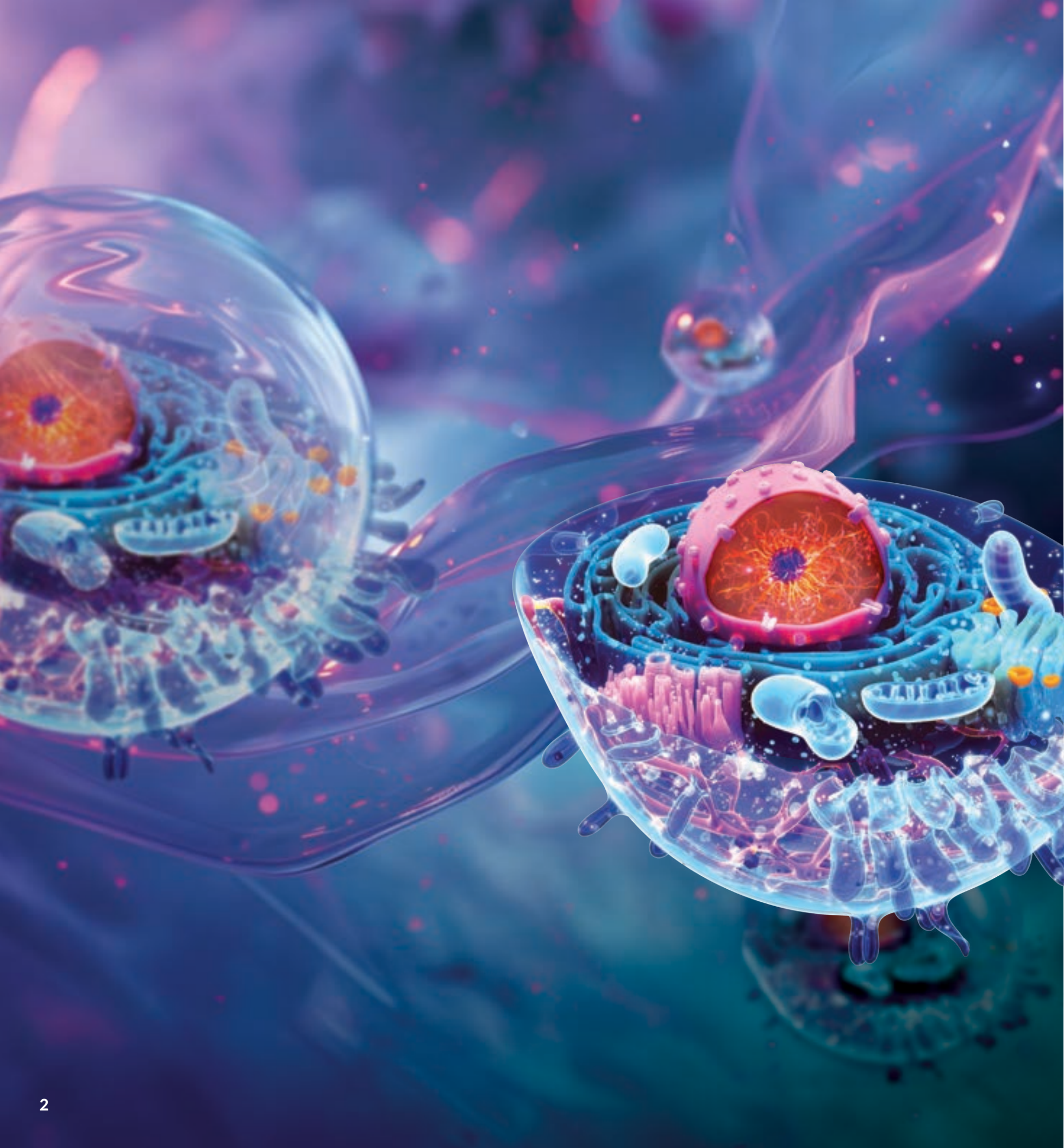




Foundational Science: Examining Ubiquinol's Role in Mitochondrial Function



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Rethinking the Powerhouse: Mitochondria at the Center of Cellular Health

