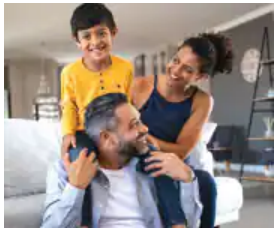


Life Extension Magazine®

PQQ Revitalizes Aging Cells

PQQ promotes production of new mitochondria. Human studies continue to validate its efficacy. A recent animal model showed PQQ increased lifespan by 30%.

Scientifically reviewed by: **Dr. Gary Gonzalez**, MD, in November 2021. Written by: Marsha McCulloch, MS, RD.



The **mitochondria** are referred to as the **energy** powerhouses of our cells.

But like parts in a machine, they become damaged and dysfunctional over time.¹

Mitochondrial dysfunction is associated with an array of age-related health problems, including insulin resistance, macular degeneration, cognitive decline, and osteoarthritis.²⁻⁴

A compound called **pyrroloquinoline quinone(PQQ)** helps stimulate the creation of healthy **new mitochondria**.⁵⁻⁸

This has the potential to ward off multiple health issues including blood sugar dysregulation, neurodegeneration, and more.⁹⁻¹²

In an animal model used to study **longevity**, PQQ increased **lifespan** by an average of **30%**.^{13,14}

Consider including **10 mg to 20 mg** of **PQQ** each day as part of an anti-aging program.

What Is PQQ?

PQQ (pyrroloquinoline quinone) is a water-soluble, vitamin-like compound found in plants, yeast, and certain bacteria.¹⁵

PQQ plays a key role in supporting **mitochondria**. These "power plants" of cells generate the energy that cells need to **function**.¹⁶

Mitochondria are critical regulators of cell processes, including immune system activation, cellular signaling, and inflammation.^{1,17,18}

As we age, mitochondria become damaged.¹⁹ That can interfere with the **function** of many organs and tissues, especially ones with **higher energy requirements**. These include the heart, brain, retina, kidney, liver, pancreas, and musculoskeletal system.²⁰

Supporting the Mitochondria

PQQ fortifies mitochondria in a few distinct ways:

- In animal models, it has been shown to enhance the action of **enzymes** that help mitochondria increase energy production.^{21,22}
- In mice, it has improved **mitophagy**, the clearing out of *defective* mitochondria.²³ Damaged mitochondria must be cleared to ensure cell survival.²⁴
- It promotes the creation of healthy, **new mitochondria** to replace those that are damaged or dysfunctional.^{5,25}



PQQ boosts the synthesis of new mitochondria by activating a protein called **PGC-1 alpha**.^{11,26}

In a study of healthy men who completed a 6-week aerobic training program, those given **20 mg** of **PQQ** daily more than **doubled** their **PGC-1 alpha** levels. Men given a **placebo** had a significantly smaller increase of this mitochondrial-generating protein.⁶

In mouse models, PQQ helps undo the harm caused by dysfunctional mitochondria, reducing the tendency of **chronic inflammation**, and preventing **oxidative damage**.^{22,27}

Promoting Longevity

The potential benefits of PQQ are so extraordinary, one of the world's leading nutritional scientists has called it a "**longevity vitamin**."²⁸

Increasing the number of healthy mitochondria can preserve youthful function and vitality.

Preclinical research has shown PQQ helps prevent or delay chronic, low-grade inflammation in cells. This inflammation is so closely tied to the aging process, it is sometimes referred to as **inflammaging**.²⁹

When human lung cells were pretreated with PQQ and then exposed to **inflammatory cytokines** (signaling proteins), fewer dysfunctional **senescent cells** developed, compared to untreated cells.²⁹ These senescent cells are a major driver of disease and accelerated aging.

