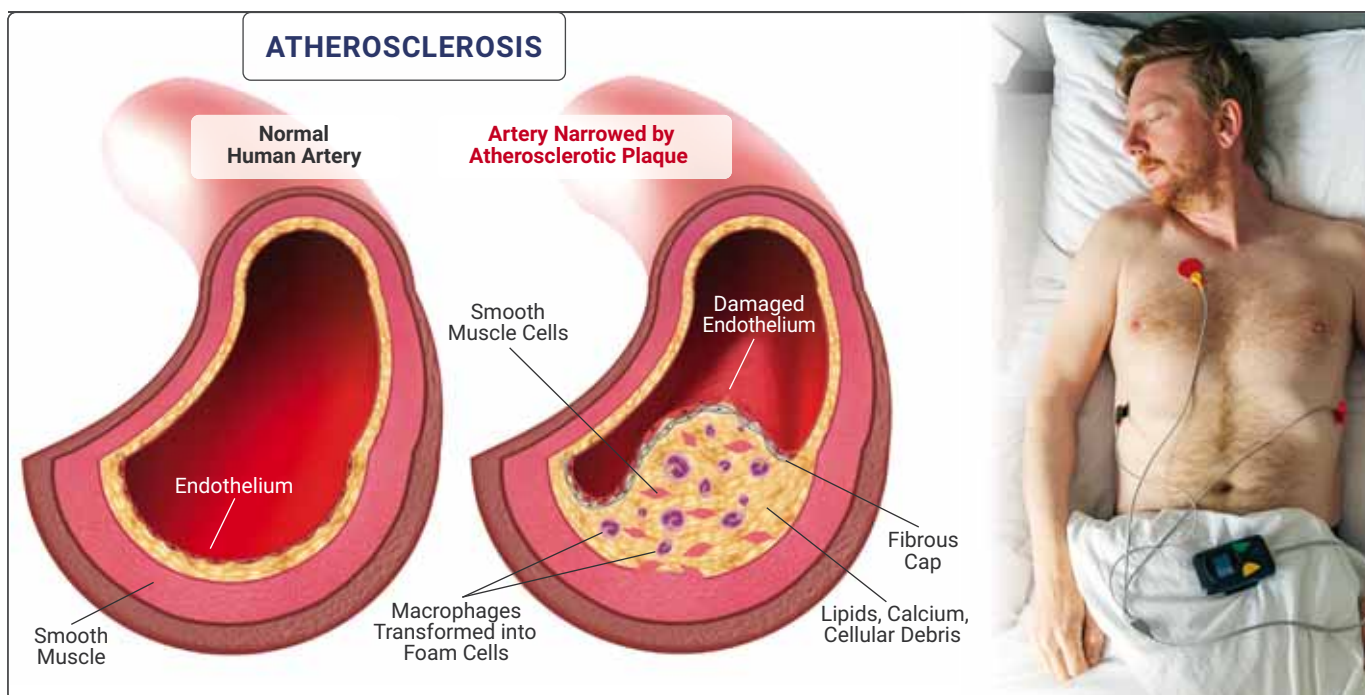
In seeking to identify what caused atherosclerosis in the mummified bodies, *The Lancet* researchers speculated that it could have been the high levels of **inflammation** these primitive people were exposed to from constant **infections** and a non-hygienic environment.¹⁵

People with severe inflammatory conditions (like rheumatoid arthritis^{19,20} and lupus^{21,22}) suffer *accelerated* atherosclerosis.

Multiple studies of modern people link **chronic inflammation** as an independent risk factor for **stroke**²² and **heart attack**.²²⁻²⁴

One study investigated a large group of older individuals who were followed for **17** years. Those with the *highest* levels of inflammatory blood markers were over **three-times** more likely to suffer sudden **cardiac death** compared to people with the lowest levels.²⁵

It's comforting to live in an era where **chronic inflammation** can be identified with **blood tests** and corrective actions taken to suppress elevated **inflammatory** markers.



Why Is Atherosclerosis So Common?

We know that multiple factors can alter arterial function and damage the delicate arterial lining (endothelium). These negative factors include excess dietary intake of the wrong kinds of fats,²⁶ sugars,²⁷ high salt intake,²⁸ and tobacco smoke.^{29,30}

Most of us are born with an intact **endothelium** that lines our arteries and protects against the formation of **atherosclerosis**.

As we age, our arterial lining endures chronic insult from internal and external factors that results in endothelial damage leading to **atherosclerosis**. Progressing blockage of blood vessels results in decreased blood flow to the heart, kidneys, and brain.^{18,31}

Health issues linked to atherosclerosis are a leading reason why most aging people die in the United States.¹⁶

Those who understand the many causes of **atherosclerosis** should not be surprised that it is so common. What's appalling is the number of people who continue to succumb to atherosclerotic diseases because of apathy and ignorance.

What Can You Do About It?

Endothelial dysfunction is one reason why **atherosclerosis** is such a common worldwide pathology.³¹

As the inner arterial wall (the **endothelium**) becomes **dysfunctional**, it enables **atherosclerotic** lesions to initiate, progress, and eventually occlude blood flow.³¹

As you will read on page 40 of this issue, a combination of low-cost nutrients has been shown to help modulate endothelial dysfunction and improve blood flow.

Annual Blood Test Sale

We at **Life Extension**® have long encouraged annual **blood tests** to screen for underlying arterial disease and take corrective actions when risk factors like homocysteine, C-reactive protein, and LDL are at dangerous levels.

Once a year, prices are **discounted** so our supporters can assess their status and initiate preemptive measures before illnesses strike.

On page 11 there is a description of the popular **Male** and **Female Blood Test** panels.

Commercial labs charge thousands for these tests, but readers of this publication obtain them for only **\$224** during the **Blood Test Super Sale**.

To order the **Male** or **Female Blood Test Panel** at these low prices, call **1-800-208-3444** (24 hours) or visit: www.LifeExtension.com/sales/lab-tests

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William Faloon, Co-Founder
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Homocysteine
Total Cholesterol
LDL (low-density lipoprotein)
HDL (high-density lipoprotein)
Triglycerides

GLUCOSE PROFILE

Insulin
Hemoglobin A1c
Glucose

MINERALS

Magnesium (serum)
Ferritin (measure of iron status)
Calcium
Potassium
Phosphorus
Sodium
Iron
Chloride

ORGAN FUNCTION

Kidney: creatinine, BUN, uric acid, BUN/creatinine ratio
Liver: AST, ALT, LDH, GGT, bilirubin, alkaline phosphatase

COMPLETE BLOOD COUNT (CBC)

Blood proteins: albumin, globulin, total protein, albumin/globulin ratio
Immune Cells including: lymphocytes, monocytes, eosinophils, neutrophils, basophils
Red Blood Cells including: hemoglobin, hematocrit, MCV, MCH, MCHC, RDW
Platelets

HORMONES

Free and Total Testosterone
DHEA-S
Estradiol (an estrogen)
TSH (thyroid function)
Vitamin D (25-hydroxyvitamin D)

CANCER MARKER

PSA (Prostate Specific Antigen)

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CARDIAC MARKERS

C-Reactive Protein (high sensitivity)
Apolipoprotein B (ApoB)
Homocysteine
Total Cholesterol
LDL (low-density lipoprotein)
HDL (high-density lipoprotein)
Triglycerides

GLUCOSE PROFILE

Insulin
Hemoglobin A1c
Glucose

MINERALS

Magnesium (serum)
Ferritin (measure of iron status)
Calcium
Potassium
Phosphorus
Sodium
Iron
Chloride

ORGAN FUNCTION

Kidney: creatinine, BUN, uric acid, BUN/creatinine ratio
Liver: AST, ALT, LDH, GGT, bilirubin, alkaline phosphatase

COMPLETE BLOOD COUNT (CBC)

Blood proteins: albumin, globulin, total protein, albumin/globulin ratio
Immune Cells including: lymphocytes, monocytes, eosinophils, neutrophils, basophils
Red Blood Cells including: hemoglobin, hematocrit, MCV, MCH, MCHC, RDW
Platelets

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Progesterone
Free and Total Testosterone
DHEA-S
Estradiol (an estrogen)
TSH (thyroid function)
Vitamin D (25-hydroxyvitamin D)

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FT. LAUDERDALE, FL, 33304**

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MALE AND FEMALE PANELS

- | | SALE PRICE |
|---|------------|
| <input type="radio"/> MALE PANEL — NOW WITH FERRITIN (LC322582)
CBC/Chemistry/Lipids Panel • DHEA-S • PSA (prostate-specific antigen)
• Homocysteine • C-Reactive Protein (high sensitivity) • ApoB • Free Testosterone • Total Testosterone • Estradiol • TSH for thyroid function
• Vitamin D (25-hydroxyvitamin D) • Hemoglobin A1c • Insulin • Magnesium | \$224 |
| <input type="radio"/> FEMALE PANEL — NOW WITH FERRITIN (LC322535)
CBC/Chemistry/Lipids Panel • DHEA-S • Estradiol • Homocysteine • ApoB
• C-Reactive Protein (high sensitivity) • Progesterone • Free Testosterone
• Total Testosterone • TSH for thyroid function • Vitamin D (25-hydroxyvitamin D)
• Hemoglobin A1c • Insulin • Magnesium | \$224 |

CARDIAC RISK ASSESSMENTS

- | | |
|---|----------|
| <input type="radio"/> NMR LIPOPROFILE® (LC123810)
The NMR Lipoprofile® directly measures LDL particle size and number as well as HDL particle number, total cholesterol, and triglycerides. It also provides a calculation of one's risk of insulin resistance by assessing abnormalities in lipoprotein markers. | \$74.25 |
| <input type="radio"/> NMR LIPOPROFILE PLUS* (LC100049)
In-depth analysis of cardiovascular risk markers including: NMR LipoProfile, C-Reactive Protein, Myeloperoxidase, and Oxidized LDL . | \$201.75 |
| <input type="radio"/> OXIDIZED LDL (LC123023)
Oxidized low-density lipoprotein (LDL) cholesterol is one of the main causes of the formation of atherosclerotic plaque in the arterial wall. This blood test measures levels of oxidized LDL. | \$56.25 |
| <input type="radio"/> MYELOPEROXIDASE (MPO)* (LC123006)
The myeloperoxidase (MPO) test measures levels of an enzyme that oxidizes low-density lipoprotein (LDL) cholesterol, which could lead to increased arterial plaque formation. | \$74.25 |
| <input type="radio"/> ADVANCED OXIDIZED LDL PANEL* (LC100035)
This panel looks at vascular inflammatory biomarkers, beginning with lifestyle choices to the development of metabolic and cardiovascular disease as well as the formation of vulnerable plaque. The panel contains the following tests: F2-Isoprostanes, Myeloperoxidase, and Oxidized LDL. | \$198.75 |
| <input type="radio"/> OMEGA-3 INDEX COMPLETE** (LC100066)
Beneficial for everyone! People <u>not</u> taking omega-3/fish oil should check their baseline Omega-3 Index to see if it is in the desirable or concerning range. Those taking Omega-3/fish oil supplements should take the test to see if they need to adjust their dosage. You want to target a range of 8%-12% for your Omega-3 Index score. | \$74.25 |

CONDITION-SPECIFIC TESTS

- | | |
|--|----------|
| <input type="radio"/> PERSONALIZED AMINO ACID HEALTH ASSESSMENT** (LC100090)
An in-depth analysis of amino acid metabolism provides insight into various health concerns, such as maldigestion, GI dysbiosis, neurological issues and more, with a personalized report of diet and supplementation suggestions. Provided as an at-home urine collection kit. | \$224.25 |
| <input type="radio"/> ENVIRONMENTAL POLLUTANTS PROFILE*** (LC100089)
Toxic pollutants affect our bodies in several different ways. This test helps assess possible exposure to several common environmental pollutants by measuring their urinary metabolites. | \$111.75 |

Know YOUR NUMBERS



- | | SALE PRICE |
|--|------------|
| <p>BLOOD METALS PANEL BLOOD SPOT KIT *** (LC100092) \$123.75</p> <p>Convenient at-home test for blood levels of 8 important metals, providing insight into toxic metal burden and nutritional status with a quick and easy finger stick. Detects toxic metals: Lead, Mercury, Cadmium, Arsenic, Antimony and nutritional elements: Copper, Zinc, Selenium</p> | |
| <p>TOXIC METALS PANEL (FECAL)*** (LC100076) \$127.50</p> <p>The results of fecal elemental analysis can help you identify and eliminate dietary exposure to toxic metals, while also assessing the body's natural excretion of metals. The panel tests Antimony, Arsenic, Beryllium, Bismuth, Cadmium, Copper, Lead, Mercury, Nickel, Platinum, Thallium, Tungsten, and Uranium.</p> | |
| <p>NEUROTRANSMITTER PANEL-COMPREHENSIVE*** (LC100085) \$221.25</p> <p>Serotonin, Dopamine, Epinephrine, Norepinephrine, GABA, Glutamate, Glycine, Histamine, PEA, DOPAC, 3-MT, Normetanephrine, Metanephrine, 5-HIAA, Tryptamine, Tyrosine, Tyramine, Taurine. Alterations in neurotransmitters play a significant role in contributing to symptoms such as cognitive disorders, depression, anxiety, diminished drive, fatigue and sleep difficulties, craving, addictions, pain, and more.</p> | |
| <p>PRO7 ADVANCED NUTRIGENOMIC PROFILE** (LC100097) \$224.25</p> <p>Analyzes genetics in 7 key areas of health (methylation, neurotransmitters, mitochondrial health, detoxification, inflammation, and more) and provides genetically targeted nutritional suggestions.</p> | |
| <p>APOE GENETIC TEST FOR ALZHEIMER'S AND CARDIAC RISK** (LC100059) \$111.75</p> <p>This simple at-home cheek swab reveals your ApoE genotype and association with risk of Alzheimer's and cardiovascular disease.</p> | |
| <p>FOOD SENSITIVITY PANEL-ELITE*** (LC100096) \$336.75</p> <p>This in-depth food sensitivity assessment measures both IgG and IgA response to 240 foods.</p> | |
| <p>GUT BARRIER PANEL** (LC900004) NEW \$123.75</p> <p>Measure immune response (IgG/C3d & IgA) to three key GI markers associated with intestinal permeability and leaky gut: Candida, Zonulin, and Occludin.</p> | |

