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# Meschino Health Comprehensive Guide to Accessory Nutrients and Essential Oils



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## About the Meschino Health Comprehensive Guide to Accessory Nutrients and Essential Oils

The Meschino Health Comprehensive Guide to Vitamins is one of four eBooks on nutrients written by Dr. James Meschino:

1. Meschino Health Comprehensive Guide to Vitamins
2. Meschino Health Comprehensive Guide to Herbs
3. Meschino Health Comprehensive Guide to Minerals
4. Meschino Health Comprehensive Guide to Accessory Nutrients and Essential Oils

All four books were written to both educate and provide an easy to use quick reference to answer important questions regarding nutrients. Users of the guide can quickly find which health conditions the nutrient can impact, proper dosage, possible effects of a deficiency or the effect any potential toxicity associated with the nutrient. Finally any drug-nutrient Interactions associated with the nutrient.

More eBook and eQuick Guides

Meschino Health is excited to be able to provide tools and resources to help you achieve your healthy living objectives. Sharing the Healthy Living message and helping anyone who is interested in living a healthy happy life is what Meschino Health is all about. Visit [www.MeschinoHealth.com](http://www.MeschinoHealth.com) to learn the latest a science based research on diet and supplementation that can prevent and treat health conditions often associated with aging. New eBooks and eGuides are added every month and can be downloaded free of charge.

## Meschino Health Natural Health Assessment

Welcome to the Nutrition, Lifestyle and Anti-aging Assessment.



The most powerful health assessment on the internet

- Easy to Complete Online Questionnaire
- Your Personal Health Assessment is generated Instantly and can be downloaded to your computer
- The Meschino Health Assessment is a 15 to 20 page comprehensive report complete with diet, lifestyle and supplement considerations that are specific to your profile.

The Meschino Health Assessment is a free service created by Dr. James Meschino. The feedback in your report is based on your answers to the questions in the Health Assessment, and highlights the dietary, lifestyle and supplementation practices that are best suited to your circumstances, according to currently available scientific studies

The Meschino Health Assessment is a Free Service

### Why take it?

We all know that we should eat better, exercise more and change some of our less than desirable lifestyle habits. Did you know that 7 out of 10 North Americans are taking some form of nutritional supplements to augment their diet? While that might sound like good news, the downside is that many people are guessing at what supplements to take! So which one should you take? Better yet, what does eating better look like?

### You need a plan.

But where would you even begin to find a health assessment that takes into account your personal health status, diet, lifestyle activities and family health history-before recommending a plan of action?

Where? [Right here.](#)

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## Coenzyme Q<sub>10</sub> (*Ubiquinone*)

### General Features

CoQ<sub>10</sub> is an essential component of the electron transfer system in the mitochondria. More specifically, it functions to shuttle hydrogen electrons from NAD to cytochrome b, facilitating the release of energy required to recouple ADP with inorganic phosphate in the synthesis of ATP. As such, CoQ<sub>10</sub> is an integral part of the bioenergetic system that enables cells to produce adequate amounts of ATP through aerobic pathways. ATP is the primary fuel required to power the body's metabolic reactions, maintain optimal function of cells and sustain life. A deficit in ATP synthesis can compromise any number of energy-dependent cellular functions and hasten the onset of dysfunction and if severe enough, cell death.

Although the body can synthesize CoQ<sub>10</sub>, deficiency states of CoQ<sub>10</sub> tend to exist and are associated with various health conditions. Moreover, supplementation studies with CoQ<sub>10</sub> have been shown to effectively treat and sometimes reverse a number of these conditions.

There is evidence that a decline in CoQ<sub>10</sub> synthesis occurs with aging, predisposing individuals to a number of CoQ<sub>10</sub> deficiency-related disorders and diseases.