As new information about mitochondrial function has become a focus of scientific research, it has led to a greater understanding of the role of ubiquinol in several processes foundational to cellular and overall wellness. Beyond their role in producing adenosine triphosphate (ATP)—the energy molecule of the cell—mitochondria are now recognized for their contribution to the following broader cellular processes that together help maintain cellular homeostasis, a critical pillar for wellness:

- | | | |
|---|--|--|
| Mitochondrial biogenesis
Increasing mitochondrial size and number to meet rising energy demands or prepare for cell division ¹ | Mitochondrial fission and fusion
Splitting and rejoining mitochondria to repair damage, recycle components, and adapt to stress ^{1,2} | Programmed cell death (apoptosis) and mitophagy
Safely removing damaged cells or mitochondria to protect overall function ^{4,5} |
| | Signaling pathways
Coordinating stress and immune responses and repair mechanisms to maintain balance ³ | Cell senescence
Permanently halting the division of damaged cells to protect surrounding cells ^{6,7} |

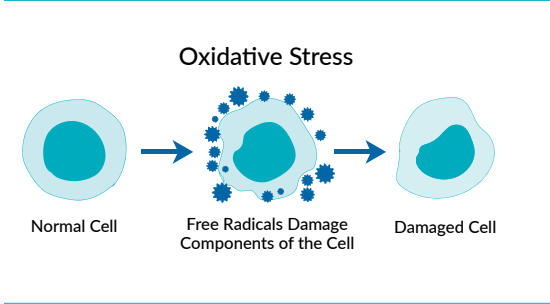
These roles position mitochondria as central to cellular resilience, maintenance, and performance. They also have an integral and wide-ranging effect on physiological processes associated with aging.^{8,9}

Mitochondria are now being described as the CEO (“Chief Executive Organelle”) of the cell, responsible for regulating complex metabolic networks, signaling, innate immunity, cell division, cell fate, and ultimately, the fate of the organism.^{3,10}

Oxidative Stress and Its Effect on Mitochondrial and Cellular Resilience

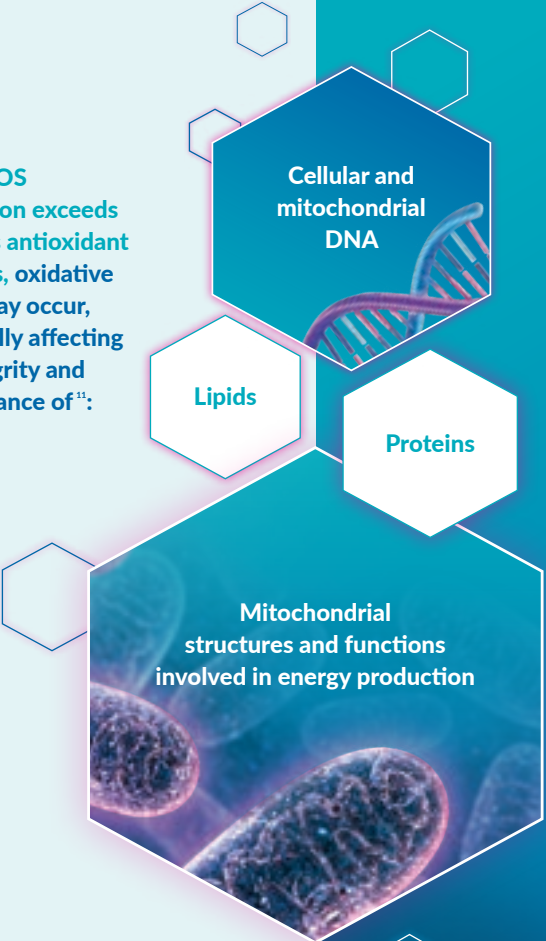
Reactive oxygen species (ROS) are natural byproducts of mitochondrial energy generation and are neutralized by antioxidants.¹¹

Oxidative stress may influence energy production and cellular repair, with age-related changes in antioxidant defenses further affecting mitochondrial function and mitochondrial DNA (mtDNA).⁸ The **oxidative stress theory of aging** states that long-term imbalances between ROS and antioxidant defenses can influence mitochondrial activity and cellular health with age.¹²



The ratio of ubiquinol to ubiquinone, commonly referred to as CoQ10 balance, is one measure used in research as an indicator of oxidative stress. When the percentage of ubiquinone increases and ubiquinol decreases, this indicates greater usage of ubiquinol to manage free radicals. The more this balance tilts toward less ubiquinol, the greater the oxidative stress.