Other research has shown PQQ promotes the activity of longevity genes.²⁹ It also promotes resistance to **oxidative stress**, which helps ward off premature aging.¹³

These benefits add up. In a roundworm model used to study longevity, PQQ was found to **increase lifespan** by an average of **30%**.^{13,14}

Brain and Nerve Protection



Mitochondrial function is vital for the health of the **brain**, which uses about **20%** of the energy of the body.¹⁶

Mitochondrial production and function are *impaired* in both **Alzheimer's** and **Parkinson's disease**.^{30,31} Research suggests **mitophagy** is also impaired in these brain conditions.²⁴

Animal models of Parkinson's disease show PQQ *reduces* the loss of neurons that produce **dopamine**.³² This nerve messenger is essential for regulating movement of the body. Dopamine shortfalls lead to shaking, stiffness, and difficulty walking, hallmarks of Parkinson's.³³

Preclinical research has also shown that PQQ *protects* nerve cells from damage from **beta-amyloid**, a harmful protein that accumulates in patients with Alzheimer's disease.³⁴

An animal model suggests PQQ may even help reduce **brain damage** when taken before or shortly after a **stroke**.³⁵

WHAT YOU NEED TO KNOW

PQQ Promotes Health and Longevity

- In aging cells, **mitochondria** become damaged, trigger inflammation, and produce insufficient energy.
- **Mitochondrial dysfunction** is associated with many age-related health problems, including insulin resistance, macular degeneration, brain disorders, and joint pain.
- **PQQ (pyrroloquinoline quinone)** is a compound that can promote the recycling of defective mitochondria and the creation of healthy, **new mitochondria**.
- In preclinical research, PQQ helps protect the health of the brain and eyes, reduce blood sugar and cholesterol, defend against kidney and liver damage, combat osteoporosis and osteoarthritis, and more



- PQQ is sometimes referred to as a "**longevity vitamin**." In roundworms, it increased lifespan by an average of **30%**.

Improved Eye Health

Age-related **macular degeneration** is a leading cause of blindness.⁸

Cell studies from eyes of human donors with macular degeneration show **under-functioning mitochondria** in the **retinal pigment epithelium**.⁸

Without enough energy, this tissue can't do its job.⁸

Abnormal mitochondrial function is also frequently found in **glaucoma** and **diabetic retinopathy**, two other conditions that can lead to blindness.³⁶

In a recent preclinical study, scientists used PQQ to treat retinal pigment epithelial cells from human organ donors who had age-related macular degeneration.⁸

About half of the tissue samples had a **50%–350%** improvement in **mitochondrial function** of retinal pigment epithelial cells when **PQQ** was administered, compared to untreated donor cells.

Cells in the PQQ group also had a **59% increase** in production of **ATP**, the energy currency that cells use.

PQQ also regulates **antioxidant genes** in the retinal cells, helping protect them from further damage.⁸

Preserving Metabolic Health

Mitochondrial dysfunction can also contribute to **metabolic syndrome**, obesity, insulin resistance, type II diabetes, and cardiovascular disease.^{21,37,38}

Scientists treated obese rats with PQQ for five weeks. These rats had been fed a high-fat, high-fructose diet to induce metabolic dysfunction.²¹

Compared to an untreated group of obese rats, **PQQ** significantly:

- Improved **blood sugar control, insulin levels, and insulin sensitivity**,
- Lowered harmful **inflammatory cytokines**, including TNF-alpha and IL-6, and
- Promoted **healthier blood lipid levels**, including total cholesterol, triglycerides, LDL ("bad") cholesterol, and HDL ("good") cholesterol.

PQQ may also help prevent unhealthy **heart enlargement**, based on promising results from a cell-based model. This type of heart enlargement, known as cardiac hypertrophy, can be caused by excessive free radicals and inflammation.³⁹

In this preclinical research, PQQ **decreased free radicals** in human heart cells by up to **50%**. That could help reduce the risk of heart failure and other cardiac conditions.³⁹

Averting Kidney and Liver Damage

High blood sugar and diabetes often lead to **kidney damage**. PQQ may help counteract these causes of kidney problems.

In an animal model of diabetes, rats given **PQQ** for four weeks had a significant **decrease in free radicals** in kidney tissue.⁴⁰ **Structural damage** to the kidneys improved significantly and the kidneys worked better with PQQ, compared to the kidneys of untreated rats.

The **liver** is also vulnerable to the effects of metabolic dysfunction associated with diabetes and obesity.⁴¹



In a preclinical study of metabolic **fatty liver disease**, PQQ protected the liver from damaging fat accumulation. It did this by improving lipid metabolism, supporting the creation of new mitochondria, and boosting antioxidant protection.⁴²

Supporting Musculoskeletal Health

Animal research has shown that PQQ can help combat the bone disease **osteoporosis**, in part by increasing **antioxidant** protection.⁴³⁻⁴⁵ Osteoporosis occurs when bone breakdown, a normal process, outpaces bone building.