GUT MICROBIOME TESTING

- | | |
|---|--|
| <p>CPP STOOL ANALYSIS (CPP) *** (LC100093) \$224.25</p> <p>Good starting point for evaluating microbiome-related gastrointestinal concerns. Identifying presence of common pathogenic microorganisms associated with acute GI distress. Includes advanced bacteria/yeast culture, pathogen detection by PCR, and parasitology.</p> | |
| <p>COMPREHENSIVE STOOL ANALYSIS (CSAP) *** (LC100083) \$299.25</p> <p>Better assessment of the gut microbiome and digestive health. Contains all the tests in the CPP Stool Analysis with additional markers of digestive function, inflammation & immune function (stool chemistry), important for ongoing chronic and acute GI concerns.</p> | |
| <p>GI360™ *** STOOL ANALYSIS (LC100088) \$379.25</p> <p>Best and most advanced assessment of microbiome dysbiosis, diversity and digestive health, the GI360 provides the most information about gut and microbiome health. Contains all the tests contained in the CPP and CSAP along with cutting-edge markers of microbiome diversity and dysbiosis.</p> | |

- | | SALE PRICE |
|---|------------|
| WHOLE-BODY HEALTH | |
| <p>MALE ELITE PANEL* (LC100016) \$431.25</p> <p>CBC/Chemistry/Lipids Panel • Free and Total Testosterone • Estradiol • Total Estrogens • DHEA-S • Progesterone • Pregnenolone • TSH • DHT • FSH • LH • Free T3 • Free T4 • Reverse T3 • IGF-1 • SHBG • Free and Total PSA • Vitamin D 25-OH • hs-CRP • Ferritin • TIBC • Homocysteine • Insulin • Hemoglobin A1c • Cortisol • ApoB • Magnesium</p> | |
| <p>MALE BASIC HORMONE PANEL (LC100012) \$56.25</p> <p>DHEA-S • Estradiol • Total and Free Testosterone • PSA</p> | |
| <p>FEMALE ELITE PANEL* (LC100017) \$431.25</p> <p>CBC/Chemistry/Lipids Panel • Free and total Testosterone • Estradiol • Total Lipids • 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 • TIBC • Homocysteine • Insulin • Hemoglobin A1c • Cortisol • ApoB • Magnesium</p> | |
| <p>FEMALE BASIC HORMONE PANEL (LC100013) \$56.25</p> <p>DHEA-S • Estradiol • Total and Free Testosterone • Progesterone</p> | |
| <p>WEIGHT LOSS PANEL-COMPREHENSIVE (LC100028) \$206.25</p> <p>CBC/Chemistry/Lipids Panel • DHEA-S • Free and Total Testosterone • Estradiol • Progesterone • Cortisol • TSH • Free T3 • Free T4 • Reverse T3 • Insulin • Hemoglobin A1c • Vitamin D 25-hydroxy • Ferritin • C-reactive protein (high sensitivity)</p> | |
| <p>HEALTHY AGING PANEL-COMPREHENSIVE* (LC100026) \$186.75</p> <p>CBC/Chemistry/Lipids Panel • C-reactive protein (high sensitivity) • Vitamin B12 • Folate • Homocysteine • Vitamin D 25-hydroxy • Hemoglobin A1c • TSH • Free T3 • Free T4 • Ferritin • Urinalysis • ApoB • Insulin</p> | |
| <p>COMPLETE HORMONE & METABOLITES PROFILE (HUMAP)*** (LC900001) \$224.25</p> <p>Comprehensive assessment of hormones and their metabolites. This urine test helps assess how your body metabolizes hormones, providing insight into endogenous hormone secretion, supplemental hormone utilization, enzyme activity, oxidative stress.</p> | NEW |

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These **CBC/Chemistry/Lipids Panel** tests are included in the popular **Male and Female Panels**, and other panels on this page so you don't have to order them separately.

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Total Cholesterol • HDL Cholesterol • LDL Cholesterol Triglycerides Cholesterol/HDL Ratio • Estimated CHD Risk • Glucose

LIVER FUNCTION

AST (SGOT) • ALT (SGPT) • LDH • Total Bilirubin • Alkaline phosphatase

KIDNEY FUNCTION

BUN • Creatinine • BUN/Creatinine Ratio • Uric Acid

BLOOD PROTEINS

Total Protein • Albumin • Globulin • Albumin/Globulin Ratio

BLOOD COUNTS

Red Blood Cell Count • White Blood Cell Count • Eosinophils Neutrophils (Absolute) • Lymphs (Absolute) • Eos (Absolute) Baso (Absolute) • RDW • Monocytes (Absolute) • Monocytes Lymphocytes • Platelet Count • Hemoglobin • Hematocrit

MCV • MCH • MCHC • Neutrophils

BLOOD MINERALS

Calcium • Potassium • Sodium • Chloride • Iron

The price for the **CBC/Chemistry/Lipids Panel** alone is **\$26.25**.
(LC381822)

This is NOT a complete listing of LE lab test services.
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* This test requires samples to be shipped to the lab on dry ice for customers using a Blood Draw Kit. Customer is responsible for obtaining dry ice.
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† Not available in NY.

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In the News



Lithium Benefits Kidneys

New research published in the *Journal of Clinical Investigation* suggests that **lithium** may help protect against kidney decline.*

Inhibiting the **glycogen synthase kinase 3-beta** (GSK3-beta) enzyme in cells supports kidney health in pre-clinical models.

Lithium has been shown to inhibit **GSK3-beta**.

Researchers have also observed that psychiatric patients treated long-term with lithium carbonate exhibited better **kidney** function than age-matched patients who did not receive it.

Their research suggested that less than a third of the dose used for bipolar disorder significantly blocks GSK3-beta activity in the kidney.

Editor's Note: Lithium has been used for many years as a treatment for bipolar disorder, and more recently was also found to extend the lifespan of roundworms and fruit flies.

* *J Clin Invest.* 2022 Feb 15. ;132(4):e141848.

N-Acetylcysteine Could Help in Older Patients' Concussion Recovery

A pilot study revealed improvement in symptoms of concussion among older men and women with mild traumatic brain injury who received supplements containing N-acetylcysteine (NAC), the *Journal of Trauma Acute Care Surgery* reported.*

The study included 65 traumatic brain injury patients who were within three hours of trauma surgery service evaluation.

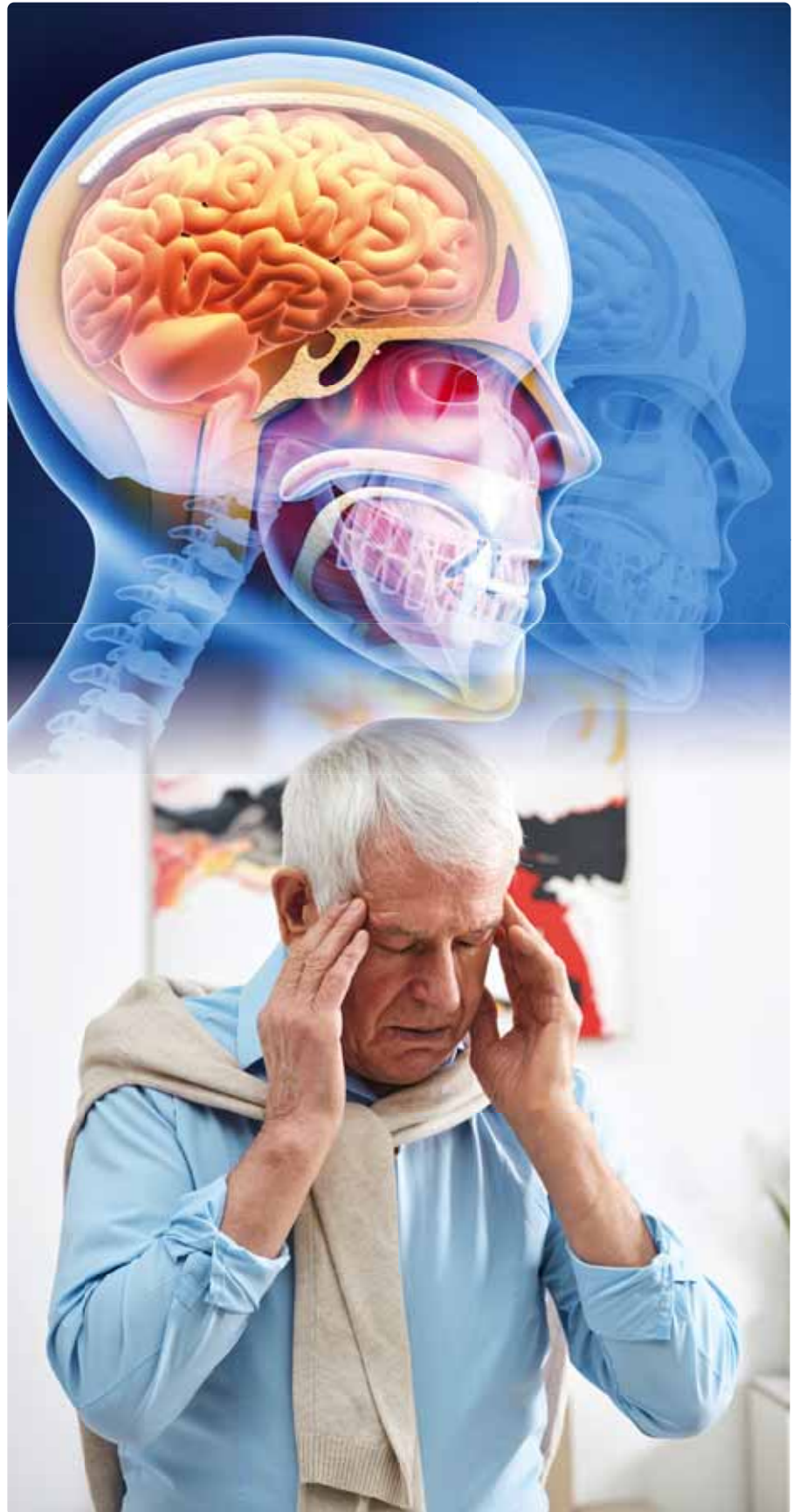
Thirty-four patients received standard treatments for traumatic brain injury plus **4 grams** of NAC within three hours of injury, followed by a 3-day period during which **2 grams** of NAC was administered twice per day and an additional 3 days during which **1.5 grams** of NAC was provided twice daily.

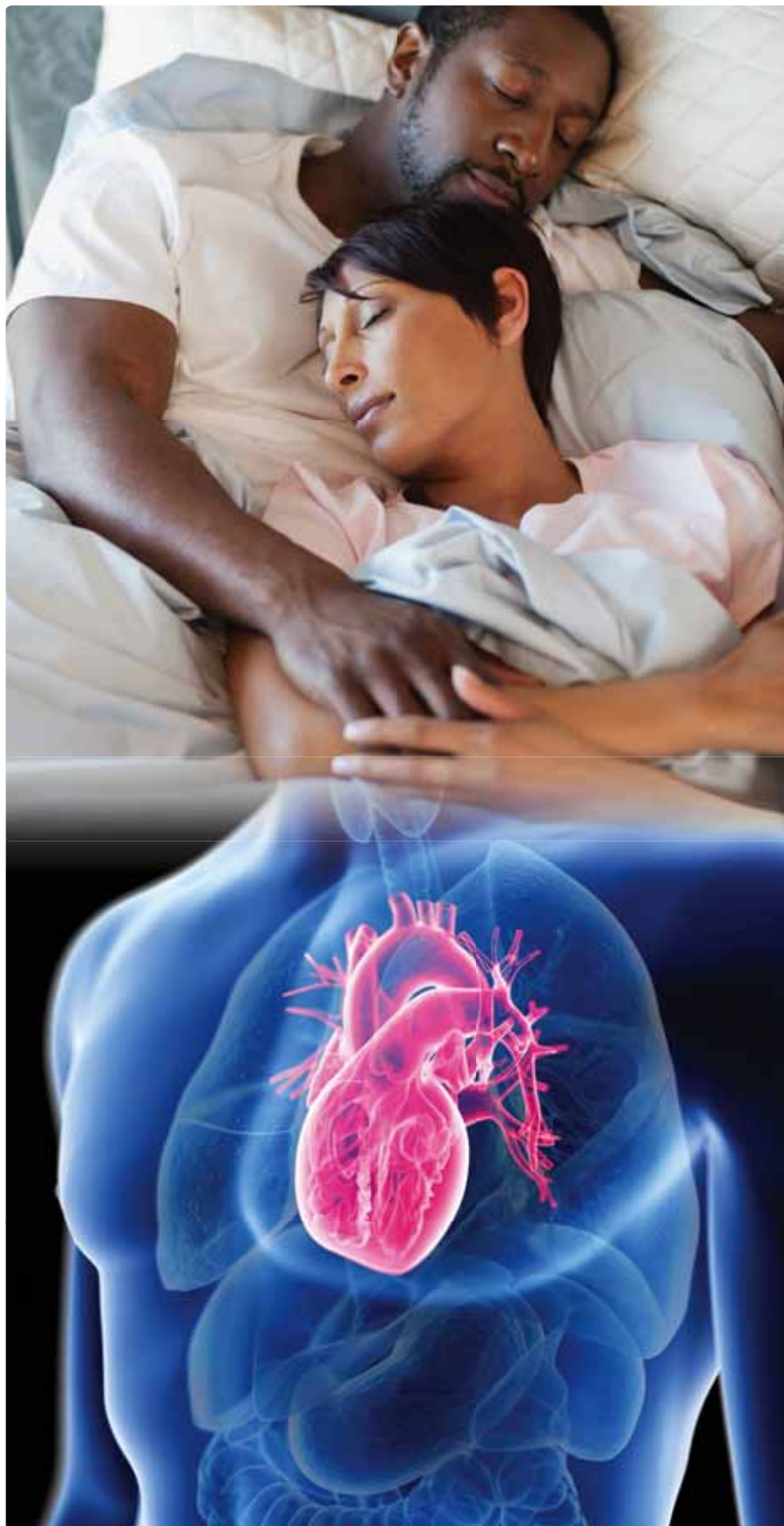
The remainder of the participants received standard treatment alone. Questionnaires that evaluated post-concussion symptoms were administered at the beginning of the study and at 7 and 30 days.