When ROS production exceeds the cell's antioxidant defenses, oxidative stress may occur, potentially affecting the integrity and performance of¹¹:



Oxidative stress can also be exacerbated by lifestyle and environmental factors, such as¹³:

- ⊗ Poor nutritional intake
- ⊗ Physical inactivity
- ⊗ Tobacco and alcohol use
- ⊗ Exposure to UV radiation, pollution, and other environmental toxins

Ubiquinol: Contributing to Cellular Energy and Antioxidant Defenses

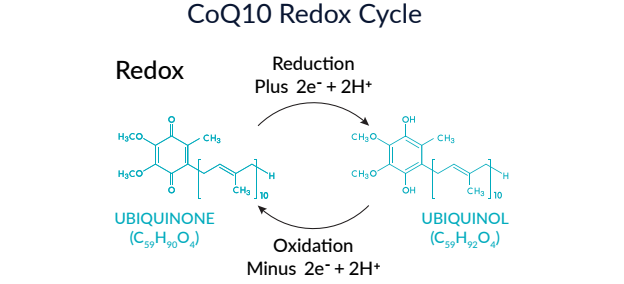
Ubiquinol is naturally produced in the body. As the active antioxidant form of CoQ10, it is used both in the production of ATP and to neutralize ROS that are produced during that process.

Lipid soluble and concentrated in the mitochondria,¹⁴ ubiquinol helps:

- Support cellular energy generation¹⁵
- Neutralize ROS at the source¹¹
- Protect lipids against peroxidation and damage^{11,16}
- Maintain mitochondrial membrane integrity



Ubiquinone and ubiquinol form a redox pair that cycles between oxidized and reduced states during energy generation and antioxidant activity. Ubiquinone is part of the electron transport chain, and once it is reduced by an enzyme complex, it becomes the antioxidant ubiquinol. When ubiquinol acts as an antioxidant and donates its electrons, it is oxidized and returns to being ubiquinone. This is known as a redox cycle.¹⁴



Research shows that ubiquinol plays a key role in both **energy generation and antioxidant protection** by being able to cycle back and forth between ubiquinone and ubiquinol.^{11,14}

The levels of ubiquinol decline with age, due to a decreased ability to convert ubiquinone to ubiquinol and an increased demand for its antioxidant activity because of greater ROS production.^{17,18}

Ubiquinol's vital role in antioxidant protection has been broadly investigated and shown to positively affect cardiovascular function,^{16,19} preconception wellness,²⁰⁻²³ menopausal well-being,²⁴⁻²⁶ healthy aging,^{19,27-29} and sports nutrition.^{30,31}

Ubiquinol is ubiquitous, as it is naturally found in almost every cell in the body.¹⁵

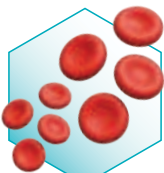
95% of the total CoQ10 in the blood is in the ubiquinol form.³²

Mitochondrial Function Through the Aging Process

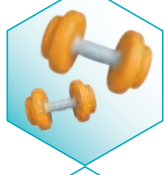
The passage of time is accompanied by the accumulation of wear and tear on all parts of the body, down to the very smallest of them. Healthy lifestyle habits such as eating a nutritious diet, getting regular exercise, prioritizing sleep, and managing stress make a profound difference, even at the mitochondrial level.²

However, even under the best of circumstances, aging is characterized by some decline in mitochondrial function. This triggers cellular changes that, over time, manifest in many conditions associated with aging.^{14,18,33}

Ubiquinol's role in maintaining cellular homeostasis supports healthy cellular function in the face of daily environmental stresses, whatever the stage of life.