Professor F.L. Crane and his colleagues at the University of Wisconsin first discovered CoQ<sub>10</sub> in 1957. Since then, Dr. Karl Folkers at the University of Texas (Austin) is most responsible for the ongoing research on CoQ<sub>10</sub>.<sup>1-4</sup>

Coenzyme Q<sub>10</sub> is also a fat soluble antioxidant, which has been shown to reduce oxidation of LDL-cholesterol and the mitochondrial DNA.<sup>5,6</sup> CoQ<sub>10</sub> supplementation has been shown to modulate immune system function, enhancing levels of immunoglobulin G (IgG), in the serum of patients provided with 60 mg CoQ<sub>10</sub> per day.<sup>7</sup>

The average person may consume about 5 mg per day of CoQ<sub>10</sub> from foods, with the main sources being meat, fish, soybeans and some vegetable oils. Clinical Coenzyme Q<sub>10</sub> studies have involved daily supplemental intake levels ranging from 60 mg to 300 mg per day; far greater than food alone can provide.<sup>8</sup>

### Supplementation Studies and Clinical Applications

#### 1. Cardiovascular Disease

Biopsy results from heart tissue in patients with various cardiovascular diseases (especially congestive heart failure) show a deficiency in CoQ<sub>10</sub> in 50 to 75 percent of cases. Low blood levels of CoQ<sub>10</sub> are also a consistent finding in the majority of these patients.<sup>9-12</sup>

Supplementation studies with patients suffering from various cardiomyopathies (i.e. ischemic cardiomyopathy, dilated cardiomyopathy, heart valve disorders) and congestive heart failure have demonstrated significant improvement in heart function (according to the New York Heart Association functional scale) in a high percentage of cases.

Many patients in these studies have been able to reduce the number of cardiac drugs required (1-3 medications reduced in 43 percent of CoQ<sub>10</sub> supplemented patients in one study of 424 patients, over an eight year period).

Heart function parameters monitored have included left ventricular wall thickness, mitral valve inflow slope, and fractional shortening.<sup>13,14,15</sup>

Congestive Heart Failure - Several controlled studies using Coenzyme Q<sub>10</sub> supplementation in patients with congestive heart failure have demonstrated significant improvement in cardiac ejection fraction, reduced shortness of breath, and increased muscle strength. Other studies have demonstrated increased stroke volume and cardiac index, improved survival and improved quality of life, in general. Of great significance is the fact that when patients discontinue CoQ<sub>10</sub> supplementation, cardiac function deteriorates. Thus, CoQ<sub>10</sub> needs to be a lifelong intervention in these cases.<sup>16,17,18</sup>

Angina - A small study has shown that CoQ<sub>10</sub> supplementation can reduce angina episodes and nitroglycerine use and improve treadmill exercise tolerance. Larger trials are required to substantiate this data.<sup>19</sup>

Hypertension - Several studies have provided evidence that CoQ<sub>10</sub> supplementation can lower systolic and diastolic blood pressure (i.e. systolic 164-146, diastolic 98-86) in hypertensive patients. Improved blood cholesterol levels also occurred in one study, with a rise in HDL and a reduction in total cholesterol from 229.9 mg/dl to 213.3 mg/dl.<sup>18-22</sup>

## 2. Periodontal Disease

Several intervention trials involving patients with periodontal disease have revealed that CoQ<sub>10</sub> supplementation can be useful in reversing the condition to various degrees.<sup>23,24</sup>

## 3. Aerobic Exercise Performance

A study of 25 cross-country skiers has provided preliminary evidence that CoQ<sub>10</sub> supplementation may improve exercise performance in endurance athletes.<sup>25</sup>

Sedentary individuals have also demonstrated improvement with work capacity, oxygen consumption, fat burning and oxygen transport after beginning an exercise program. The group supplemented with CoQ<sub>10</sub> demonstrated greater improvement in these aerobic parameters compared to the placebo group, in a 4-8 week trial period.<sup>26</sup>

## Dosage Ranges

1. Cardiovascular Conditions: typical dosage is 50-60 mg, three times per day. Large doses (up to 300 mg) may be needed in severe heart disease. Some studies use a dosage of 2 mg CoQ<sub>10</sub> for each kilogram of body weight.
2. Periodontal Disease: 50 mg per day has been used in clinical trials
3. Exercise Performance Studies: 60 mg per day<sup>27</sup>

## Toxicity and Contraindications

Coenzyme Q<sub>10</sub> is well tolerated, and no serious adverse effects have been reported with long-term use.<sup>27</sup>

## Drug-Nutrient Interactions

### 1. Warfarin

CoQ<sub>10</sub> supplementation has been shown to antagonize the anti-coagulant effects of warfarin, requiring dose adjustment.