One potential contributing factor for osteoporosis is an excess of free radicals that hastens bone breakdown. Hormone changes that happen with aging accelerate this process.⁴⁵

In a postmenopausal animal model, PQQ was as effective **as estrogen replacement** at preventing **bone loss**—but without the side effects of hormone therapy.⁴⁵

The antioxidant benefits of PQQ may also support the healing of **bone fractures**, according to another study using a postmenopausal animal model.²⁷

Similarly, other research shows that the progression of **osteoarthritis** is related to **oxidative stress**.⁴³

Cartilage cells are rich in mitochondria. Disruption of **mitochondrial function** increases oxidative stress and inflammation and damages the cartilage that cushions joints.^{46,47}

In preclinical research, PQQ significantly *decreased* mitochondrial damage and dysfunction caused by inflammatory cytokines in the mitochondria of **cartilage**.⁴⁸

Supporting mitochondrial health is crucial for healthy aging.

Summary



Mitochondria, the energy factories of cells, play a key role in regulating the **aging** process and promoting longevity.

As we age, our mitochondria tend to wear out and become less likely to be replaced.

PQQ is a vitamin-like compound that promotes the production of healthy **new mitochondria**. It is also a potent **antioxidant**.

By supporting mitochondria, PQQ promotes the health of organs that are especially vulnerable to energy deficits and oxidative stress, including the **brain, eyes, and heart**.

In an animal model, it increased **longevity** by **30%**.

Consider taking **10 mg to 20 mg** of **PQQ** daily as part of an anti-aging program.

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

References

1. Rossmann MP, Dubois SM, Agarwal S, et al. Mitochondrial function in development and disease. *Dis Model Mech*. 2021 Jun 1;14(6).
2. Kaarniranta K, Uusitalo H, Blasiak J, et al. Mechanisms of mitochondrial dysfunction and their impact on age-related macular degeneration. *Prog Retin Eye Res*. 2020 Nov;79:100858.
3. He Y, Wu Z, Xu L, et al. The role of SIRT3-mediated mitochondrial homeostasis in osteoarthritis. *Cell Mol Life Sci*. 2020 Oct;77(19):3729-43.
4. Potenza MA, Sgarra L, Desantis V, et al. Diabetes and Alzheimer's Disease: Might Mitochondrial Dysfunction Help Deciphering the Common Path? *Antioxidants (Basel)*. 2021 Aug 6;10(8).