Clinical studies show that supplementation with Kaneka Ubiquinol® **increases plasma ubiquinol levels.**^{19,27}



Research demonstrates a positive correlation between total CoQ10 status and percentage plasma ubiquinol and **overall physical functioning** in older adults.^{28,29}



Higher blood ubiquinol levels also promote **cardiovascular health**¹⁹ and **muscle health.**²⁸

With age, the body is less able to convert ubiquinone into ubiquinol.^{17,32}



Mitochondrial Efficiency and Cardiovascular Health

Cardiomyocytes—heart muscle cells—contain a large number of mitochondria to meet the high energy demands placed on them.³⁵ Their high metabolic rate also generates greater amounts of ROS, increasing susceptibility to oxidative stress.^{11,36}

Changes in mitochondrial efficiency may affect energy output and, in turn, affect heart function.^{14,35}

Oxidative balance is an important factor in maintaining vascular tone and healthy blood flow in blood vessels.³⁷

Excess ROS can directly affect the function of endothelial cells (which form the inner lining of vessels) from the mitochondrial to the cellular and tissue levels.³⁸

In addition, ROS can oxidize circulating LDL particles in the blood, which has been associated with changes in vascular health.^{16,19,39}

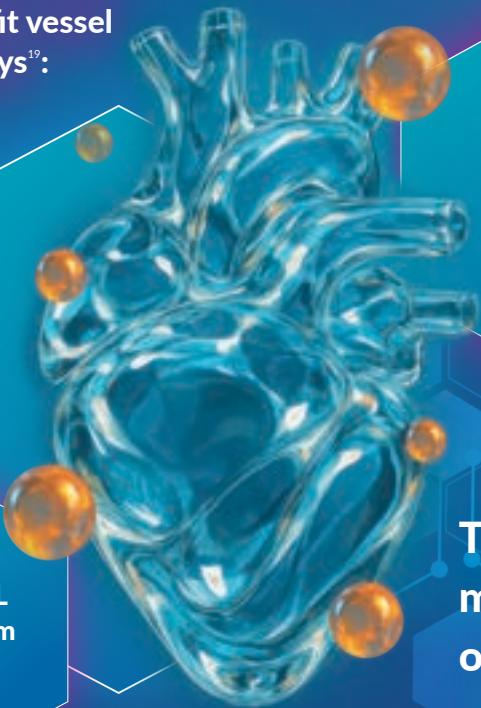
Clinical research shows that Ubiquinol supplementation improves blood markers associated with heart health.^{40,41} Additional studies highlight the relevance of these markers as indicators associated with cardiovascular health.^{42,43}

In addition, ubiquinol helps to maintain mitochondrial membrane integrity, which is now understood to be foundational for maintaining heart and blood vessel structure and function, particularly with age.^{18,44,45}

Kaneka Ubiquinol® has been shown in a clinical study to benefit vessel health in the following ways¹⁹:

Facilitating proper vasodilation and blood circulation

Protecting LDL cholesterol from oxidation



Enhancing nitric oxide (NO) production

The heart is one of the most energy-demanding organs in the body.³⁵

The Energy and Antioxidant Needs of Reproductive Cells

Healthy mitochondrial function is essential for the proper function of reproductive cells,⁴⁴ which have high energy demands²⁰ and are sensitive to oxidative stress.

Age-related changes in oxidative stress and mitochondrial function have been associated with changes in sperm health,⁴⁷ oocyte function, and ovarian reserve—particularly in women over 35.⁴⁸



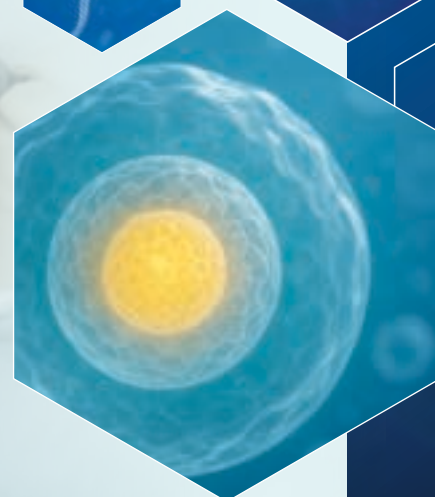
Female preconception support

Ubiquinol's potent antioxidant activity supports mitochondrial function and cellular energy generation—essential for oocyte quality and overall egg health.^{21,48,49}



Male preconception support

Kaneka Ubiquinol® supports healthy sperm function, including morphology, concentration, and motility,^{22,23} and provides antioxidant protection to seminal fluid.⁵⁰



Menopause and Redox Balance

During the menopausal transition, declining estrogen levels are associated with reduced antioxidant defenses and higher oxidative stress.^{24,51-53}

Higher oxidative stress can put a strain on ubiquinol levels which, over time, may impact mitochondrial function.