### 2. Beta-Blockers and HMG

CoA Reductase (statin) drugs for cholesterol lowering are known to inhibit the endogenous synthesis of CoQ<sub>10</sub>. CoQ<sub>10</sub> supplementation can compensate for this inhibition effect and is indicated as a concurrent therapy when these drugs are in use (100 mg of CoQ<sub>10</sub> per day is recommended in these cases).

### 3. Psychotropic Drugs

Coenzyme Q<sub>10</sub> supplementation has been shown to reduce the cardiac side effects of phenothiazines and tricyclic antidepressant drugs.

### 4. Chemotherapy

Q<sub>10</sub> supplementation can mitigate the cardiac side effects and cardiotoxicity of the chemotherapy drug known as adriamycin (100 mg per day of CoQ<sub>10</sub> supplementation). Even children with lymphoblastic leukemia or non-Hodgkin lymphoma realized this benefit compared with the placebo group.<sup>28-34</sup>

### 5. Anticoagulants

As noted above, there are reported cases of CoQ<sub>10</sub> countering the action of warfarin. Thus, the physician prescribing warfarin may need to adjust the warfarin dose if CoQ<sub>10</sub> is to be used and therefore must be consulted.<sup>34,35</sup>

The following drugs may reduce the body's levels of CoQ<sub>10</sub>:

1. Orlistat: reduces CoQ<sub>10</sub> absorption.<sup>36</sup>
2. Beta blockers: decreases CoQ<sub>10</sub> synthesis.<sup>37</sup>
3. Biguanides: decreases CoQ<sub>10</sub> synthesis.<sup>38</sup>
4. Clondine: decreases CoQ<sub>10</sub> synthesis.<sup>39</sup>
5. Gemfibrozil: (cholesterol-lowering drug).<sup>40</sup>
6. Haloperidol: decreases CoQ<sub>10</sub> synthesis.<sup>41</sup>
7. HMG-CoA Reductase inhibitors: decreases CoQ<sub>10</sub> synthesis.<sup>42</sup>
8. Hydralazine: decreases CoQ<sub>10</sub> synthesis.<sup>37</sup>
9. Methylopa: decreases CoQ<sub>10</sub> synthesis.<sup>39</sup>
10. Phenothiazines: decreases CoQ<sub>10</sub> synthesis.<sup>41</sup>
11. Sulfonyleureas: some of these drugs decrease CoQ<sub>10</sub> synthesis (acetohexamide, glyburide, tolazamide).<sup>38</sup>
12. Thiazide Diuretics: decrease CoQ<sub>10</sub> synthesis.<sup>39</sup>
13. Tricyclic Antidepressants: decrease CoQ<sub>10</sub> synthesis.<sup>41</sup>

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1. Folkers K, Yamamura Y, editors. Biomedical and clinical aspects of coenzyme Q. Vol 1. Amsterdam: Elsevier/North-Holland Biomedical Press; 1977.
  2. Yamamura Y, Folkers K and Ito Y, editors. Biomedical and clinical aspects of coenzyme Q. Vol 2. Amsterdam, Holland; Elsevier/North-Holland Biomedical Press; 1980.
  3. Folkers K, Yamamura Y, editors. Biomedical and clinical aspects of coenzyme Q. Vol 3. Amsterdam: Elsevier/North-Holland biomedical Press; 1981.
  4. Folkers K, Yamamura Y, editors. Biomedical and clinical aspects of Coenzyme Q. Vol 4. Amsterdam: Elsevier Science Publ; 1984.
  5. Frei B, Kim MC, Ames BN. Ubiquinol-10 is an effective lipid-soluble antioxidant at physiological concentrations. *Proc Natl Acad Sci* 1990;87:4879-83.
  6. [Weber C](#), [Jakobsen TS](#), [Mortensen SA](#), [Paulsen G](#), [Holmer G](#). Effect of dietary coenzyme Q10 as an antioxidant in human plasma. *Mol Aspects Med* 1994;15(Suppl):97S-102S.
  7. Folkers K, et al. Increase in levels of IgG in serum of patients treated with coenzyme Q10. *Res Commun Chem Pathol Pharmacol* 1982;38:335.
  8. Reavely N. *New encyclopedia of vitamins, minerals, supplements and herbs*. New York: Evans M and Company, Inc.; 1998. p. 353-61.
  9. Kitamura N, Yamaguchi A, Otaki M, Sawatani O, Minoji T, Tamura H, et al. Myocardial tissue level of coenzyme Q10 in patients with cardiac failure. In: Folkers K, Yamamura Y, editors. *Biomedical and Clinical Aspects of Coenzyme Q*, Vol 4. Amsterdam, Holland: Elsevier Science, Publ; 1984. p. 243-52.
  10. Littarru GP, Ho L, Folkers K. Deficiency of coenzyme Q10 in human heart disease, Part II. *Int J Vit Nutr Res* 1972;42:413.
  11. Folkers K, et al. Evidence for a deficiency of coenzyme Q10 in human heart disease. *Int J Vit Res* 1970;40:380.
  12. Folkers K, Vadhavavikit S, Mortensen SA. Biochemical rationale and myocardial tissue data on the effective therapy of cardiomyopathy with coenzyme Q10. *Proc Natl Acad Sci* 1985;82:901.
  13. Langsjoen H, Langsjoen P, Langsjoen P, Willis R, Folkers K. Usefulness of coenzyme Q10 in clinical cardiology: A long-term study. *Mol Aspects Med* 1994;1(Suppl):165S-75S.