5. Naveed M. The Life History of Pyrroloquinoline Quinone (PQQ): A Versatile Molecule with Novel Impacts on Living Systems. *International Journal of Molecular Biology*. 2016;1(1).
6. Hwang PS, Macheck SB, Cardaci TD, et al. Effects of Pyrroloquinoline Quinone (PQQ) Supplementation on Aerobic Exercise Performance and Indices of Mitochondrial Biogenesis in Untrained Men. *J Am Coll Nutr*. 2020 Aug;39(6):547-56.
7. Harris CB, Chohanadisai W, Mishchuk DO, et al. Dietary pyrroloquinoline quinone (PQQ) alters indicators of inflammation and mitochondrial-related metabolism in human subjects. *J Nutr Biochem*. 2013 Dec;24(12):2076-84.
8. Ebeling MC, Polanco JR, Qu J, et al. Improving retinal mitochondrial function as a treatment for age-related macular degeneration. *Redox Biol*. 2020 Jul;34:101552.
9. Lu J, Chen S, Shen M, et al. Mitochondrial regulation by pyrroloquinoline quinone prevents rotenone-induced neurotoxicity in Parkinson's disease models. *Neurosci Lett*. 2018 Nov 20;687:104-10.
10. Jornayvaz FR, Shulman GI. Regulation of mitochondrial biogenesis. *Essays Biochem*. 2010;47:69-84.
11. Ames BN. Prolonging healthy aging: Longevity vitamins and proteins. *Proc Natl Acad Sci U S A*. 2018 Oct 23;115(43):10836-44.
12. Wang Z, Chen GQ, Yu GP, et al. Pyrroloquinoline quinone protects mouse brain endothelial cells from high glucose-induced damage in vitro. *Acta Pharmacol Sin*. 2014 Nov;35(11):1402-10.
13. Wu JZ, Huang JH, Khanabdali R, et al. Pyrroloquinoline quinone enhances the resistance to oxidative stress and extends lifespan upon DAF-16 and SKN-1 activities in *C. elegans*. *Exp Gerontol*. 2016 Jul;80:43-50.
14. Sasakura H, Moribe H, Nakano M, et al. Lifespan extension by peroxidase and dual oxidase-mediated ROS signaling through pyrroloquinoline quinone in *C. elegans*. *J Cell Sci*. 2017 Aug 1;130(15):2631-43.
15. Jonscher KR, Rucker RB. Pyrroloquinoline Quinone. In: Watson RR, Preedy VR, editors. *Dietary Interventions in Liver Disease*: Academic Press; 2019:157-73.
16. Woo J, Cho H, Seol Y, et al. Power Failure of Mitochondria and Oxidative Stress in Neurodegeneration and Its Computational Models. *Antioxidants (Basel)*. 2021 Feb 3;10(2).
17. Suarez-Rivero JM, Pastor-Maldonado CJ, Povea-Cabello S, et al. From Mitochondria to Atherosclerosis: The Inflammation Path. *Biomedicines*. 2021 Mar 5;9(3).
18. Su YJ, Wang PW, Weng SW. The Role of Mitochondria in Immune-Cell-Mediated Tissue Regeneration and Ageing. *Int J Mol Sci*. 2021 Mar 6;22(5).
19. Chistiakov DA, Sobenin IA, Revin VV, et al. Mitochondrial aging and age-related dysfunction of mitochondria. *Biomed Res Int*. 2014;2014:238463.
20. Krako Jakovljevic N, Pavlovic K, Jotic A, et al. Targeting Mitochondria in Diabetes. *Int J Mol Sci*. 2021 Jun 21;22(12).
21. Devasani K, Kaul R, Majumdar A. Supplementation of pyrroloquinoline quinone with atorvastatin augments mitochondrial biogenesis and attenuates low grade inflammation in obese rats. *Eur J Pharmacol*. 2020 Aug 15;881:173273.
22. Jonscher KR, Stewart MS, Alfonso-Garcia A, et al. Early PQQ supplementation has persistent long-term protective effects on developmental programming of hepatic lipotoxicity and inflammation in obese mice. *FASEB J*. 2017 Apr;31(4):1434-48.
23. Ma W, Zhang R, Huang Z, et al. PQQ ameliorates skeletal muscle atrophy, mitophagy and fiber type transition induced by denervation via inhibition of the inflammatory signaling pathways. *Ann Transl Med*. 2019 Sep;7(18):440.
24. Burtcher J, Millet GP, Place N, et al. The Muscle-Brain Axis and Neurodegenerative Diseases: The Key Role of Mitochondria in Exercise-Induced Neuroprotection. *Int J Mol Sci*. 2021 Jun 17;22(12).
25. Zhang Q, Zhou J, Shen M, et al. Pyrroloquinoline Quinone Inhibits Rotenone-Induced Microglia Inflammation by Enhancing Autophagy. *Molecules*. 2020 Sep 23;25(19).