Kaneka Ubiquinol® supports general health and well-being during and after menopause.

80%

In a consumer use study, **80% of menopausal women** taking 200 mg of Kaneka Ubiquinol® per day reported decreased irritability, sensitivity, stress, and mood swings after 60 days of supplementation.^{25,26}



Antioxidant capacity is decreased in postmenopausal women.⁵¹

Oxidative stress markers increase after menopause.^{24,52,53}

Ubiquinol for Exercise Performance

During exercise, contraction of the heart and muscles increases, upping the demand for ATP as much as 10-fold.⁵⁴ This is why heart muscle cells contain three to five times more mitochondria than most other cells,⁵⁴ and other muscle cells are dense with mitochondria in specialized arrangements for maximal ATP generation and access. The increase in ATP generation also elevates the production of ROS, leading to greater oxidative stress during and following exercise.⁵⁵

Athletic ability is influenced by training intensity, age, and baseline fitness,⁵⁶ as well as stressors such as high altitude. Exercising at high altitude can further challenge oxidative balance by increasing hypoxia-driven ROS production and placing an additional strain on oxygen delivery and cardiorespiratory capacity.³⁰

In a clinical study, participants taking 200 mg of Kaneka Ubiquinol® per day reduced the overall decrease in cardiorespiratory fitness experienced when exercising at high altitude by almost half compared to the placebo—a decrease of 11% in the Ubiquinol group versus 21% for the placebo group.⁵⁷

Recent research shows that Kaneka Ubiquinol® supports physical performance by:



Promoting a healthy oxidative balance during exercise³¹



Supporting energy metabolism during exercise¹⁵



Helping the body adapt to rigorous exercise^{57,58}



Enhancing peak power production in elite athletes when taken at 300 mg per day⁵⁹



Promoting healthy cardiovascular performance when exercising at high altitudes^{57,58}



The Kaneka Ubiquinol® Difference

At Kaneka Nutrients, our commitment to scientific rigor and quality assurance drives the development, safety, and efficacy of our ingredients.

Kaneka Ubiquinol® is 2x better absorbed than conventional CoQ10 supplements⁶⁰ and does not require conversion to be absorbed into the blood and to function as an antioxidant.^{19,61}

Research demonstrates that 200 mg of Kaneka Ubiquinol® increases ubiquinol levels by approximately 8x compared to baseline in healthy adults when taken daily for at least 30 days.²⁷



Kaneka Ubiquinol®: Backed by Science.
Trusted by 200+ Brands.



Made in the USA

18+

18+ years of positive consumer experience with Kaneka Ubiquinol® supplementation

100+

Backed by 100+ clinical studies using Kaneka Ubiquinol®

50 years

50 years of ubiquinone and ubiquinol research and testing



Free of impurities commonly found in synthetic CoQ10



Bioidentical to the body's own ubiquinol

Get the Kaneka Ubiquinol® Advantage.

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Mitochondria and Ubiquinol

01

Ubiquinol's lipid solubility enables it to concentrate within the mitochondrial membrane.^{14,19}

02

Ubiquinol helps transfer electrons between complexes on the electron transport chain.¹⁵

03

ROS are natural byproducts of mitochondrial energy generation.¹¹

04

Ubiquinol helps to neutralize ROS created by the electron transport chain.¹¹

05

By keeping ROS in check, ubiquinol helps maintain the integrity of the mitochondrial membrane.^{11, 18}

References

1. Adebayo M, Singh S, Singh AP, Dasgupta S. Mitochondrial fusion and fission: the fine-tune balance for cellular homeostasis. *FASEB J*. 2021;35(6):e21620.

2. Liu YJ, McIntyre RL, Janssens GE, Houtkooper RH. Mitochondrial fission and fusion: a dynamic role in aging and potential target for age-related disease. *Mech Ageing Dev*. 2020;186:111212.

