

Pharmacological Regimen of Coenzyme Q10

Simranjeet Kaur*

Faculty of Pharmaceutical Sciences, PCTE Group of Institutes, Baddowal- 141012, Ludhiana, Punjab, India

***Corresponding Author:** Simranjeet Kaur, Faculty of Pharmaceutical Sciences, PCTE Group of Institutes, Baddowal- 141012, Ludhiana, Punjab, India.

Received: February 15, 2023; **Published:** March 30, 2023

The coenzyme Q-10 is possessing antioxidant property, high degree of hydrophobicity and occurrence in biological system, it plays an important role in cellular defense against oxidative damage. The coenzyme Q-10 is mainly present in brain, kidney, heart and liver and it is indicated in the treatment of cardiovascular disease, neurodegenerative disorders, cancer, diabetes mellitus, aging, alzheimer disease and kidney disease.

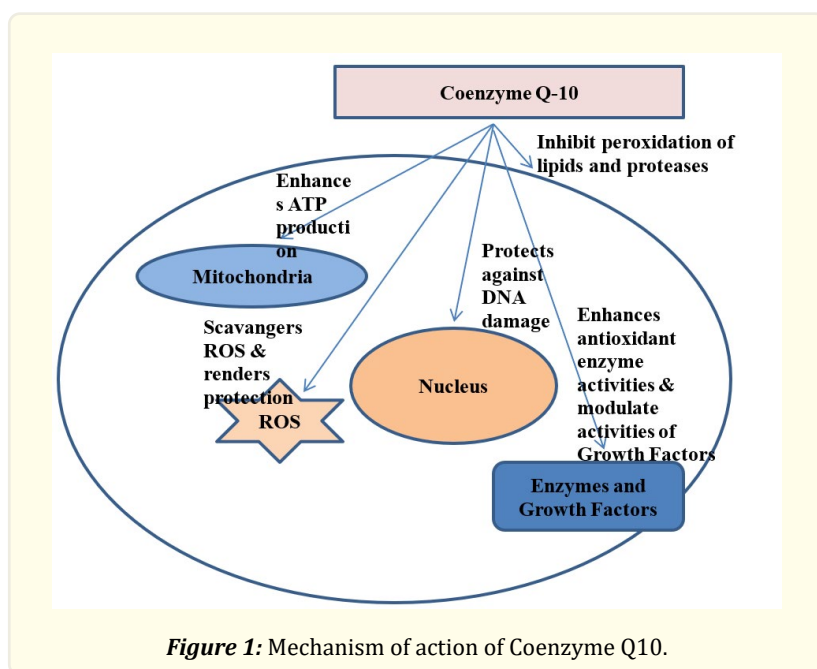
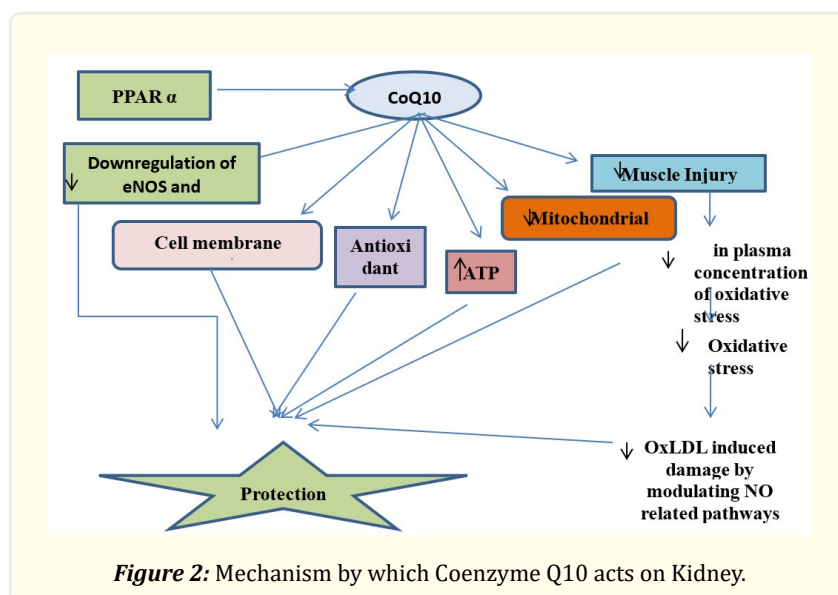


Figure 1: Mechanism of action of Coenzyme Q10.

Coenzyme Q-10 is a fat soluble vitamin like quinone commonly known as ubiquinone, CoQ, Vit Q 10. It is used in the treatment of a variety of disorders primarily related to suboptimal cellular energy metabolism and oxidative injury [1, 2]. Coenzyme Q-10 or ubiquinone is ubiquitous in nature and is widely distributed in plants, animals and micro-organisms. Coenzyme Q 10 was first isolated in 1957 in beef mitochondria and is found in highest concentrations in tissues with high energy turn over such heart, brain, liver and kidney [3, 4]. The primary role of coenzyme Q-10 is to facilitate the electron transfer between redox components of electron transport chain in order to create a proton gradient across the inner mitochondrial membrane thereby facilitating the ATP formation [5]. Additional biological functions maintenance of membrane fluidity, recycling of radicals forms of vitamin C and E [6, 7]. It is most importantly antioxidant protection against membrane lipid peroxidation [8]. It is the only endogenously occurring lipid soluble antioxidant among all the coenzymes ubiquitously synthesized [9, 10].