14. Tsuyusaki T, Noro C, Kikawada R. Mechanocardiography of ischemic or hypertensive heart failure. In: Yamamura Y, Folkers K, Ito Y, editors. *Biomedical and clinical aspects of coenzyme Q*. Vol 2. Amsterdam, Holland; Elsevier/North-Holland Biomedical Press; 1980. p. 273-88.
15. Judy WV, Hall JT, Toth PD, Folkers K. Myocardial effects of co-enzyme Q10 in primary heart failure. In: Folkers K, Yamamura Y, editors. *Biomedical and clinical aspects of Coenzyme Q*. Vol 4. Amsterdam: Elsevier Science Publ; 1984. p. 281-90.
16. Langsjoen PH, Vadhanavikit S, Folkers K. Response of patients in classes III and IV of cardiomyopathy to therapy in a blind and crossover trial with coenzyme Q10. *Proc Natl Acad Sci* 1985;82:4240-4.
17. Morisco C, Trimarco B, Condorelli M. Effect of CoQ10 therapy in patients with congestive heart failure: A long-term multicenter randomized study. *Clin Invest* 1993;71(Suppl):134S-6S.
18. Baggio E, Gandini R, Plancher AC, Passeri M, Carosino G. Italian multicenter study on the safety and efficacy of coenzyme Q10 as adjunctive therapy in heart failure. *CoQ10 Drug Surveillance Investigators. Mol Aspects Med* 1994;15(Suppl):287S-94S.
19. Kamikawa T, Kobayashi A, Yamashita T, Hayashi H, Yamazaki N. Effects of coenzyme Q10 on exercise tolerance in chronic stable angina pectoris. *Am J Cardiol* 1985;56:247-51.
20. Digiesi V, Cantini F, Brodbeck B. Coenzyme Q10 in essential hypertension. *Mol Aspects Med* 1994;15(Suppl):257S-63S.
21. Langsjoen P, et al. Treatment of essential hypertension with coenzyme Q10. *Mol Aspects Med* 1994;15:265-72.
22. Digiesi V, Cantini F, Bisi G, et al. Mechanism of action of coenzyme Q10 in essential hypertension. *Curr Thes Res* 1992;51:668-72.
23. Nakamura R, Littrau GP, Folkers K, Wilkinson EG. Study of Co Q10 enzymes in gingiva from patients with periodontal disease and evidence for a deficiency of coenzyme Q10. *Proc Natl Acad Sci* 1974;71:1456.
24. Wilkinson EG, Arnold RM, Folkers K. Treatment of periodontal and other soft tissue diseases of the oral cavity with coenzyme Q10. In: Folkers K, Yamamura Y, editors. *Biomedical and clinical aspects of coenzyme Q*. Vol 1. Amsterdam: Elsevier/North-Holland Biomedical Press; 1977. p. 251-65.
25. Ylikoski T, Piirainen J, Hanninen O, Penttinen J. The effect of coenzyme Q10 on the exercise performance of cross-country skiers. *Mol Aspects Med* 1997;18(Suppl):283S-90S.
26. Vanfraechem JHP, Folkers K. Coenzyme Q10 and physical performance. In: Folkers K, Yamamura Y, editors. *Biomedical and clinical aspects of coenzyme Q*. Vol 3. Amsterdam: Elsevier/North-Holland biomedical Press; 1981. p. 235-41.
27. Murry M. *Encyclopedia of Nutritional Supplements*. Rocklin, CA: Prima Publishing; 1996. p. 296-308.
28. Mortensen SA, Leth A, Agner E, Rohde M. Dose-related decrease of serum coenzyme Q10 during treatment with HMG-CoA reductate inhibitors. *Mol Aspects Med* 1997;18(Suppl):137S-44S.