3. Picard M, Shirihaï OS. Mitochondrial signal transduction. *Cell Metab*. 2022;34(11):1620-53.

4. Wang S, Long H, Hou L, et al. The mitophagy pathway and its implications in human diseases. *Sig Transduct Target Ther*. 2023;8:304.

5. Tait SW, Green DR. Mitochondrial regulation of cell death. *Cold Spring Harb Perspect Biol*. 2013;5(9):a008706.

6. Kumari R, Jat P. Mechanisms of cellular senescence: cell cycle arrest and senescence associated secretory phenotype. *Front Cell Dev Biol*. 2021;9:645593.

7. Varesi A, Chirumbolo S, Campagnoli LIM, et al. The role of antioxidants in the interplay between oxidative stress and senescence. *Antioxidants* (Basel). 2022;11(7):1224.

8. Lima T, Li TY, Mottis A, Auwerx J. Pleiotropic effects of mitochondria in aging. *Nat Aging*. 2022;2(3):199-213.

9. Asejeje FO, Ogunro OB. Deciphering the mechanisms, biochemistry, physiology, and social habits in the process of aging. *Arch Gerontol Geriatrics Plus*. 2024;1(1):100003.

10. Lee-Glover LP, Picard M, Shutt TE. Mitochondria—the CEO of the cell. *J Cell Sci*. 2025;138(9):jcs263403.

11. Bentinger M, Brismar K, Dallner G. The antioxidant role of coenzyme Q. *Mitochondrion*. 2007;7(Suppl):S41-50.

12. Beckman KB, Ames BN. The free radical theory of aging matures. *Physiol Rev*. 1998;78(2):547-81.

13. Srivastava KK, Kumar R. Stress, oxidative injury and disease. *Indian J Clin Biochem*. 2015;30(1):3-10.

14. Pallotti F, Bergamini C, Lamperti C, Fato R. The roles of coenzyme Q in disease: direct and indirect involvement in cellular functions. *Int J Mol Sci*. 2021;23(1):128.

15. Martini FH. Metabolism, nutrition and energetics. In: *Fundamentals of Anatomy and Physiology*. 12th ed. Pearson; 2024:943-50.

16. Stocker R, Bowry V, Frei B. Ubiquinol-10 protects human low-density lipoprotein more efficiently against lipid peroxidation than does alpha-tocopherol. *Proc Natl Acad Sci USA*. 1991;88:1646-50.

17. Wada H, Goto H, Hagiwara S, Yamamoto Y. Redox status of coenzyme Q10 is associated with chronological age. *J Am Geriatr Soc*. 2007;55(7):1141-2.

18. Navas P, Villalba JM, de Cabo R. The importance of plasma membrane coenzyme Q in aging and stress responses. *Mitochondrion*. 2007;7(Suppl):S34-40.

19. Sabbatinelli J, Orlando P, Galeazzi R, et al. Ubiquinol ameliorates endothelial dysfunction in subjects with mild-to-moderate dyslipidemia: a randomized clinical trial. *Nutrients*. 2020;12(4):1098.

20. Bentov Y, Casper RF. The aging oocyte—can mitochondrial function be improved? *Fertil Steril*. 2013;99(1):18-22.

21. Ben-Meir A, Burstein E, Borrego-Alvarez A, et al. Coenzyme Q10 restores oocyte mitochondrial function and fertility during reproductive aging. *Aging Cell*. 2015;14(5):887-95.

22. Thakur AS, Litarru GP, Funahashi I, et al. Effect of ubiquinol therapy on sperm parameters and serum testosterone levels in oligoasthenozoospermic infertile men. *J Clin Diagn Res*. 2015;9(9):BC01-3.

23. Cakiroglu B, Eyyupoglu SE, Gozukucuk R, Uyanik BS. Ubiquinol effect on sperm parameters in subfertile men who have astheno-teratozoospermia with normal sperm concentration. *Nephro Urol Mon*. 2014;6(3):e16870.

24. Sánchez-Rodríguez MA, Zacarías-Flores M, Arronte-Rosales A, Correa-Muñoz E, Mendoza-Núñez VM. Menopause as risk factor for oxidative stress. *Menopause*. 2012;19(3):361-7.