26. Scarpulla RC. Metabolic control of mitochondrial biogenesis through the PGC-1 family regulatory network. *Biochim Biophys Acta*. 2011 Jul;1813(7):1269-78.
27. Wu X, Zhou X, Liang S, et al. The mechanism of pyrroloquinoline quinone influencing the fracture healing process of estrogen-deficient mice by inhibiting oxidative stress. *Biomed Pharmacother*. 2021 Jul;139:111598.
28. Zhu W, Klinman JP. Biogenesis of the peptide-derived redox cofactor pyrroloquinoline quinone. *Curr Opin Chem Biol*. 2020 Dec;59:93-103.
29. Hao J, Ni X, Giunta S, et al. Pyrroloquinoline quinone delays inflammaging induced by TNF-alpha through the p16/p21 and Jagged1 signalling pathways. *Clin Exp Pharmacol Physiol*. 2020 Jan;47(1):102-10.
30. Popov LD. Mitochondrial biogenesis: An update. *J Cell Mol Med*. 2020 May;24(9):4892-9.
31. Simmons EC, Scholpa NE, Schnellmann RG. Mitochondrial biogenesis as a therapeutic target for traumatic and neurodegenerative CNS diseases. *Exp Neurol*. 2020 Jul;329:113309.
32. Cheng Q, Chen J, Guo H, et al. Pyrroloquinoline quinone promotes mitochondrial biogenesis in rotenone-induced Parkinson's disease model via AMPK activation. *Acta Pharmacol Sin*. 2021 May;42(5):665-78.
33. Sonninen TM, Hamalainen RH, Koskivi M, et al. Metabolic alterations in Parkinson's disease astrocytes. *Sci Rep*. 2020 Sep 2;10(1):14474.
34. Kim J, Kobayashi M, Fukuda M, et al. Pyrroloquinoline quinone inhibits the fibrillation of amyloid proteins. *Prion*. 2010 Jan-Mar;4(1):26-31.
35. Zhang Y, Feustel PJ, Kimelberg HK. Neuroprotection by pyrroloquinoline quinone (PQQ) in reversible middle cerebral artery occlusion in the adult rat. *Brain Res*. 2006 Jun 13;1094(1):200-6.
36. Carrella S, Massa F, Indrieri A. The Role of MicroRNAs in Mitochondria-Mediated Eye Diseases. *Front Cell Dev Biol*. 2021;9:653522.
37. Gonzalez-Franquesa A, Patti ME. Insulin Resistance and Mitochondrial Dysfunction. *Adv Exp Med Biol*. 2017;982:465-520.
38. Morciano G, Patergnani S, Bonora M, et al. Mitophagy in Cardiovascular Diseases. *J Clin Med*. 2020 Mar 24;9(3):892.
39. Wen J, Shen J, Zhou Y, et al. Pyrroloquinoline quinone attenuates isoproterenol hydrochloride-induced cardiac hypertrophy in AC16 cells by inhibiting the NFkappaB signaling pathway. *Int J Mol Med*. 2020 Mar;45(3):873-85.
40. Zhang M, Zhang J, Xiong Y, et al. Pyrroloquinoline Quinone Inhibits Oxidative Stress in Rats with Diabetic Nephropathy. *Med Sci Monit*. 2020 Jun 27;26:e924372.
41. Karkucinska-Wieckowska A, Simoes ICM, Kalinowski P, et al. Mitochondria, oxidative stress and nonalcoholic fatty liver disease: A complex relationship. *Eur J Clin Invest*. 2021 May 29:e13622.
42. Qiu K, Zhao Q, Wang J, et al. Effects of Pyrroloquinoline Quinone on Lipid Metabolism and Anti-Oxidative Capacity in a High-Fat-Diet Metabolic Dysfunction-Associated Fatty Liver Disease Chick Model. *Int J Mol Sci*. 2021 Feb 1;22(3).
43. Qin R, Sun J, Wu J, et al. Pyrroloquinoline quinone prevents knee osteoarthritis by inhibiting oxidative stress and chondrocyte senescence. *Am J Transl Res*. 2019;11(3):1460-72.
44. Wu X, Li J, Zhang H, et al. Pyrroloquinoline quinone prevents testosterone deficiency-induced osteoporosis by stimulating osteoblastic bone formation and inhibiting osteoclastic bone resorption. *Am J Transl Res*. 2017;9(3):1230-42.
45. Geng Q, Gao H, Yang R, et al. Pyrroloquinoline Quinone Prevents Estrogen Deficiency-Induced Osteoporosis by Inhibiting Oxidative Stress and Osteocyte Senescence. *Int J Biol Sci*. 2019;15(1):58-68.
46. de Sire A, Marotta N, Marinario C, et al. Role of Physical Exercise and Nutraceuticals in Modulating Molecular Pathways of Osteoarthritis. *Int J Mol Sci*. 2021 May 27;22(11).



47. Mao X, Fu P, Wang L, et al. Mitochondria: Potential Targets for Osteoarthritis. *Front Med (Lausanne)*. 2020 November-26;7(808):581402.
48. Han GT, Cai WS, Zhang YB, et al. Protective Effect of Pyrroloquinoline Quinone on TNF-alpha-induced Mitochondrial Injury in Chondrocytes. *Curr Med Sci*. 2021 Feb;41(1):100-7.

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