29. Bargossi AM, Grossi G, Fiorella PL, Gaddi A, Di Giulio R, Battino M. Exogenous CoQ10 supplementation prevents plasma ubiquinone reduction induced by HMG-CoA reductate inhibitors. *Mol Aspects Med* 1994;15(Suppl):187S-93S.
30. Hamada M, Kazarani Y, Ochi T, Ito T, Kokubu T. Correlatio between serum CoQ10 level and myocardial contractility in hypertensive patients. In: Folkers K, Yamamura Y, editors. *Biomedical and clinical aspects of Coenzyme Q*. Vol 4. Amsterdam: Elsevier Science Publ; 1984. p. 263-70.
31. Judy WV, Hall JH, Dugan W, Toth PD, Folkers K. Coenzyme Q10 reduction in adriamycin cardiotoxicity. In: Folkers K, Yamamura Y, editors. *Biomedical and clinical aspects of Coenzyme Q*. Vol 4. Amsterdam: Elsevier Science Publ; 1984. p. 231-41.
32. Iarussi D, Auricchio U, Agretto A, Murano A, Giuliano M, Casale F, et al. Protective effect of coenzyme Q10 on anthracyclines cardiotoxicity: Control study in children with acute lymphoblastic leukemia and non-Hodgkin lymphoma. *Mol Aspects Med*, 1994;15(Suppl):207S-12S.
33. Kishi T, Makino K, Okamoto T, Kishi H, Folkers K. Inhibition of myocardial respiration by psychotherapeutic drugs and prevention by coenzyme Q. In: Yamamura Y, Folkers K and Ito Y, editors. *Biomedical and clinical aspects of coenzyme Q*. Vol 2. Amsterdam, Holland; Elsevier/North-Holland Biomedical Press; 1980. Pg. 139-54.
34. Heck AM, Deweitt BA, Lukes AL. Potential interactions between alternative therapies and warfarin. *Am J Health Syst Pharm* 2000;57(13):1221-30.
35. Landbo C, Almdal TP. Interaction between warfarin and coenzyme Q10. *Ugeskr Laeger*. 1998;160(22):3226-7.
36. Xenical (orlistat), Product Prescribing Information. Nutley, NJ: Roche Laboratories, Inc., Sept 2000.
37. Kishi T, Watanabe T, Folkers K. Bioenergetics in clinical medicine XV. Inhibition of coenzyme Q10-enzymes by clinically used adrenergic blockers of beta-receptors. *Res Commun Chem Pathol Pharmacol*. 1977;17(1):157-64.
38. Kishi T, Kishi H, Watanabe T, Folkers K. Bioenergetics in clinical medicine XI. Studies on coenzyme Q and Diabetes Mellitus. *J Med* 1976;7(3):307-21.
39. Kishi H, Kishi T, Folkers K. Bioenergetics in clinical medicine. III. Inhibition of coenzyme Q10-enzymes by clinically used anti-hypertensive drugs. *Res Commun Chem Pathol Pharmacol* 1975;12(3):533-40.
40. [Aberg F](#), [Appelkvist EL](#), [Bröijersén A](#), [Eriksson M](#), [Angelin B](#), [Hjemdahl P](#), et al. Gemfibrozil-induced decrease in serum Ubiquinone and alpha- and gamma-tocopherol levels in men with combined hyperlipidaemia. *Eur J Clin Invest* 1998;28(3):235-42.
41. Kishi T, Makino K, Okamoto T, Kishi H, Folkers K. Inhibition of myocardial respiration by psychotherapeutic drugs and prevention by coenzyme Q10. In: *Biomedical and clinical aspects of coenzyme Q10*. Yamamura Y, Folkers K, Ito Y, editors. Vol 2. Amsterdam: Elsevier/North-Holland Biomedical Press; 1980. p. 139-54.