25. Palacios S, Ramírez M, Lilue M, et al. Estudio clínico para conocer la eficacia de la coenzima Q10 sobre la calidad de vida en mujeres postmenopáusicas. *Toko-Gin Pract*. 2019;78(1):3-7. (Proprietary English translation on file.)

26. Kaneka Internal Report. Real-life UBIQUINOL study on 200 postmenopausal women. Expansion Consulteam. 2024.

27. Hosoe K, Kitano M, Kishida H, et al. Study on safety and bioavailability of ubiquinol (Kaneka QH) after single and 4-week multiple oral administration to healthy volunteers. *Regul Toxicol Pharmacol*. 2007;47(1):19-28.

28. Fischer A, Onur S, Niklowitz P, et al. Coenzyme Q10 status as a determinant of muscular strength in two independent cohorts. *PLoS One*. 2016;11(12):e0167124.

29. de la Bella-Garzón R, Fernández-Portero C, Alarcón D, et al. Levels of plasma coenzyme Q10 are associated with physical capacity and cardiovascular risk in the elderly. *Antioxidants* (Basel). 2022;11(2):279.

30. Gaur P, Prasad S, Kumar B, Sharma SK, Vats P. High-altitude hypoxia induced reactive oxygen species generation, signaling, and mitigation approaches. *Int J Biometeorol*. 2021;65(4):601-15.

31. Sarmiento A, Diaz-Castro J, Pulido-Moran M, et al. Short-term ubiquinol supplementation reduces oxidative stress associated with strenuous exercise in healthy adults: a randomized trial. *Biofactors*. 2016;42(6):612-22.

32. Yamashita S, Yamamoto Y. Simultaneous detection of ubiquinol and ubiquinone in human plasma as a marker of oxidative stress. *Anal Biochem*. 1997;250(1):66-73.

33. Bratic A, Larsson NG. The role of mitochondria in aging. *J Clin Invest*. 2013;123(3):951-7.

34. Niklowitz P, Onur S, Fischer A, et al. Coenzyme Q10 serum concentration and redox status in European adults: influence of age, sex, and lipoprotein concentration. *J Clin Biochem Nutr*. 2015;58(3):240-5.

35. Li A, Shami GJ, Griffiths L, Lal S, Irving H, Braet F. Giant mitochondria in cardiomyocytes: cellular architecture in health and disease. *Basic Res Cardiol*. 2023;118(1):39.

36. Forsmark-Andrée P, Lee CP, Dallner G, Ernster L. Lipid peroxidation and changes in the ubiquinone content and the respiratory chain enzymes of submitochondrial particles. *Free Radic Biol Med*. 1997;22(3):391-400.

37. Rajendran P, Rengarajan T, Thangavel J, et al. The vascular endothelium and human diseases. *Int J Biol Sci*. 2013;9(10):1057-69.

38. Chiong M, Cartes-Saavedra B, Norambuena-Soto I, et al. Mitochondrial metabolism and the control of vascular smooth muscle cell proliferation. *Front Cell Dev Biol*. 2014;2:72.

39. Mohr D, Bowry VW, Stocker R. Dietary supplementation with coenzyme Q10 results in increased levels of ubiquinol-10 within circulating lipoproteins and increased resistance of human low-density lipoprotein to the initiation of lipid peroxidation. *Biochim Biophys Acta*. 1992;1126(3):247-54.

40. Onur S, Niklowitz P, Jacobs G, et al. Ubiquinol reduces gamma glutamyl transferase as a marker of oxidative stress in humans. *BMC Res Notes*. 2014;7:427.

41. Onur S, Niklowitz P, Jacobs G, et al. Association between serum level of ubiquinol and NT-proBNP, a marker for chronic heart failure, in healthy elderly subjects. *Biofactors*. 2015;41(1):35-43.

42. Lee DS, Evans JC, Robins SJ, et al. Gamma glutamyl transferase and metabolic syndrome, cardiovascular disease, and mortality risk: the Framingham Heart Study. *Arterioscler Thromb Vasc Biol*. 2007;27(1):127-33.

43. Dhingra R, Gona P, Wang TJ, et al. Serum gamma glutamyl transferase and risk of heart failure in the community. *Arterioscler Thromb Vasc Biol*. 2010;30(9):1855-60.