While questionnaire scores were similar at the beginning of the study, they were significantly better in the **NAC group** on days 7 and 30.

Editor's Note: The authors concluded that, "N-acetylcysteine was associated with significant improvements in concussion symptoms in elderly patients with mild traumatic brain injury. These results justify further research into using NAC to treat traumatic brain injury."

* *J Trauma Acute Care Surg.* 2022 Apr 8.





Melatonin Improves Heart Failure Patients' Lives

Stable **heart failure** patients with reduced ejection fraction (when left ventricle of the heart fails to contract normally) who consumed a nightly **melatonin** supplement, experienced a reduction in a marker of heart failure and better **quality of life** compared to patients who received a **placebo**, the journal *Clinical Cardiology* reported.*

The clinical trial included 85 patients who received **10 mg** melatonin or a placebo nightly for 24 weeks.

Levels of NT-pro BNP which, when elevated, may be an indicator of heart failure, were similar in the melatonin-supplemented group and the placebo group at the beginning of the trial.

At the end of the trial, NT-pro BNP declined to **221.1 ng/L** among participants who received **melatonin** compared to **332.1 ng/L** among those who received a **placebo**.

Lower **NT-pro BNP** blood levels are an indicator of reduced heart failure severity.

Editor's Note: Melatonin-supplemented participants also experienced significant improvements in clinical outcome, quality of life, and New York Heart Association classification of heart failure, compared to the placebo group.

* *Clin Cardiol.* 2022 Apr;45(4):417-426.

Compounds for Preserving Bone Health

Bone health is maintained by two key cells called osteoblasts, which differentiate into osteocytes (bone cells), and osteoclasts, which break down old bone so that osteoblasts can form new bone.

The health and function of osteoblasts and osteoclasts is regulated by pathways which have been found to be activated by ‘cellular’ proteins such as sirtuin 1, AMPK, and Nrf2.

Researchers have reviewed bone preserving mechanisms of specific nutrients that regulate these cellular proteins.*

SIRT1 is increased by melatonin, nicotinamide riboside, glucosamine and thymoquinone, found in black cumin seed oil. (Resveratrol long ago demonstrated potent SIRT1 activation.)

Nutrients such as gynostemma pentaphyllum and the drug metformin are compounds that activate AMPK.

Nrf2 regulates the cells’ defense against oxidative stress. Lipoic acid, melatonin, thymoquinone, astaxanthin and sulforaphane can promote Nrf2 activity.

Editor’s Note: “Regimens providing a selection of these nutraceuticals in clinically meaningful doses may have an important potential for preserving bone health,” the authors concluded. “Concurrent supplementation with taurine, N-acetylcysteine, vitamins D and K2, and minerals, including magnesium, zinc, and manganese, plus a diet naturally high in potassium, may also be helpful in this regard.”

* *Int J Mol Sci.* 2022 April 26;23(9):4776.



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*European Journal of Nutrition. 2011;50(5):387-389.

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A woman with grey hair, wearing a white wide-brimmed hat, a blue long-sleeved top with a lace-up front, and white pants, is walking on a stone-paved path in a village. She is smiling and looking upwards. The background shows a stone wall on the left and yellow buildings on the right.

A Solution for Urinary Symptoms in Women



BY MICHAEL DOWNEY

For many women, frequent trips to the bathroom or needing to get up at night to urinate are a drain on quality of life.^{1,2}

Even a good laugh can lead to urinary incontinence.

In an eight-week **clinical trial**, a blend of three **plant extracts** improved urination issues in women as follows:³

- Incontinence episodes were **reduced** by **65%**,
- Urinary urgency episodes **decreased** by **57%**,
- Nighttime urination episodes were **reduced** by **43%**, and
- Daytime urination frequency returned to **normal**.

A robust **79%** of women who took the blend reported a significant **benefit**, compared to **17%** who took a **placebo**.



Bladder Problems Worsen with Age

Urinary issues tend to become more common in women as they age. Onset of symptoms is usually observed over the age of 40.⁴

The prevalence and severity of symptoms are greater in women than in men. A population study with 40- to 99-year-old women participants from the U.S., UK, and Sweden revealed that:⁵

- **56%** experience **incontinence**,
- **36%** experience **urinary urgency**,
- **34%** experience **nighttime urination**, and
- **25%** experience **frequent urination**.

The median daytime urinary frequency is 3-4 hours (6-8 times daily).⁶ Those afflicted with an overactive bladder have to go to the bathroom more frequently (>8 times during the day and >1 time at night).²

According to a study, *only 46%* of symptomatic women have discussed any urinary concerns with their health care provider.⁴ Many are self-conscious about these symptoms but assume they are an unavoidable part of aging.⁷

Those who do seek medical advice are often prescribed drugs with **minimal benefits** at best. For example, only about **13%** of participants taking drugs achieve urinary continence, and side effects prompt some patients to discontinue medication.³

In contrast, a clinical study found that a blend of **three plant extracts**³ was well-tolerated by participating women, *improved* urinary symptoms, and significantly *improved* participants' quality of life.

How Does this Herbal Combination Work?

In a rat model of **overactive bladder**, the herbal blend reversed the alterations in various biomarkers in the urine and lining of bladder and muscle, leading to improvement of urinary symptoms in:⁷

- Storage phase (e.g., urgency, frequency, nocturia)
- Voiding/Post-voiding phase (e.g., hesitancy, intermittency, weak stream, dribbling post-voiding)

The researchers suggest that the ability of these **plant extracts** to favorably alter markers of urinary changes in the rat model may explain the clinically significant benefits observed in the **human** study.⁷

Twenty Years in Development

More than **20 years** ago, naturopath and medical herbalist Dr. Tracey Seipel began researching an effective solution to **urinary problems** for her patients.⁸ She sifted through the medical literature on **plant compounds** that had been traditionally used to treat bladder issues and gradually began incorporating some into her clinical practice.³

Eventually, Dr. Seipel's experiential research allowed her to refine a treatment that included a blend of **three plant extracts**, each with a history of effectiveness and each from a different area of health care:

- **Horsetail** (*Equisetum arvense*) from Western herbal medicine,
- **Lindera** (*Lindera aggregata*) from Chinese medicine, and
- **Three-leaf caper** (*Crateva nurvala*) from Ayurvedic medicine.

Reduced Urinary Symptoms

To validate Dr. Seipel's research on the blend of three plant extracts, scientists designed a randomized, **placebo-controlled** trial.³

They enlisted 88 women with an average age of **62 years** who had at **least two** of the following symptoms:

- Daytime urination episodes of **10 or more** a day,
- Incontinence episodes of **one or more** per day,
- Urinary urgency episodes of **two or more** a day, and
- Nighttime urination episodes of **two or more** a night.



HORSETAIL

WHAT
YOU
NEED
TO
KNOW

The treatment group took **840 mg** of the extract blend in the form of two capsules once daily. After just **eight weeks**:³

- Daytime urination episodes were restored to normal levels, decreasing from an average of **11.59 times** a day to an average of just **7.88 times** a day.
- Incontinence episodes were reduced by **65%**, from an average of **3.49 times** a day to an average of only **1.21 times** a day,
- Urinary urgency episodes decreased by **57%**, from an average of **3.77 times** a day to an average of just **1.61 times** a day, and
- Nighttime urination episodes were reduced by **43%**, from an average of **3.76 times** a night to an average of **2.15 times** a night.

Improved Quality of Life

Women taking the plant extract also reported impressive improvements in quality-of-life questionnaires, including a:³

- **50% decrease** in overactive bladder,
- **39% decrease** in incontinence, and
- **39% decrease** in urogenital distress.

A remarkable **79%** of women who took the blend reported feeling a **significant benefit**, compared to **17%** of women in the placebo group.

The extract blend produced these results **without the side effects** commonly seen with medications.³

Help For Women's Bladder Problems

- **Urinary symptoms** are common in women, especially after age 40. These include frequent urination, urinary incontinence, urinary urgency, and nighttime urination.
- A formula has been developed from extracts of **three** plants with a history of traditional use in addressing bladder disorders: **horsetail, lindera, and three-leaf caper**.
- In an animal model of overactive bladder, the herbal combination treatment reversed the detrimental shifts in bladder function parameters and in the levels of several tested biomarkers in the bladder epithelium and muscle.
- A placebo-controlled **clinical** trial has validated that this three-extract blend significantly improves urinary symptoms in women *without* harsh side effects.
- After eight weeks, this combination reduced daytime urination frequency to the **normal level** and incontinence episodes by **65%**. It also significantly reduced urinary urgency and nighttime urination and improved quality of life.

Summary

Urinary problems are common in women as they age and can impact the quality of their life.

In a clinical study, a blend of extracts from the plants **horsetail**, **lindera**, and **three-leaf caper** was shown to *reduce* daytime urinary frequency, urinary incontinence, urinary urgency, and nighttime urination in women.

These benefits translate into improved quality of life, without the adverse side effects of drugs commonly prescribed for these problems. •

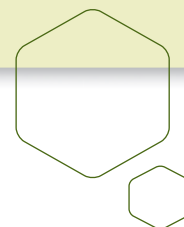
If you have any questions on the scientific content of this article, please call a **Life Extension** Wellness Specialist at 1-866-864-3027.

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Lower Urinary Tract Symptoms

- **Lower urinary tract symptoms (LUTS)** are divided into storage and voiding/ micturition (urination) phases.
- Storage phase involves **overactive bladder (OAB)** and **stress incontinence (SI)**.
- **Overactive bladder (OAB)** is a group of urinary symptoms, defined as having an urgent need to empty the bladder during the day or night, with or without incontinence.
- Median daytime urinary frequency is **3-4** hours **6-8** times daily.⁶
- Those afflicted with an overactive bladder have to go to the bathroom frequently (**>8** times during the day and **>1** time at night), may leak urine into their clothes, and report feeling depressed, stressed, and sleep deprived.
- This could be due to overactivity of the bladder detrusor (a muscle lining the urinary bladder, that manages storage and voiding of urine).²
- **Stress incontinence** is another common bladder problem in which women leak urine while sneezing, laughing, or doing other physical activities.
- Voiding symptoms include hesitancy, straining, terminal dribbling, intermittency, and slow, weak and/or interrupted stream.²



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New Study Demonstrates Significant **PAIN RELIEF**

BY MICHAEL DOWNEY

Some nutrients can reduce existing inflammation, while others may help resolve inflammation.

Winding down the flow of inflammation at its source is vital to maintain healthy, functional tissues.¹

Compounds that actively promote the resolution of inflammation are known as **specialized pro-resolving mediators** or **SPMs**.²

A clinical study published in **2022** demonstrates that taking **SPM precursors** along with bioavailable **curcumin** deliver significant relief of pain and discomfort.³

In just 30 days, the combination significantly reduced:

- **Total** pain,
- Pain **intensity**, and
- Pain **severity**.

After 60 days, a remarkable **79%** of participants had an improvement in **total pain**.





New Human Study

Curcumin, a compound found in the **turmeric** plant, is well-established as a particularly powerful **anti-inflammatory** and an important nutrient to reduce inflammation.^{4,5}

Specialized pro-resolving mediators (SPMs) are compounds produced in the body that resolve inflammation, helping return inflamed tissues back to their healthy state.²

Scientists wondered whether **combining SPM precursors** with **curcumin** might more thoroughly reduce inflammation *and* thus have an impact on **pain**.

A **2022** open-label **clinical** pilot study recruited healthy male and female adults with mild to moderate pain.³

Every day for 60 days, 29 participants were asked to take:

- One softgel containing **500 mg** of a marine oil enriched with **three SPM precursors**, and
- One capsule containing **500 mg** of a highly bioavailable (absorbable) **curcumin**.

Participants completed **three** well-known questionnaires used to measure pain, quality of life, and overall health:

- Short-Form McGill Pain Questionnaire (**SF-MPQ**),
- Short-Form 36 Health Survey (**SF-36**), and
- Medical Symptoms Questionnaire (**MSQ**).

Compelling Results

The results of the study were published in *Translational Medicine Communications*, a peer-reviewed medical journal.

The **SF-MPQ** (Short-Form McGill Pain Questionnaire) responses showed significant *improvements* in all aspects of the questionnaire within **30 days**, especially in:³

- **Total** pain,
- Pain **intensity**, and
- Pain **severity**.

The **SF-36** (Short-Form 36 Health Survey) questionnaire showed significant *improvements* in:³

- **Four** aspects of physical health, especially **pain** and **physical function**, and
- Perceived health change.

The **MSQ** (Medical Symptoms Questionnaire) results showed:³

- Significant *reduction* in **joint/muscle pain**.

An impressive **62%** of participants had an improved **total pain** score at **30 days**, and **79%** of participants showed improvement in **total pain** at **60 days**.

No adverse events were reported.³

This strongly suggests that taking **SPM precursors** with a bioavailable form of **curcumin** delivers significant relief of **pain** and discomfort associated with inflammation.

Difference Between Anti-Inflammatories and SPMs

Curcumin and **SPM precursors** target inflammation in completely different ways.

Curcumin and other **anti-inflammatories** work to reduce body inflammation levels.