42. [Ghirlanda G](#), [Oradei A](#), [Manto A](#), [Lippa S](#), [Uccioli L](#), [Caputo S](#), et al. Evidence of plasma Co<sub>10</sub>- lowering effect by HMG-CoA reductase inhibitors: a double-blind, placebo-controlled study. *J Clin Pharmacol* 1993;33(3):226-9.

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## Creatine

### General Features

It is now widely accepted that Creatine supplementation can increase muscle strength and mass.<sup>1,2,3,4</sup> Creatine is an amino acid that is stored in muscle in the form of Creatine phosphate. During explosive or intensive exercise, Creatine phosphate is broken down by a specific enzyme to yield Creatine, plus phosphate, plus free energy. The free energy released from the breakdown of Creatine phosphate is used to regenerate ATP, which is the fuel that powers muscle contraction.<sup>2</sup>

The normal daily requirement for Creatine is about 2 grams for a person weighing 70 kg. Animal protein (especially meats) normally provides at least half that amount, with approximately one gram per day synthesized by the liver. A half-pound of raw meat contains about 1 gram of Creatine, but fish is also a good source.

A number of recent studies have demonstrated that short-term Creatine supplementation increases Creatine phosphate stores in skeletal muscle by 10% to 40%.<sup>3</sup> This in turn leads to an increase in muscle mass, which is thought to occur from increased protein synthesis, as the muscle lays down an increased number of contractile myofilaments (protein bands that contract and generate force). Increased muscular fluid retention may also participate in muscle volume gains with Creatine use.<sup>5,6,7</sup> Creatine has also been shown to provide antioxidant properties. This may be of some significance as free radicals generated from exercise can affect muscle fatigue and protein turnover.<sup>24</sup>

It also appears that Creatine supplementation may allow athletes to train harder (due to increased available energy for muscle concentration), which promotes strength gains, and increases muscle size due to hypertrophy (larger muscle fiber size).<sup>2,3</sup>

The established protocol for Creatine supplementation used by athletes involves a loading dosage of 20 to 25 grams per day for the first 5 to 7 days. Typically an athlete will mix a heaping teaspoon of Creatine monohydrate crystals into a glass of juice to obtain about 5 grams of Creatine. During the loading phase the athlete does this on 4 or 5 occasions throughout the day to attain an intake of 20-25 grams. After the loading phase is completed, the maintenance daily dosage is usually 5 to 10 grams per day. Recent reports suggest that taking Creatine with glucose (a simple carbohydrate) may increase the amount of Creatine absorbed by the muscles. As such, some manufacturers combine Creatine with carbohydrates in a premix product to help improve Creatine delivery to muscles.<sup>25</sup>

### Clinical Application and Mechanism of Action

#### 1. Increased Strength and Performance In Athletes

Several studies have shown that Creatine supplementation improves performance in repeated bouts of high intensity strength work and repeated sprints, which are requirements for many sports.<sup>8,9,10,11,12,13,14,16,17,18</sup> In short, substantial evidence suggests that Creatine supplementation increases lean body mass, muscular strength, and sprint power.<sup>24</sup>

Significant gains in strength and lean mass often occur in the first 6 weeks of Creatine supplementation, when combined with proper training and diet. In one study, college football players who took Creatine supplements for 28 days during resistance and agility training had significant gains in lean mass when compared to players who took the placebo.<sup>15</sup>

Individuals may vary in their response to Creatine supplementation, but it is not uncommon to see a 5 to 10 lb. increase in weight within the first six weeks.

Approximately 80% of Creatine studies have reported a performance-enhancing effect. This is quite impressive when you consider the fact that Creatine is not structurally or functionally related to anabolic steroids, and that Creatine supplements are not banned by the International Olympic Committee or the National Collegiate Athletic Association. Creatine use is based on the same principle as carbohydrate loading in that an athlete is manipulating their dietary intake to optimize muscle Creatine phosphate stores for more explosive power and enhanced performance.