44. Lin KL, Chen SD, Lin KJ, et al. Quality matters? The involvement of mitochondrial quality control in cardiovascular disease. *Front Cell Dev Biol*. 2021;9:636295.

45. Poznyak AV, Kirichenko TV, Borisov EE, Shakhpazyan NK, Kartuesov AG, Orekhov AN. Mitochondrial implications in cardiovascular aging and diseases: the specific role of mitochondrial dynamics and shifts. *Int J Mol Sci*. 2022;23(6):2951.

46. Mihalas BP, Redgrove KA, McLaughlin EA, Nixon B. Molecular mechanisms responsible for increased vulnerability of the ageing oocyte to oxidative damage. *Oxid Med Cell Longev*. 2017;2017:4015874.

47. Agarwal A, Parekh N, Pannel Selvam MK, et al. Male oxidative stress infertility (MOSI): proposed terminology and clinical practice guidelines for management of idiopathic male infertility. *World J Men's Health*. 2019;37(3):296-312.

48. Zhu Z, Xu W, Liu L. Ovarian aging: mechanisms and intervention strategies. *Med Review* (2021). 2022;2(6):590-610.

49. Zhang M, ShiYang X, Zhang Y, et al. Coenzyme Q10 ameliorates the quality of postovulatory aged oocytes by suppressing DNA damage and apoptosis. *Free Radic Biol Med*. 2019;143:84-94.

50. Alleva R, Scararmucci A, Mantero F, Bompadre S, Leoni L, Littarru GP. The protective role of ubiquinol-10 against formation of lipid hydroperoxides in human seminal fluid. *Mol Aspects Med*. 1997;18(1):221-8.

51. Doshi SB, Agarwal A. The role of oxidative stress in menopause. *J Midlife Health*. 2013;4(3):140-6.

52. Vincent J, Inassi J. Comparison of oxidative stress between premenopausal and postmenopausal women. *Nat J Physiol Pharm Pharmacol*. 2020;10(5):359-62.

53. Heravi AS, Michos ED, Zhao D, et al. Oxidative stress and menopausal status: the Coronary Artery Risk Development in Young Adults cohort study. *J Women's Health* (Larchmt). 2022;31(7):1057-65.

54. Wang X, Zhang X, Wu D, et al. Mitochondrial flashes regulate ATP homeostasis in the heart. *Elife*. 2017;6:e23908.

55. Clemente-Suárez VJ, Bustamante-Sanchez Á, Mielgo-Ayuso J, Martínez-Guardado I, Martín-Rodríguez A, Tornero-Aguilera JF. Antioxidants and sports performance. *Nutrients*. 2023;15(10):2371.

56. Militello R, Luti S, Gamberi T, Pellegrino A, Modesti A, Modesti PA. Physical activity and oxidative stress in aging. *Antioxidants* (Basel). 2024;13(5):557.

57. Liu Z, Yang J, Yang B, et al. Effect of ubiquinol on electrophysiology during high-altitude acclimatization and de-acclimatization: a substudy of the Shigatse CARDiorespiratory fitness (SCARF) randomized clinical trial. *Int J Cardiol*. 2024;401:131817.

58. Lv H, Liu Z, Sun M, et al. CARDiorespiratory fitness and effects of ubiquinol during high-altitude acclimatization and deacclimatization: the SCARF trial. *iScience*. 2025;28(3):11211.

59. Alf D, Schmidt ME, Siebrecht SC. Ubiquinol supplementation enhances peak power production in trained athletes: a double-blind, placebo controlled study. *J Int Soc Sports Nutr*. 2013;10:24.

60. Langsjoen PH, Langsjoen AM. Comparison study of plasma coenzyme Q10 levels in healthy subjects supplemented with ubiquinol versus ubiquinone. *Clin Pharmacol Drug Dev*. 2014;3(1):13-7.

61. Kubo H, Yamamoto Y, Fujisawa A. Orally ingested ubiquinol-10 or ubiquinone-10 reaches the intestinal tract and is absorbed by the small intestine of mice mostly in its original form. *J Clin Biochem Nutr*. 2023;72(2):101-6.

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