This is helpful but not enough to completely restore health, since inadequate resolution can lead to chronic inflammation, excessive tissue damage, and dysregulation of tissue healing, and may also lead to fibrosis.⁶

Hence, inflammation needs to be resolved, to get tissues back to their healthy, functional state.

Resolution of inflammation is a complex, active process guided by specific signaling compounds produced in the body.^{7,8} Among these compounds are **specialized pro-resolving mediators**.

How SPMs Resolve Inflammation

SPM precursors are predominantly derived from **EPA** and **DHA**, the **omega-3** fatty acids found in fish oil.

The **precursors** needed to produce SPMs in the body include:⁹

- **18-HEPE** (18-hydroxyeicosapentaenoic acid),
- **17-HDHA** (17-hydroxydocosahexaenoic acid), and
- **14-HDHA** (14-hydroxydocosahexaenoic acid).

The marine-blend softgels used in the **2022** study provided a total of **300 mcg** of these three SPM precursors.³

These precursors are converted in the body into three different types of SPMs: **resolvins**, **protectins**, and **maresins**.^{3,10}

These make up the bulk of the SPMs that target **inflammation**. They do so through three mechanisms:¹⁰⁻¹²

- **Removing** dead and dying cells, helping to clean up the aftermath of inflammatory cascades,
- **Restoring** inflammation balance by decreasing pro-inflammatory mediators and increasing **anti-inflammatory** compounds, and
- **Renewing** damaged tissue by promoting cellular **regeneration**.

WHAT
YOU
NEED
TO
KNOW

Targeting Inflammation to Stop Pain

- **Inflammation** is a major risk factor for age-related disease and degenerative disorders. It is also a source of pain.
- **Curcumin** is a well-known anti-inflammatory, working to *reduce* inflammation.
- Other compounds called **specialized pro-resolving mediators (SPMs)** *resolve* inflammation, shutting off inflammation and returning tissues to a healthy state.
- A new clinical trial shows that combining a highly bioavailable **curcumin** with marine oil enriched with **SPM precursors** significantly reduces subjective levels of **pain and discomfort**.



Evidence for Curcumin and SPM Benefits

Before researchers tested the *combined* effects of SPM precursors and curcumin, many studies had shown that each had benefits alone.

SPMS

Animal data showed promising results from the use of **SPMs**, including improvements in obesity-related **osteoarthritis**¹³ and in inflammation-induced **neuropathic pain**.¹⁴

A clinical trial found that oral intake of **omega-3s** increased SPM levels in the body by **229%** and significantly lowered levels of the inflammatory marker **C-reactive protein**.¹⁵

Another clinical study showed that taking marine oil enriched with a combination of **SPM precursors** (including **18-HEPE**, **17-HDHA**, and **14-HDHA**) increased SPM levels and helped **resolve** inflammation.¹⁶

CURCUMIN

Curcumin is known for the **curcuminoids** and polyphenols found in the **turmeric** plant.¹⁷

A meta-analysis of eight human trials involving 606 patients found that curcuminoids significantly reduced pain severity from a variety of causes, including arthritis and muscle soreness.¹⁸

Another review paper concluded that curcumin was safe and has significant **anti-inflammatory** activity.⁵

Combining SPM precursors with a powerful anti-inflammatory like curcumin targets inflammation in multiple ways and results in clear pain reduction.

Summary

Curcumin is a powerful **anti-inflammatory** that reduces existing inflammation.

Specialized pro-resolving mediators (SPMs) help *resolve* inflammation, guiding tissues back to their healthy state.

A new human study shows that *combining* marine oil enriched with **SPM precursors** with a highly bioavailable **curcumin** improves subjective measures of **pain** and discomfort associated with inflammation. ●





Building a Better Curcumin

Due to poor **bioavailability**, a large portion of curcumin taken orally never gets into the bloodstream or reaches tissues.

In a major advance, scientists combined **curcumin** with a fiber called **galactomannan** that protects curcumin in the gut.

Clinical research demonstrates that taking the curcumin-galactomannan combination results in blood levels of free curcuminoids that are over **45-times greater** than in those who take pure curcumin alone.¹⁹

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

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*Int J Med Sci. 2019;16(6):845-53.

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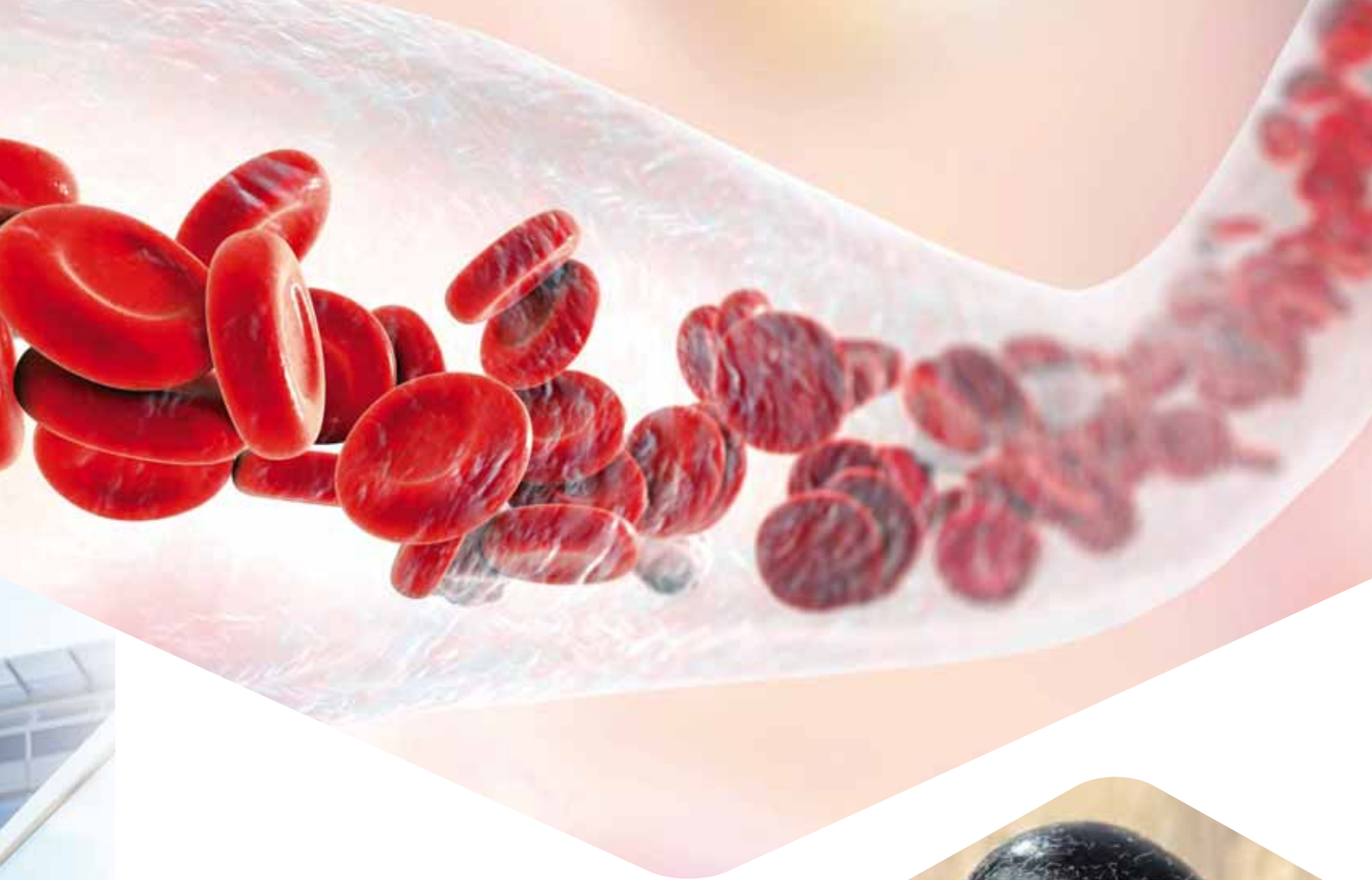
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Nitric Oxide Improves Blood Flow





BY CHANCELLOR FALOON

Nitric oxide plays a major role in the *dilation* of blood vessels.

It is vital for maintaining healthy **blood flow**,¹ **blood pressure**,² and **platelet function**.¹

The problem is that internal **nitric oxide** production *decreases* with age.^{3,4}

Low nitric oxide is associated with an increased risk of **cardiovascular diseases**,⁴ **cognitive decline**, and **dementia**.^{4,5}

Nitric oxide has a short half-life. It gets metabolized and quickly eliminated from the bloodstream.⁶

Aronia berry and a long-acting form of **L-arginine** each work to boost and sustain *higher nitric oxide* production.



ARONIA BERRY



Importance of Nitric Oxide

Nitric oxide is produced by nearly every cell in the body.^{7,8} It's needed to dilate **blood vessels**, allowing them to open up to increase blood flow.⁹

When we exercise^{9,10} or travel to higher altitudes,¹¹ our bodies release *more* nitric oxide to relieve blood vessel constriction.¹⁰

Nitric oxide is also produced in our nasal cavity when we breathe through our nose. This helps to combat **viruses** and **bacteria**.^{12,13}

The 1998 **Nobel Prize in Physiology or Medicine** was awarded to scientists who discovered its role in maintaining **cardiovascular health**.^{14,15}

With age, nitric oxide levels decline. This can result in **endothelial dysfunction**, when these cells lining the inner walls of arteries don't work properly.⁴ Blood vessels can't widen when needed, *reducing* blood flow.

That can lead to high blood pressure, atherosclerosis, abnormal clotting, and increased risk of **heart attacks**, **strokes**, and **sudden cardiac death**.

Endothelial dysfunction also increases the risk of **dementia** and **cognitive dysfunction**.^{3,16,17}

Long-Acting Form of L-Arginine

For the body to produce nitric oxide, no compound is more important than the amino acid **L-arginine**. It is the direct **precursor** nutrient that blood vessels use to make **nitric oxide**.

In a clinical trial, patients with high blood pressure were given either a placebo or a single dose of **L-arginine**. Measurements were taken of **flow-mediated dilation**, how much a blood vessel **dilates** (widens) in response to an increase in blood flow.

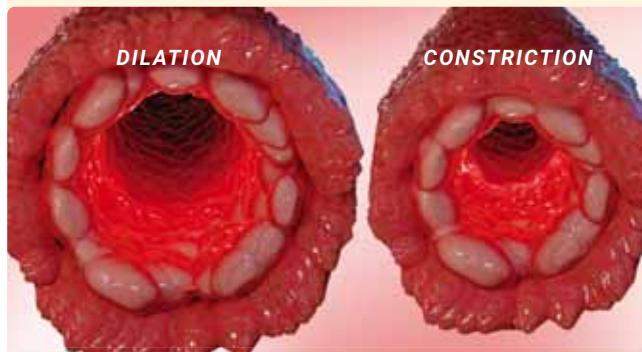
Participants who received the placebo had no change. But those given **L-arginine** had an average improvement in flow-mediated dilation from **1.7%** to **5.9%**.¹⁸

Considering that each **1%** improvement in flow-mediated dilation correlates with a **12%** lower risk of cardiovascular events, this suggests a potential **50%** reduction in risk for cardiovascular events.^{19,20}

A drawback with L-arginine has been that it takes about an hour to take effect and it does not stay in the bloodstream for long.²¹

But scientists have developed a more **bioavailable** (more enters circulation and can have an effect) form of L-arginine called **inositol-stabilized arginine silicate**. It works within just **15 minutes** and sustains L-arginine levels for up to **six hours**.^{21,22}

Flow-Mediated Dilation



Flow-mediated dilation is the change in diameter of blood vessels in response to increased blood flow (such as exercise or high blood pressure).

It can assess vascular function and cardiovascular risk in an individual.

Research indicates that for every **1%** increase in brachial artery flow-mediated dilation, the risk of cardiovascular events is reduced by **0.87%**.

Inositol-stabilized arginine silicate appears to keep L-arginine levels *higher* for longer periods because it inhibits the enzyme **arginase**, which breaks down L-arginine.²³

This stabilized L-arginine form has demonstrated clinical benefits that standard L-arginine hasn't been shown to achieve.

In three recent randomized, controlled trials, **inositol-stabilized arginine silicate** improved working memory, processing speed, concentration, and other measurements of **cognition** in young adults.²⁴⁻²⁶

Effects of Aronia Berry

Aronia berries are native to North America and resemble cranberries. They have been thought of as a superfood due to their high content of vitamin C, **anthocyanins**, and other polyphenols.²⁷

WHAT
YOU
NEED
TO
KNOW

Keeping Blood Vessels Healthy

Research indicates that low flow-mediated dilation of blood vessels is associated with a high risk of cardiovascular events.²⁰ Research also suggests that **aronia berries** boost nitric oxide production by increasing the activation of an *enzyme* that converts **L-arginine** into endothelial **nitric oxide**.²⁸

In a randomized, controlled trial, a daily intake of **500 mg** of aronia whole fruit berry and extract powder for 12 weeks resulted in a **1.2%** improvement in **flow-mediated dilation** compared to placebo.²⁹ This corresponds to a nearly **11% reduction** in the risk for cardiovascular events.¹⁹

In another clinical trial, 101 adults aged 40-60 years old were randomized to receive aronia berry extract or a placebo for **24 weeks**.

Cognitive function was assessed using tests including the **grooved pegboard test**, in which pegs must be precisely rotated to match a slot before they can be inserted.³⁰ This measures **psychomotor speed**, the ability to quickly think and then perform a motor action.

The results showed that those who received **aronia berry extract** daily had significantly *higher* scores on the test after just **six weeks** than those taking a placebo.

Taking **aronia** extract along with the new form of **L-arginine** may lead to even greater levels of nitric oxide production, maximizing the cardiovascular and cognitive benefits.