Athletes requiring repeated bouts of explosive power may also benefit from Creatine supplementation as demonstrated by M. Izquierdo et al. Among other positive benefits revealed in their study of nineteen trained athletes, they showed that short-term Creatine supplementation (20 gms per day for 5 days) enhanced repeated sprint performance and attenuated decline in jumping ability after repetitive high-power-output exercise bouts (MRPB).<sup>22</sup> Similar results have been documented by G. Cottrell et al, in subjects performing repeated sprint cycling.<sup>23</sup> These studies have important implications for many sports such as hockey, basketball, soccer, volleyball, lacrosse, football, tennis and any sport requiring repeated bouts of all-out lower extremity explosive power and/or jumps.

## 2. Neuromuscular Diseases

Creatine supplementation in humans has been reported to enhance power and strength both in normal subjects and in patients with various neuromuscular diseases.<sup>14</sup> Clinical studies in patients with ALS (amyotrophic lateral sclerosis).<sup>14</sup> Huntington's disease, Parkinson's disease, Duchenne muscular dystrophy, McArdles disease<sup>15</sup> and Myasthenia Gravis<sup>16</sup> have shown that Creatine supplementation can produce an increase in strength and thus, provide symptomatic treatment and improved quality of life for many of these patients.<sup>14,15,16</sup>

## 3. Heart Failure

Creatine supplementation has been shown to improve exercise capacity in patients with heart failure in some studies. Along with Coenzyme Q10, hawthorn extract and L-carnitine, Creatine is one of few natural health products that is shown to reverse certain parameters of heart failure. As reported by K. Witte et al, there is evidence for a possible role for micronutrient deficiency in heart failure, of which Creatine may be one of the principle factors.<sup>10,11,17</sup>

## 4. Musculoskeletal Rehabilitation

Creatine was shown to speed recovery of muscular power in a double blind, placebo-controlled study involving 20 male and female students whose right legs were immobilized in casts for a period of two weeks. Those given Creatine supplementation during and after leg immobilization displayed more muscular power and greater muscle size after three to ten weeks of physical rehabilitation than did subjects who took the placebo.<sup>18</sup>

## 5. Anti-Aging in Older Subjects

Creatine supplementation provided to active subjects over 70 years of age, and subjects 59-72 years of age, have resulted in significant gains in several indices of muscle performance including increased maximal dynamic and isometric strength, lower body mean power, lower extremity functional capacity, increased fat-free mass, increased lean mass and endurance power. These studies suggest that Creatine supplementation may help to forestall or reverse muscular atrophy and progressive weakness that occurs during aging, and that Creatine may be useful as an intervention to improve the ability of certain elderly individuals to perform functional living tasks, decreasing dependency and, enhancing their quality of life.<sup>19,20</sup>

Other studies have noted that younger individuals respond to Creatine supplementation more efficiently than do older subjects in that muscular phosphocreatine stores were shown to increase on average by 35% in young subjects (~24 years of age) and 7% in older subjects (~70 years of age) after five days of Creatine supplementation

(20 gms per day). As such, it may take a longer period to maximize Creatine stores in older subjects with Creatine supplementation.<sup>21</sup>

### Absorption and Utilization

Creatine absorption from the intestinal tract is very efficient. Studies show that a 6-8 gm oral load of Creatine results in approximately 50% of the ingested Creatine being excreted in the urine. Thus, researchers are still working to identify the ideal single, daily and cumulative doses of Creatine for various applications.<sup>26</sup> Other studies demonstrate that a 5 gm oral load of Creatine, followed by 93 gm oral load of simple carbohydrate in solution ( water ) at 30 minutes post-Creatine intake (4-times per day), resulted in a 60% increase in total muscle Creatine compared to subjects ingesting the same amount of Creatine in the absence of a simple carbohydrate drink. Subjects ingesting Creatine and the simple carbohydrate drink had higher insulin levels and significantly less Creatine lost in their urine, indicating that higher insulin levels are likely a key to greater muscle uptake and utilization of Creatine, and a reduction in urinary loss. Thus, it is accepted that Creatine utilization is enhanced by concurrent ingestion of a simple carbohydrate drink (e.g. fruit juice).<sup>25</sup>