Coenzyme Q-10 also appears to increase adenosine triphosphate levels by preventing the loss of adenine nucleotide pool from cardiac cells [11]. Additionally, coenzyme Q-10 has demonstrated activity in preventing lipid peroxidation as an antioxidant scavenger and an indirect stabilizer of calcium channels to decrease calcium overload [12, 13]. Coenzyme Q-10 has been cleared for the treatment of congestive heart failure. (CHF) These levels appear to correlate with CHF severity in the animal and human model, with coenzyme Q-10 supplementation protecting against ischemia and reperfusion injury in animal studies [14, 15] CoQ 10 affects the function of all cells in the body, making it essential for the health of all tissues and organs. CoQ 10 also functions as an intracellular antioxidant at the mitochondrial level, perhaps accounting for its benefit in neurodegenerative disease, male infertility, periodontal disease, cardiovascular disease, diabetes mellitus, cancer, asthma, renal failure, migraine and immune [9] fenofibrate is a PPAR- α agonist, it increases the expression of lipolytic enzymes and reduces oxidative stress and lipid accumulation thus inhibit the development of albuminuria and glomerular fibrosis [16] (Tanaka et al, 2011).

CoQ10 decreases the oxLDL mediated down-regulation of eNOS and up-regulation of iNOS [17] (Tsai KL et al, 2011).



References

1. Greenberg S and Frishman WH. "Co-enzyme Q10; a new drug for cardiovascular disease". *J Clin Pharmacol* 30 (1990): 596-608.
2. Tran MT, et al. "Role of coenzyme Q10 in chronic heart failure, angina and hypertension". *Pharmacotherapy* 21 (2001): 797-806.
3. Crane FL, et al. "Isolation of a quinone from beef heart mitochondria". *Biochim Biophys Acta* 25 (1957): 220-221.
4. Shunk, C.H., et al. Coenzyme Q and its biosynthesis 59 (1998): 150-159.
5. Fernandez-Ayala, et al. "Specificity of coenzyme Q10 for a balanced function of respiratory chain and endogenous ubiquinone biosynthesis in human cells". *Biochem Biophys Acta* 1706 (2005): 174-183.
6. Beyer RE., et al. "Tissue coenzyme Q and protein concentrations over the life span of the laboratory rat". *Mech Ageing Dev* 32 (1995): 267-281.
7. Malchair P, et al. "Coenzyme Q10: biochemistry, pathophysiology of its deficiency and potential benefit of an increased intake". *Rev Med Liege* 60 (2005): 45-51.
8. Al-Thakafy HS., et al. "Alterations of erythrocyte free radical defense system, heart tissue lipid peroxidation and lipid concentration in streptozotocin induced diabetic rats under coenzyme Q10 supplementation". *Saudi Med J* 25 (2005): 1824-1830.
9. Battino M Giunta, et al. "Coenzyme Q10, antioxidant status and ApoE isoforms". *Biofactors* 18 (2003): 299-305.

10. Molyneux SL, et al. "Biological variations of coenzyme Q10". Clin Chem 51 (2005): 445-457.
11. Ito H, et al. "Coenzyme Q10 attenuates cyanide activation of the ATP sensitive potassium channel current in single cardiac myocytes of the guinea pig". Nahrungsmittelwissenschaft Arch Pharmacol 344 (1991): 133-6.
12. Sugiyama S, et al. "Antioxidant effect of coenzyme Q10". Experientia 36 (1980): 10002-3.
13. Nayler WG. "The use of coenzyme Q10 to protect ischemia heart muscle". In: Yamamura Y; Folkers K, Ito Y, eds. Biomedical and clinical aspects of coenzyme Q Amsterdam: Elsevier 2 (1980): 409-25.
14. Boler JB, et al. "Deficiency of coenzyme Q10 in the rabbit". Int Z vitaminforsch 39 (1969): 281-8.
15. Mortensen SA. "Perspective on therapy of cardiovascular disease with coenzyme Q10". Clin Investing 71 (1993): 116-23.
16. Tanaka Y, Shinji kume and Shin chi araki. "Fenofibrate, a PPAR α agonist, renoprotective effect". kidney international 79 (2011): 871-882.
17. Tsai KL, et al. "A novel mechanism of coenzyme Q10 protects against human endothelial cells from oxidative stress-induced injury by modulating NO-related pathways". Journal of nutritional chemistry 23 (2011): 458-468.

Volume 4 Issue 4 April 2023

© All rights are reserved by Simranjeet Kaur.