- The endothelium is a layer of cells lining the inside of blood vessels. It produces **nitric oxide**, a molecule that signals the vessel to dilate, allowing blood to flow through.
- With age, nitric oxide production tends to decline, contributing to **endothelial dysfunction**. This can increase the risk of heart attacks, strokes, cognitive decline, and dementia.
- **L-arginine** is a precursor the body needs to make nitric oxide. A new form, **inositol-stabilized arginine silicate**, improves its bioavailability and supports higher levels in the bloodstream for a longer time.
- **Aronia berries** help activate the enzyme that converts L-arginine into nitric oxide.
- Taking oral L-arginine improves blood vessel dilation enough to correlate with a **50%** reduction in risk for **cardiovascular events**. It also improves several measures of cognition.
- In a clinical study, **aronia berry extract** also improved blood vessel dilation and significantly boosted scores on a test of cognitive function.
- Taking these two ingredients **together** may significantly boost nitric oxide production, reducing the risk for cardiovascular events and enhancing cognitive function.



Summary

Nitric oxide is needed to maintain the health of blood vessels. As production in the body decreases with age, the risk of **cardiovascular diseases** and **cognitive deficits** rises.

L-arginine and **aronia berry extract** have each been shown to *increase* nitric oxide production.

Research suggests that aronia berry increases the activation of an *enzyme* that converts the amino acid L-arginine into nitric oxide.

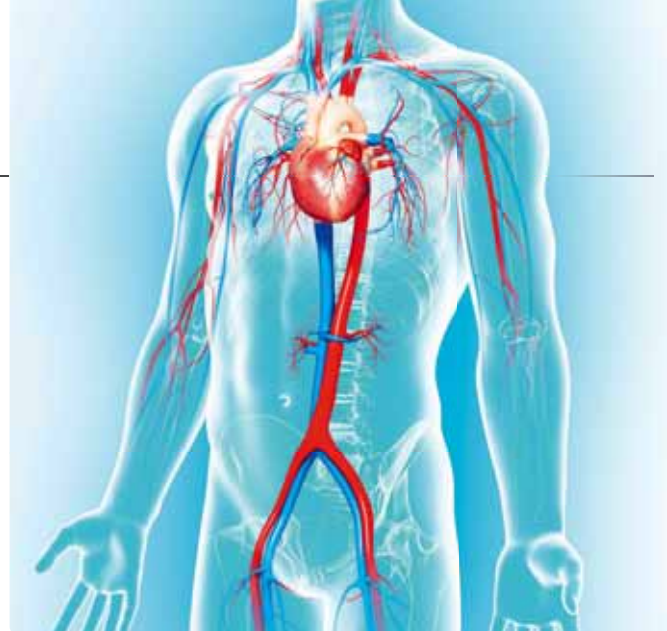
Clinical trials have shown that taking each has benefits for both **cardiovascular** and **cognitive** health.

Taking both together may result in ideal levels of nitric oxide production in the body for aging individuals. ●

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

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Fish oil is well known to support **heart**¹ and **brain** health.^{1,2}

Few know that **human** population studies show a correlation between **fish oil** and **longer lifespans**.¹⁻³

Thanks to a novel technology, **Omega-3 Fish Oil Gummy Bites** are available that melt in your mouth.

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These tasty **gummy bites** can serve as an alternative or addition to fish oil softgels.

These **new Omega-3 Fish Oil Gummy Bites** pack the same **EPA/DHA** into a concentrated form to provide a small and delicious **tropical fruit-flavored sugar-free* gummy bite**.⁴ No fishy aftertaste.

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36 gummy bites

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* *Not a low-calorie food.*



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- Hundreds of published studies describe **resveratrol's** health and longevity potential.¹
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Optimized Resveratrol Elite™ provides **bioavailable resveratrol** plus highly **absorbable quercetin** to provide complementary biological functions.

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Enhance Arginine to Boost NITRIC OXIDE



"It does give me that
extra boost."

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When L-arginine is ingested, about 40% is degraded in the digestive tract by the *arginase* enzyme.

After absorption, arginine encounters further degradation, leaving little for conversion into nitric oxide.

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NitroVasc™ provides a combination of inositol-stabilized L-arginine silicate and aronia berry extract.

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Fish Oil

Super Omega-3 provides components found in **Mediterranean diets**, including **sesame lignans** to enhance the health benefits of fish oil.



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SUPER OMEGA-3
EPA/DHA Fish Oil,
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IFOSTM certification mark is a registered trademark of Nutrasource Diagnostics, Inc. These products have been tested to the quality and purity standards of the IFOSTM program conducted at Nutrasource Diagnostics, Inc.

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

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.

What is D-Ribose?

BY LAURIE MATHENA



Normal aging results in a decline in muscle mass and energy production.^{1,2}

This energy *deficit* has an impact on heart health, cognitive function, and lifespan.²

D-ribose helps restore **energy** production in the body.³

It is a building block of adenosine triphosphate (ATP), the energy source for every cell.⁴

By supporting the production of ATP, **D-ribose** can help replenish the metabolic energy needed by all cells, including those in major organs such as the **heart** and **brain**.³

ATP, Energy, and D-Ribose

Energy in the body is produced in the form of adenosine triphosphate, ATP.

This takes place in mitochondria (powerhouse in each cell). ATP is the primary energy source for most biochemical and physiological processes, such as growth, movement and homeostasis.⁵

Mitochondrial function declines with age^{5,6} and other health conditions, such as heart failure, among others.³ This results in loss of ATP production and decreased energy levels.³

Hope for Heart Failure

Heart failure means the heart muscle is failing to pump enough blood to meet the body's metabolic requirements.

In heart failure, D-ribose production falls in heart muscle cells.^{3,7}

This leads to a *decrease* in **ATP** production, resulting in cellular energy deficiency in the muscle cells of the organs that need energy the most.⁷

Taking **oral D-ribose** can help create new ATP molecules and *restore* cardiac energy levels.^{7,8}

Clinical trials have shown that D-ribose taken orally can **improve heart function** in heart failure patients.⁹⁻¹¹



Lifestyle Modifications to Fight Fatigue¹⁸

A constant feeling of tiredness or weakness is called fatigue. It can affect anyone, and most adults will experience fatigue at some point in their life. Some lifestyle modifications that may help:

- Exercise: even a 15-minute walk can give you an energy boost.
- Lose weight if overweight.
- Get optimal sleep.
- Relieve stress: meditate, work out, do yoga, listen to music, have social support.
- Limit alcohol and caffeine intake.
- Stay hydrated.

People with heart failure taking D-ribose were shown to improve **blood flow** through the heart and body *and* boost the exchange of oxygen and CO₂ through the lungs, leading to improvements in breathing parameters.

Another study showed the ability of D-ribose to reduce symptoms and improve quality of life in **heart failure** patients.¹⁰

In a review of studies in animals and humans, D-ribose has been shown to increase **ATP production** in heart muscle cells and improve cardiac function.

In clinical trials D-ribose enhanced cardiac function and improved quality of life in patients with heart failure.⁷

Fibromyalgia

There is evidence that **defective production of ATP** is the one potential culprit behind **fibromyalgia** (a condition that causes pain throughout the body)¹² and chronic fatigue syndrome.^{13,14}

In an open-label, early study, patients with fibromyalgia or chronic fatigue syndrome took **5 grams** of **D-ribose** three times daily until they reached a total of **280 grams**.¹⁴

The participants reported significant improvement in all five categories on a standard questionnaire: energy, sleep, mental clarity, pain intensity, and well-being.

On average, patients reported a stunning 45% increase in self-reported energy levels.

Restless Leg Syndrome

Restless leg syndrome is a disorder causing discomfort and pain in the legs. This condition progresses with age and often leads to insomnia. Disordered energy metabolism has been suggested as one possible cause of **restless leg syndrome**.

Based on that observation, researchers gave individuals with restless leg syndrome **5 grams** of **D-ribose**, three times per day. Remarkably, daytime symptoms were **completely eliminated**, and nighttime symptoms were *significantly reduced*.¹⁵

Exercise Performance

D-ribose is a building block of **ATP**.³ It may help speed muscle recovery after high-intensity exercise.

- In a double-blind cross-over study of 26 athletes, subjects were given either a dextrose sugar control or **10 grams** of D-ribose for two days. This was followed by three additional days of supplementation. During these three days, both groups underwent 60 minutes per day of high-intensity exercise. After five days, significant improvement in exercise performance and lower perceived exercise exertion were observed in the D-ribose group, compared to the **placebo** arm of the study.¹⁶
- In a study of healthy, active individuals, supplying fatigued muscle cells with D-ribose quickly restored **ATP levels** to normal.¹⁷

Summary

By restoring the body's ability to produce energy, **D-ribose** leads to improved function for organs such as the heart and muscles.

D-ribose intake is especially valuable for **heart failure** patients and has been shown to produce meaningful improvements in cardiovascular function.

Because **high doses of D-ribose** are needed, most people find it more efficient to take **5 grams** or more each day in a neutral-tasting **powder** form. •

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

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What is ATP (adenosine triphosphate)?^{3,4}

- ATP is a molecule carrying energy, found in every cell of the body; it is vital for energy production.
- It has a nitrogenous base (adenine), and a sugar (ribose), attached to three phosphate molecules.
- Cells need energy to perform cellular functions, such as growth, nerve impulse propagation, and muscle cell contraction.
- The energy is stored in phosphate bonds of ATP and is released when these bonds are broken by chemical processes.



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VITAMIN



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Mary

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Buffered **ascorbate** encased in two plant extracts (liposomes plus hydrogel fenugreek) increases blood (plasma) exposure nearly **seven times more** compared to an equivalent dose of regular vitamin C.

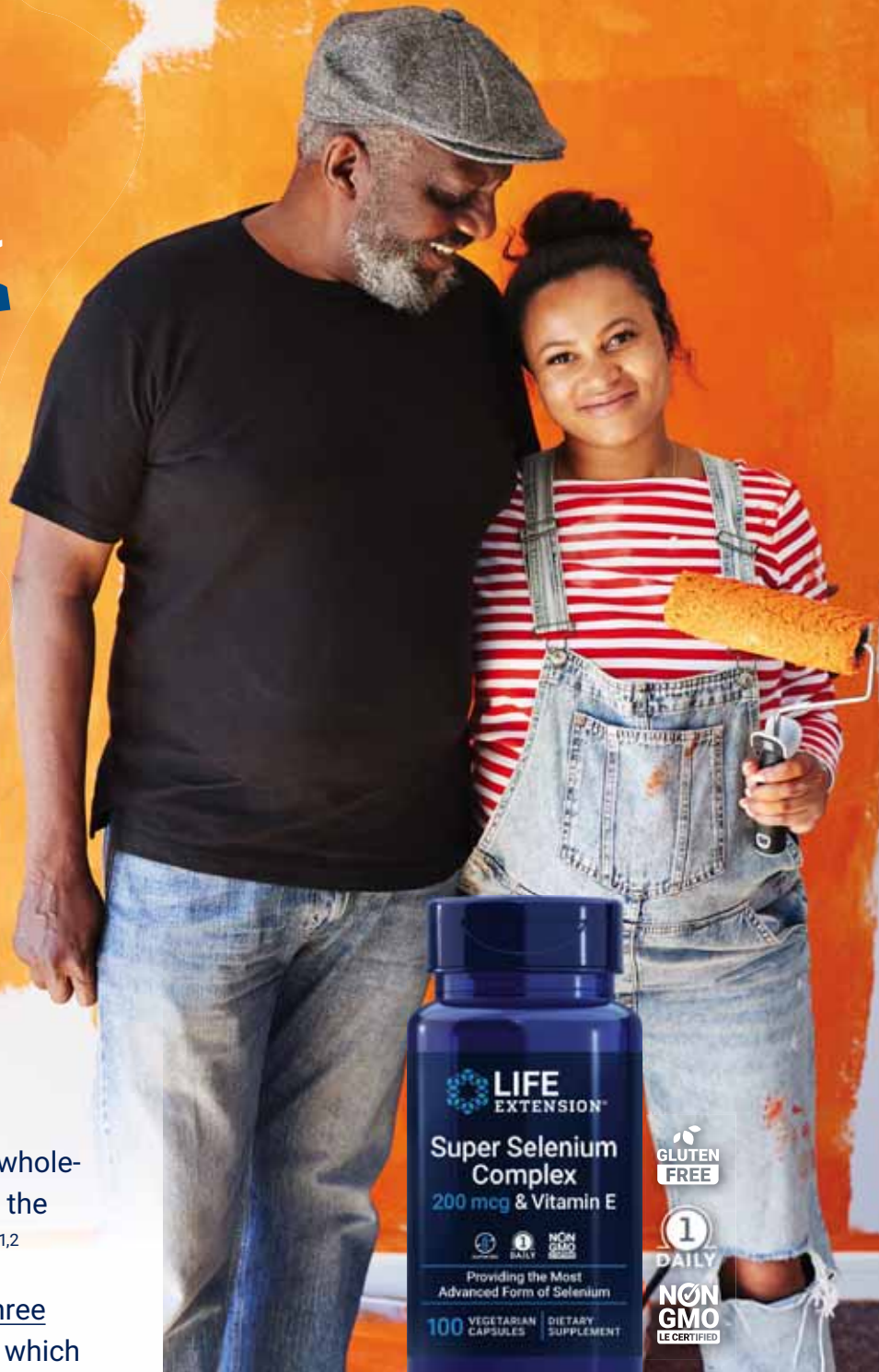
It also maintains *higher* vitamin levels throughout the day.¹

Just one vegetarian tablet daily provides **around-the-clock** vitamin C support.

1. Akay Internal Study. Liposomal hydrogel vitamin C pharmacokinetics. Data on file. 2021.

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Selenium has been shown to confer whole-body protection including supporting the heart, brain, and healthy cell division.^{1,2}

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2. **L-selenomethionine**
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References

1. *Biol Trace Elem Res.* 2004 Oct;101(1):73-86.
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Item #01778 • 100 vegetarian capsules

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Each bottle provides a supply that lasts more than three months.

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Rick

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"Great for energy."

Loni

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100 vegetarian tablets
1 bottle \$24
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Item #00972
150 grams powder (.33lb. or 5.29 oz)
1 jar \$27
4 jars \$24 each

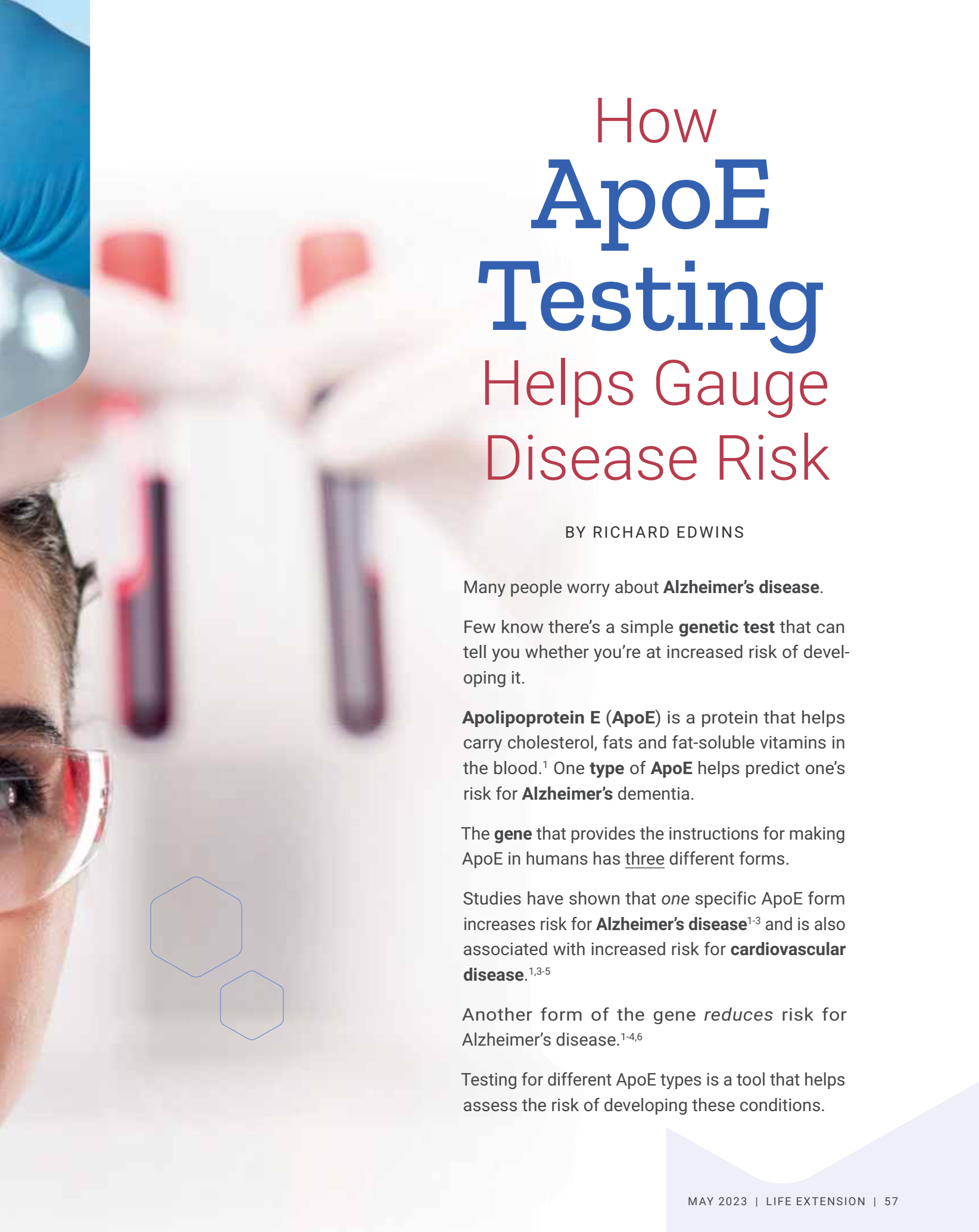
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How ApoE Testing Helps Gauge Disease Risk

BY RICHARD EDWINS

Many people worry about **Alzheimer's disease**.

Few know there's a simple **genetic test** that can tell you whether you're at increased risk of developing it.

Apolipoprotein E (ApoE) is a protein that helps carry cholesterol, fats and fat-soluble vitamins in the blood.¹ One **type** of **ApoE** helps predict one's risk for **Alzheimer's** dementia.

The **gene** that provides the instructions for making ApoE in humans has three different forms.

Studies have shown that *one* specific ApoE form increases risk for **Alzheimer's disease**¹⁻³ and is also associated with increased risk for **cardiovascular disease**.^{1,3-5}

Another form of the gene *reduces* risk for Alzheimer's disease.^{1-4,6}

Testing for different ApoE types is a tool that helps assess the risk of developing these conditions.

ApoE Types

The ApoE gene exists in three different forms:²

- **ApoE e3** is the most common type. About **60%** of the population has two copies of **ApoE e3**, one from each parent.
- **ApoE e4** is less common, and it is the problematic type. About **20%-30%** of people have one or two copies of this form.
- **ApoE e2** is the least common form. Only about **10%-20%** of people carry at least one copy of this gene.



Who Should Test?

The problematic **ApoE e4** form is present in approximately **25%** of the U.S. population.⁹ Genetic testing for ApoE status can be done with a simple cheek swab from home.

ApoE status does not tell you if you will or will not develop Alzheimer's disease or other conditions, only whether you are at **greater risk** of having them.

If you are concerned about your risk of Alzheimer's or cardiovascular disease or have a family history of either, consider testing your ApoE status, the sooner the better, so preventative measures can be taken if ApoE e4 is detected.

Your physician can also discuss how you can use the test results to make lifestyle changes to minimize your overall risk.

ApoE Functions

ApoE is mainly synthesized in the liver, and it helps transport lipids, fat-soluble vitamins, and cholesterol into the lymph system and then into the blood.

In the brain, ApoE is mainly produced by **astrocytes** that help transport essential lipids and maintain healthy neuronal signaling.⁷

The three forms of ApoE have different lipid-binding capacities that involve their abilities to bind or clear **amyloid beta plaques** that form around nerve cells.^{1,8,9}

ApoE genes function to help control brain inflammation,^{7,9} and support removal of degenerated cell membranes and lipids that accumulate with aging.⁸

ApoE binds to LDL receptors in the liver to remove remnant lipoprotein particles (chylomicrons and VLDL) from circulation. Defects in ApoE result in an increase in plasma cholesterol and triglycerides due to impaired clearance of chylomicrons, VLDL and LDL.¹⁰

One distinct feature of Alzheimer's disease is the build-up of clumps of beta-amyloid in the brain. **ApoE** enhances beta-amyloid break-down, both within and between cells.

Unfortunately, the **ApoE e4** genetic variant is not as effective as the others at breaking down the beta-amyloid clusters, controlling brain inflammation, and protecting brain and nerve cells.¹¹

Alzheimer's Risk

In population studies, the **ApoE** genetic type correlates with risk for **Alzheimer's disease** in the following ways:

- **ApoE e4:** The presence of even *one* copy of the **e4** form of the gene increases the risk of developing Alzheimer's disease.^{2,3,6} Typical estimates suggest one **e4** gene increases the risk of Alzheimer's disease **two to three fold**, whereas having **two** copies (one from each parent) is associated with **8 to 12 fold** greater risk than average.¹²
- **ApoE e3:** This gene form is not considered to have any impact on risk for Alzheimer's disease.
- **ApoE e2:** The rarest form of the gene has been found to be protective, reducing risk of developing Alzheimer's disease.¹⁻³

A study published in **2022** found that **ApoE e4** is linked to faulty cholesterol processing in the brain, which leads to defects in the protective coating around nerve fibers. That, in turn, may cause memory problems.¹³



WHAT
YOU
NEED
TO
KNOW

What ApoE Genetic Testing Can Tell You

- **Apolipoprotein E (ApoE)** is a protein that helps carry lipids, fat soluble vitamins and cholesterol in the blood.
- Three forms of the gene for ApoE exist in humans: e4, e3, and e2.
- The **ApoE e4** form is associated with an *increased* risk of developing **Alzheimer's disease**.
- The rare ApoE **e2** form is associated with a *reduced* Alzheimer's risk.
- The most common form of the gene, ApoE **e3**, does not appear to have an impact on risk.
- The ApoE **e4** form is also associated with increased risk for other conditions, including **cardiovascular disease**.
- **Genetic testing** can identify which form of the ApoE gene people have.
- Those at higher risk for these conditions can consult with a doctor about making lifestyle changes to minimize overall risk.

Having the **e4 gene** does not mean someone *will* develop Alzheimer's disease. But it does mean there's an increased risk of developing it. It also means there's a greater risk of developing it at a **younger age**, and of having a **faster decline** in cognitive function as it progresses.^{2,6}

A **genetic test** can identify which form of the ApoE gene a person has. Those who discover they're at *higher* risk can make changes to reduce *other risk factors* for Alzheimer's disease.¹

Other Health Outcomes

ApoE status is associated with other health conditions.

Most notably, having the **ApoE e4** gene form is associated with *higher* cholesterol levels and increased risk of cardiovascular diseases.^{1,3,5,10} It plays a role in glucose transport, neuronal signaling, and mitochondrial function. Independently, **ApoE e4** may trigger inflammatory response causing neurovascular dysfunction.⁷

Genetic testing can identify those at higher risk for these conditions, allowing them to monitor their health and make changes that could help ward off disease.

Summary

Genetic testing to see which form of the **apolipoprotein E (ApoE)** gene you have can help assess your risk for certain chronic diseases, particularly **Alzheimer's disease**.

Having the **ApoE e4** form of the gene is considered a risk factor for Alzheimer's and heart disease. On the other hand, possessing a copy of the **e2** form indicates reduced risk of Alzheimer's disease.

Genetic testing can allow people at greatest risk for developing Alzheimer's and cardiovascular diseases to make healthy lifestyle choices to reduce the dangers. •

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

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What Dietary and Lifestyle Changes Can Be Beneficial for Alzheimer's Disease? ¹⁴

- The Mediterranean diet has been linked to a reduced risk of Alzheimer's and other neurodegenerative diseases. ¹⁴⁻¹⁶
- Aggressive dietary interventions demonstrated to be efficacious in reversing early Alzheimer's dementia include eliminating red meat, processed foods, simple sugars and excess carbohydrates while increasing intake of healthy plant foods, fiber, overnight fasting, and exercise. ^{17,18}
- Low-calorie diets have been linked to a reduced risk of cognitive decline. ¹⁵
- Avoiding smoking and protecting against hypertension, diabetes, dyslipidemia, and obesity, may help in the prevention of dementia. ¹⁵
- Regular exercise may directly benefit brain cells by increasing blood and oxygen flow in the brain. ¹⁴
- Limiting alcohol consumption. ¹⁶
- Engaging in late-life cognitive activities—being intellectually engaged by keeping the mind active—may benefit the brain. ¹⁶
- Supplement the brain with basic cognitive support—nutrients such as omega-3 fatty acids, curcumin, resveratrol, and magnesium. ¹⁵

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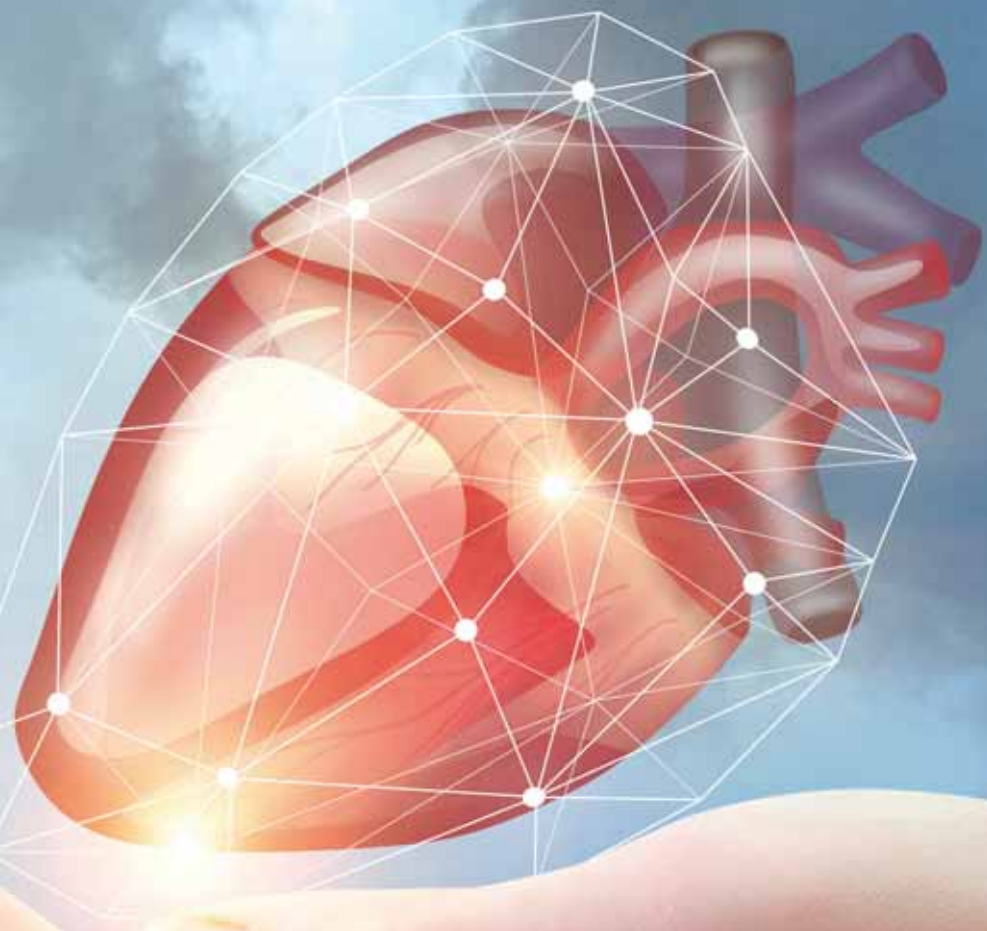
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KATIE COURIC

A Tireless Advocate for Colon Cancer Awareness

BY LAURIE MATHENA

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From 1991-2006, Katie Couric was the face of morning television as the co-host of the *TODAY* show on NBC.

Today, she has become the face of something even more important: **colon cancer awareness**.

In 1998, colon cancer changed Couric's life forever when it claimed the life of her husband, Jay Monahan, at just **42** years old.

He was not diagnosed until it was in the advanced stages, and he died nine short months after.

Couric learned that **routine colon screening** could have saved her husband's life, and for

the past 20-plus years, she has devoted her life to spreading awareness, promoting routine screening, and raising money for cancer research.

More recently, due to an alarming increase in colorectal cancer cases in people under 50, the recommendations for routine screening have been lowered from age 50 down to 45.

Now, Couric is stepping out in new ways to promote screening to a younger generation.

Spreading Awareness

In Couric's memoir, *Going There*, she gets deeply personal about the anger, frustration, hurt, and loss she experienced after her husband's diagnosis and untimely death.

She laments the signs she overlooked—the sensitive stomach, the fatigue, the weight loss.

She talks about her search for a cure, her bargains with God, and the regret she had that she and her young daughters (ages five and one at the time of Jay's death) wouldn't have more time with their father.

Ultimately, she determined to honor Jay's memory by becoming an advocate for cancer screening.

In 2000, Couric appeared before the U.S. Senate Select Committee on Aging in an effort to highlight the dangers of colon cancer and the importance of colon cancer screening.

"During this terrible struggle, I got a quick and painful education about this devastating disease. I learned that colon cancer is the second leading [cancer] killer. It kills more people than any other cancer,

with the exception of lung cancer,” Couric said in testimony before the Committee. “But I also learned that it has a **90%** or better cure rate if detected early. That means that colon cancer screening is a critical weapon in the fight against a disease no one needs to die from.”

That same year, she co-founded the National Colorectal Cancer Research Alliance (NCCRA) with anti-cancer activist Lilly Tartikoff and the Entertainment Industry Foundation.

The goal of the NCCRA was to fund research to develop better tests, treatments, and ultimately a cure—as well as to promote the life-saving value of screening.

Nothing brought more awareness to the value of screening than when Couric herself underwent a colonoscopy live on the *TODAY* show.

“Of everything I’ve done in my career, here is the thing of which I’m most proud: Studying the impact a public figure can have on health

issues, the University of Michigan reported a **20%** jump in colonoscopy screenings as a result of my airing the procedure. They called it the Couric Effect,” she said in *Going There*. “But even more gratifying was unsolicited feedback from [my daughter] Ellie when she was just nine. ‘Mom, I’m so proud of the work you’re doing with colon cancer.’”

New Risks for a New Generation

Screening is critical because colorectal cancer typically doesn’t cause symptoms until it has already grown or spread, according to the American Cancer Society.

By the time symptoms appear—such as a change in bowel habits, blood in the stool, persistent abdominal discomfort, and unexplained weight loss—the cancer is often in advanced stages.

“Jay never really had any symptoms of colon cancer,” Couric said, “other than being tired all of the time, which we chalked up to a busy lifestyle, having young children, and his coast-to-coast trips providing legal analysis for NBC.”

This is why colon cancer continues to be one of the deadliest cancers—but it is also one of the most treatable when caught early with screening.

Colon cancer rates have been declining *overall* since the 1980s, no doubt due in part to Couric’s relentless campaigning for screening.

Unfortunately, during that same time period, there has been an alarming increase in younger adults.

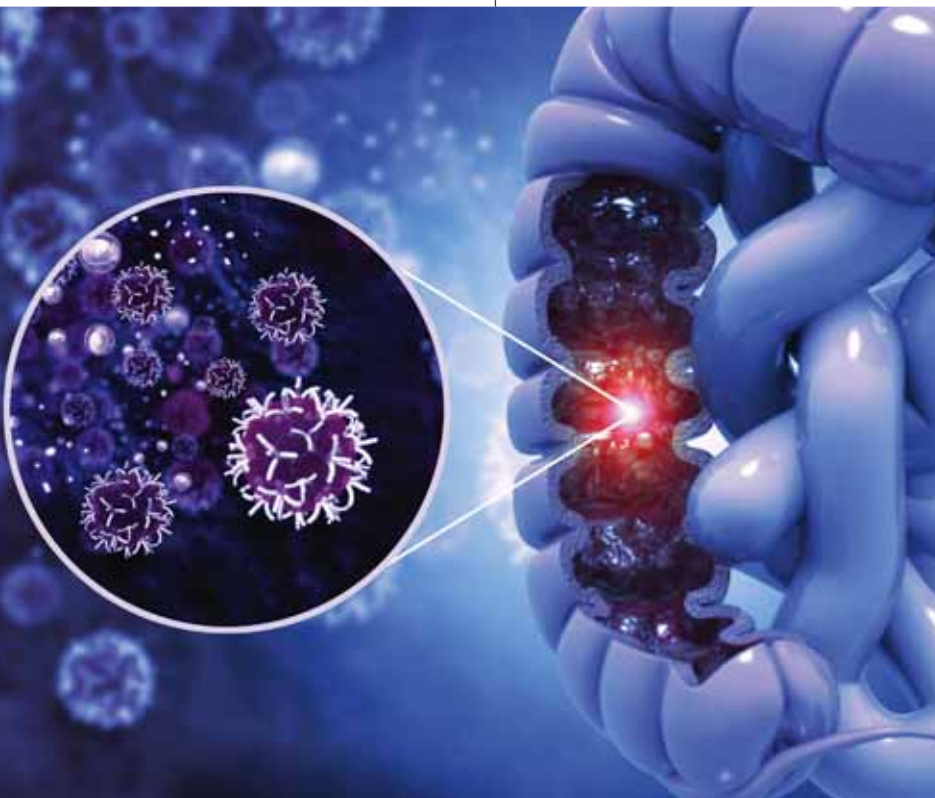
In adults ranging from 20-55 years of age, colon cancer incidence has been increasing by **1%-2%** per year. Rectal cancer rates are rising even faster, at **2%-3%** per year.

According to research conducted by the American Cancer Society, people born after **1990** have **double** the risk of colon cancer and **quadruple** the risk of rectal cancer, compared to those born in 1950.

These increases have led the **U.S. Preventive Services Task Force** to change their screening guidelines from **50 years** old down to **45 years** old.

This change makes **45 million Americans** above the age of **45** eligible for screening.

(NOTE: LifeExtension® has always advocated for colonoscopy screening to begin at age 40. We did not agree with conventional medicine’s guidelines to begin screening at age 50 nor do we concur with U.S. Preventive Services Health Task Force’s new recommendation to begin screening at age 45 instead of age 50.)



Colon Cancer Screening at Home

With a new generation facing alarmingly increased rates of colorectal cancer, Couric is on a renewed mission to make the younger generation aware of the importance of early screening.

“Screening wouldn’t have crossed our minds for Jay when he was diagnosed at 41,” said Couric. “My hope is that we figure out a way to do even broader screening and that we can save even more lives. But until then, people need to pay attention, they need to talk to their doctors... And they need to get screened.”

Couric, along with one of the organizations she co-founded, Stand Up To Cancer, is now promoting the at-home screening test, Cologuard®, in a national TV commercial.

This at-home test detects altered DNA or blood in stools and is **92%** accurate at detecting cancers. However, it only detects about **42%** of precancerous polyps. It also had a 13% false positive rate.

While this could be a stepping-stone for increasing awareness and compliance (due to its less invasive nature), a traditional **colonoscopy** remains the gold standard of testing.

Colonoscopies are more than **90%** effective at identifying colon cancer and the presence of precancerous polyps that can turn into cancers.

One study in the *British Medical Journal* found that getting a colonoscopy was associated with a **67%** reduced risk of dying from colorectal cancer.

In a recent interview with *TODAY*, Couric said, “The bottom line is that [early] screening saves lives. Early detection saves lives.”

ALARMING TREND

While colon cancer death rates have fallen in the general population, largely due to better screening, it has risen in younger people.¹

According to the American Cancer Society, **20%** of new colon cancer diagnoses are people under 55. In 2023, it is estimated that there will be **19,500** new cases in people under 50. To make matters worse, these cases in younger people are being diagnosed at a more advanced stage of the disease.² Doctors are perplexed by this alarming trend.

One answer is obesity and high body mass index. A recent meta-analysis found **30%** increased risk of colon cancer in males and **12%** in females with each 5-unit increase in body mass index.^{3,4}

And obesity is increasingly prevalent in the young.⁵

More screening is needed at a younger age along with diet and lifestyle modifications. Screening recommendations by the American Cancer Society includes annual stool tests followed by a colonoscopy every five years.²

A Legacy to Be Proud Of

Beyond promoting early detection through screening, Couric has poured her efforts into finding a cure.

In 2008, she cofounded Stand Up To Cancer, a program whose mission is to raise funds to accelerate the pace of groundbreaking translational research that can get new therapies to patients quickly and save lives *now*.

Couric says that, to date, they’ve raised over **\$600 million**, and have helped gain FDA approval for nine new cancer drugs.

“I don’t like to think about my obituary too often,” said Couric. “But when I do, I hope the first line will be ‘Katie Couric was a tireless advocate for cancer awareness and research.’” •

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

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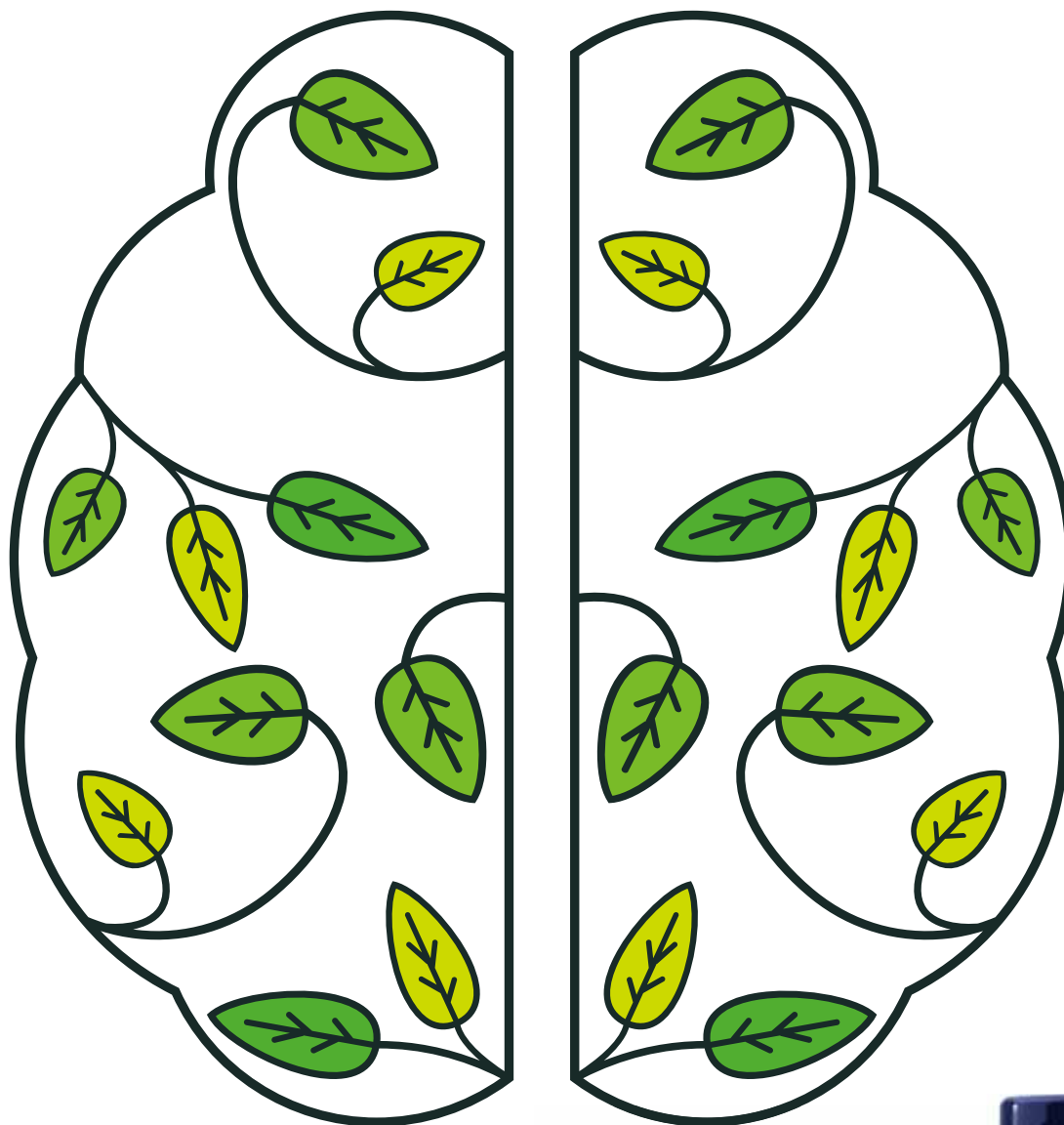
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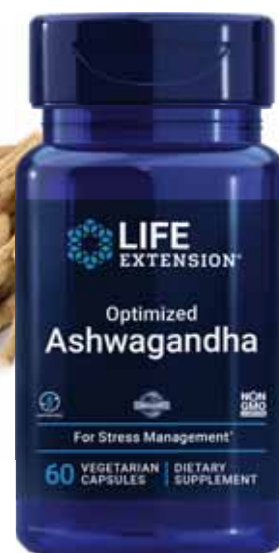
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Astaxanthin and the Liver

BY LISA STUART



Chronic liver disease is reported as one of the **15** leading causes of death in the U.S.

In the year **2020** it contributed to over **50,000** fatalities.¹

These liver diseases include **nonalcoholic fatty liver disease**, alcoholic liver disease, viral hepatitis, and other toxic conditions.

A large proportion of the population is at risk of long-term liver damage.

Researchers are intrigued by animal, preclinical and human data about the carotenoid **astaxanthin** and its potential ability to protect the liver in multiple ways.²⁻⁵

Studies show that this plant carotenoid can help defend the liver against injury and may help **reverse damage** that has already been done.^{2,3}

Astaxanthin has been studied for potential clinical benefits, including for eye,⁶⁻⁸ heart,⁸⁻¹⁰ and brain health.^{8,11}

Protecting the liver is one more area where astaxanthin may prove valuable.²



Astaxanthin

Astaxanthin is a red carotenoid produced by marine microorganisms such as microalgae and phytoplankton. It is responsible for the pinkish color of shrimp, salmon, and flamingos, as a result of how much astaxanthin they consume.¹⁰

Astaxanthin is an anti-**inflammatory** and **antioxidant**.¹⁰

More recent research has found that astaxanthin protects against various forms of injury that can cause **chronic liver disease**.^{2,3}

Role in Liver Fibrosis

Several factors can cause **liver damage**, including poor diet, metabolic disease, alcohol consumption, hepatitis viruses, and others. These lead to a cycle of chronic inflammation, and tissue destruction that over time can cause **liver fibrosis** (scarring).¹²

In the late stages of liver disease, this scarring becomes permanent. When large sections of the liver are replaced by scar tissue and can no longer function, it is known as cirrhosis. The end-stage result is **liver failure**. At this point, the only effective treatment is a liver transplant.¹²

Several *in vivo* and *in vitro* studies have found that **astaxanthin** is protective against **fibrosis**.^{2,3,13,14}

With fibrosis, a certain type of liver cell becomes activated and begins synthesizing fibrous tissue. In animal models and human liver cells, astaxanthin **prevents and reverses** this process, blocking the activation of these cells *and* shutting down cells that have already been activated.^{15,16}

The chemical carbon tetrachloride is a powerful **liver toxin**. It causes severe liver damage, leading to fibrosis.¹⁷

In an animal trial, with **liver damage** induced by this liver toxin, rats were given **astaxanthin** at a dose of **10 mg/kg**. After two weeks it was observed that the liver in the treatment group was protected from oxidative damage induced by the toxin, with a significant reduction of inflammation and fibrosis.¹⁷

Role in Fatty Liver Disease

A condition called **nonalcoholic fatty liver disease** causes the liver to accumulate **fat**, resulting in inflammation and damage that can lead to liver fibrosis and loss of function. More than **20%** of adults in the U.S. and North America may suffer from this condition.¹⁸

Preclinical studies show that **astaxanthin** may help to prevent or even reverse it.

Several rodent model studies have shown that astaxanthin can **prevent and reverse** the development of fatty liver changes. It also improved markers of metabolism and **insulin sensitivity**.^{5,13,14,19-21}

One group of scientists studying different models of **fatty liver disease** in mice found that giving them **astaxanthin**.⁵

- Prevented fatty liver changes,
- Reversed existing changes, decreasing liver fat,
- Reduced inflammation and fibrosis, and
- Improved insulin sensitivity.

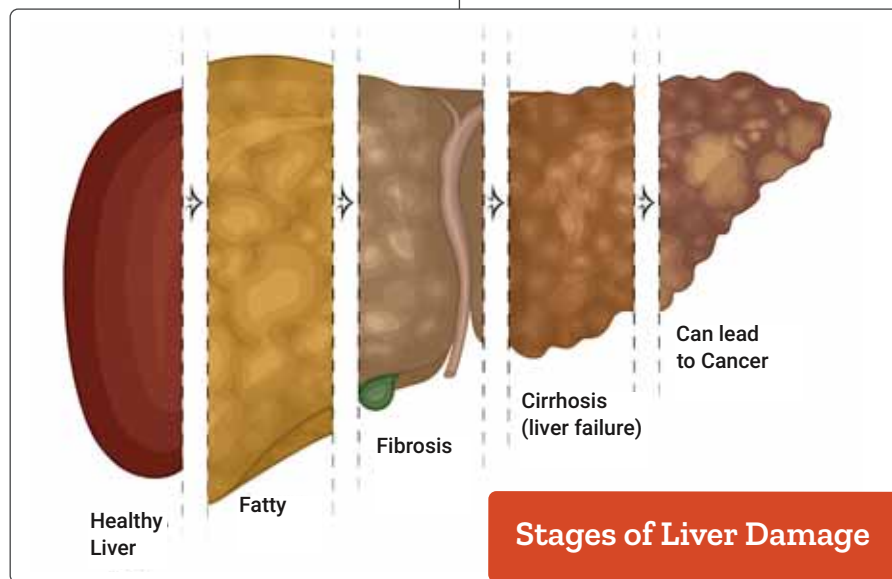
The same research group performed a preliminary study in **humans** with existing **fatty liver**. Subjects were randomized to receive either **12 mg** of **astaxanthin** per day for 24 weeks or a placebo. Compared to the **placebo**, astaxanthin was able to reduce liver fat and **slow the progression** of fatty liver.⁵

Several studies have illustrated some ways that astaxanthin can protect the liver.² In addition to shielding against oxidative damage and inflammation, it inhibits the signaling proteins **NF-κB** and **PPARs**, which are both drivers of liver inflammation and formation of fatty deposits. These effects can even help decompose existing fat droplets in the liver.^{20,21}

Other Liver Benefits

Preclinical studies have found that astaxanthin administration holds promise for a wide range of liver disorders.²

- One notable example is **alcoholic liver disease**—multiple changes that are normally seen in the liver with excessive **alcohol** intake—including inflammation, fatty change, fibrosis, and mitochondrial dysfunction.



Excessive alcohol intake over time is one of the most common causes of cirrhosis and liver failure. Several animal model studies have shown that **astaxanthin** protects against alcohol-induced liver damage.²²⁻²⁷ It was shown in a mouse model that by improving gut health and reducing inflammation and oxidative stress caused by alcohol, astaxanthin guarded against liver damage.²⁶

- In a study of mice on a high-fat diet, astaxanthin inhibited the increases in body weight, and reduced liver weight, liver triglyceride, plasma triglyceride, and total cholesterol.²⁸ This suggests astaxanthin might be of value in reducing the likelihood of obesity and metabolic syndrome
- Even in a study of healthy older women, astaxanthin reduced liver enzymes in the blood, which can be markers of **liver damage**.⁴ This provides evidence that astaxanthin intake may protect the liver in people *with or without* any obvious signs of liver damage.

Conclusion

The carotenoid **astaxanthin** is a powerful antioxidant as well as an anti-inflammatory.

Preclinical models show that astaxanthin is protective of the **liver**, and some preliminary research in humans is also promising. Astaxanthin may guard against fatty liver changes, damage due to alcohol and other toxins, and liver fibrosis. •

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

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300 mcg, 100 veg tablets
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750 mcg, 60 veg tablets
- 01721 Optimized Tryptophan Plus
- 01444 Quiet Sleep Melatonin • 3 mg, 60 veg capsules
- 01445 Quiet Sleep Melatonin • 5 mg, 60 veg capsules
- 02502 Rest & Renew

VITAMINS

- 01533 Ascorbyl Palmitate
- 00920 Benfotiamine with Thiamine
- 01945 BioActive Complete B-Complex
- 00102 Biotin
- 00084 Buffered Vitamin C Powder
- 02229 Fast-C® and Bio-Quercetin Phytosome

- 02075 Gamma E Mixed Tocopherol Enhanced with Sesame Lignans
- 02070 Gamma E Mixed Tocopherol & Tocotrienols
- 01913 High Potency Optimized Folate
- 01674 Inositol Caps
- 02244 Liquid Vitamin D3 • 50 mcg (2000 IU)
- 02232 Liquid Vitamin D3 (Mint) • 50 mcg (2000 IU)
- 01936 Low-Dose Vitamin K2
- 00373 No Flush Niacin
- 01939 Optimized Folate (L-Methylfolate)
- 01217 Pyridoxal 5'-Phosphate Caps
- 01400 Super Absorbable Tocotrienols
- 02334 Super K
- 01863 Super Vitamin E
- 02422 Vegan Vitamin D3
- 02028 Vitamin B5 (Pantothenic Acid)
- 01535 Vitamin B6
- 00361 Vitamin B12 Methylcobalamin
- 01536 Vitamin B12 Methylcobalamin • 1 mg, 60 veg lozenges
- 01537 Vitamin B12 Methylcobalamin • 5 mg, 60 veg lozenges
- 02228 Vitamin C and Bio-Quercetin Phytosome • 60 veg tablets
- 02227 Vitamin C and Bio-Quercetin Phytosome • 250 veg tablets
- 01753 Vitamin D3 • 25 mcg (1000 IU), 90 softgels
- 01751 Vitamin D3 • 25 mcg (1000 IU), 250 softgels
- 01713 Vitamin D3 • 125 mcg (5000 IU), 60 softgels
- 01718 Vitamin D3 • 175 mcg (7000 IU), 60 softgels
- 01758 Vitamin D3 with Sea-Iodine™
- 02040 Vitamins D and K with Sea-Iodine™

WEIGHT MANAGEMENT & BODY COMPOSITION

- 02479 7-Keto® DHEA Metabolite • 100 mg, 60 veg capsules
- 02207 AMPK Metabolic Activator
- 02504 Body Trim and Appetite Control
- 02478 DHEA Complete
- 01738 Garcinia HCA
- 02506 Mediterranean Weight Management
- 01432 Optimized Saffron
- 00818 Super CLA Blend with Sesame Lignans
- 02511 Thermo Weight Control
- 02509 Waistline Control™

WOMEN'S HEALTH

- 01942 Breast Health Formula
- 01626 Enhanced Sex for Women 50+
- 01894 Estrogen for Women
- 02204 Menopause 731™
- 02319 Prenatal Advantage
- 01649 Super-Absorbable Soy Isoflavones
- 02513 Women's Bladder Support
- 02507 Youthful Woman 40+ with B-Complex

A Therapeutic Foot Massage with every step

Wearing Kenkohs daily, even for short periods, can help improve blood circulation, keep your body aligned, relieve pain in the feet, ankles, knees, legs, hips and back, reduce swelling, relieve stress and enhance your overall mood. Kenkoh revitalizes and rejuvenates your whole body!

"Massage Improves Circulation..."
UNIVERSITY OF MIAMI SCHOOL OF MEDICINE

"Massage aids muscle recovery and speeds recovery times..."

McMASTER UNIVERSITY, ONTARIO

"A daily foot massage lowers blood pressure and lowers triglyceride levels..."

PUSAN NATIONAL UNIVERSITY, SOUTH KOREA



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- 25 times the VITAMIN B6
- 10 times the BIOTIN
- 10 times the SELENIUM
- 8 times the VITAMIN C
- 2.5 times the VITAMIN B3
- 2 times the VITAMIN D
- 3 times the VITAMIN E
- 3 times the VITAMIN B12
- 2 times the ZINC



Two-Per-Day Multivitamin Tablets

Item #02315 • 120 tablets (two-month supply)
1 bottle \$18.38 • 4 bottles \$16.25 each

Two-Per-Day Multivitamin Capsules

Item #02314 • 120 capsules (two-month supply)
1 bottle \$19.13 • 4 bottles \$17 each

Each bottle provides a two-month supply.

"Covers all the bases."

Brian
 VERIFIED
 CUSTOMER
 REVIEW

For full product description and to order **Two-Per-Day Multivitamin**, call **1-800-544-4440** or visit **www.Life Extension.com**

*2023 Consumer Satisfaction, Rated #1 Catalog/Internet Merchant. Ratings based on results of the 2023 ConsumerLab.com Survey of Supplement Users. More information at www.consumerlab.com/survey.

* Compared to CENTRUM Silver Adults 50*

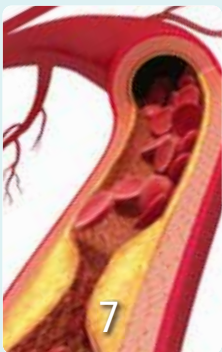
CAUTION: Individuals consuming more than 50 mcg (2000 IU)/day of vitamin D (from diet and supplements) should periodically obtain a serum 25-hydroxy vitamin D measurement. Vitamin D supplementation is not recommended for individuals with high blood calcium levels.

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IN THIS EDITION OF *LIFE EXTENSION* MAGAZINE®



7 **ATHEROSCLEROSIS IN SWEDEN**

A startling **42%** of Swedish study subjects without known heart disease were found to have **atherosclerosis**. Prevalence in **Americans** is likely *higher*. Blood tests can detect risk factors for **atherosclerosis** before major cardiovascular events strike.



22 **A SOLUTION FOR URINARY SYMPTOMS IN WOMEN**

A blend of **plant extracts** has been shown to *reduce* urinary episodes by **65%** in women, with **79%** reporting significant benefit.



30 **RESOLVE PERSISTENT PAIN**

A new study shows that marine oil-derived **SPM precursors** combined with bioavailable **curcumin** *resolve and reduce* inflammation-induced discomforts.



40 **NITRIC OXIDE IMPROVES BLOOD FLOW**

A form of **L-arginine** has been shown to *boost* endothelial **nitric oxide** production for improved cardiovascular health.



63 **KATIE COURIC: A TIRELESS ADVOCATE FOR COLON CANCER AWARENESS**

In her memoir *Going There*, Katie Couric describes her relentless battle to honor her late husband's memory by advocating for **cancer screening** and finding a cure.



69 **ASTAXANTHIN AND THE LIVER**

Known for its eye and brain benefits, preclinical studies show that **astaxanthin** can help protect the **liver**.