Additionally, concurrent administration of Creatine and glycogen reveal that Creatine supplementation enhances muscle levels of glycogen (glycogen supercompensation) beyond that attainable from glycogen loading alone. As supercompensation of muscle glycogen is also an ergogenic factor in exercise performance, the combination of Creatine and carbohydrate loading appear to improve performance by increasing muscle Creatine and muscle glycogen.<sup>27</sup>

### Dosage and Standardized Grade (2:1 powdered extract)

1. Athletic Performance (strength, sprint and repeated sprint power, lean mass etc.) - The usual protocol is 5 gm, 4-5-times per day for five consecutive days during the loading phase, followed by 5 gm, twice daily as the maintenance dose. Many athletes cycle one month on, one month off to prevent any possibility of toxicity and to prevent the body from compensating by reducing its own endogenous synthesis of Creatine in the liver.<sup>28,22,23,26,27</sup>
2. Neuromuscular Diseases / Amyotrophic Lateral sclerosis - One study used 20 gm per day in 5 gm divided doses for 7 days, followed by 3 gm per day for 3-6 months.<sup>14</sup> This dose may be appropriate for all neuromuscular diseases mentioned above in regards to adult supplementation, <sup>34</sup> although one patient with myasthenia gravis demonstrated significant improvement with 5 gm of Creatine per day combined with resistance training, 3 times per week.<sup>16</sup> In McArdles disease, a daily dose of Creatine of approximately 10 gm per day, followed by a maintenance dose of approximately 4 gm per day has been used successfully to increase muscular strength. <sup>35</sup> Other researchers have shown that a daily adult dose of 10 gm of Creatine per day, and a daily dose of 5 gm of Creatine for children, have been beneficial for individuals with various muscular dystrophies.<sup>39,40</sup>
3. Heart Failure - A daily dose of 20 gm of Creatine per day, in 5 gm divided doses, has shown good results over the 5-10 day test period. Participants showed improvement in strength, endurance and improved skeletal function upon exertion.<sup>17,36,37</sup>
4. Musculoskeletal Rehabilitation During and After Immobilization - Same dose as for athletic performance.<sup>18</sup>
5. Anti-aging in Older Patients - Same dose as for athletic performance.<sup>19,20</sup>

### Adverse Side Effects, Toxicity and Contraindications

As for the safety of Creatine supplementation, a 1997 study showed that short-term Creatine use (20 grams per day for 5 days) did not increase markers of kidney stress in five healthy men.<sup>13</sup> A study comparing Creatine users, for up to five years duration, to control subjects has shown that Creatine users have no remarkable differences in their Creatine,

urea, and plasma albumin clearances compared to controls. The researchers conclude that neither short-term, medium-term, nor long-term oral Creatine supplements induce detrimental effects in the kidney of healthy individuals.<sup>29,30,31</sup> To date no liver abnormalities have been evident in short-term Creatine challenge studies.<sup>30</sup> However, individuals with pre-existing kidney disease should be cautious as evidenced by the development of kidney dysfunction in a 25 year old soccer player taking Creatine who previously had been treated for focal segmental glomerulosclerosis of the kidney. His kidney function returned to normal after discontinuing Creatine supplementation.<sup>28</sup>

Some experts suggest that compulsory regular kidney and liver monitoring should accompany the use of Creatine due to the increased burden placed upon the liver and kidneys.<sup>30</sup> As pointed out by other experts, Creatine is normally found in cardiac muscle, brain, and testes, as well as skeletal muscle, and these former tissues have been largely unstudied with respect to the effects of Creatine supplementation.<sup>32</sup> The Food and Drug Administration (FDA) has advised athletes to consult a physician or a health care professional before embarking on any scheme of Creatine loading or supplementation.<sup>28</sup> Nevertheless, few reported adverse side effects from Creatine use have been reported despite its widespread use among young athletes, with Creatine sales reaching \$200 million in the U.S. in 1998.<sup>1</sup>

Other infrequently reported side effects include gastrointestinal disturbances and muscle cramps.<sup>30</sup>

In regards to children and younger athletes, the safety of Creatine supplementation has not yet been investigated in these individuals. Until all safety issues have been evaluated, experts strongly recommend against use of Creatine among children and adolescent athletes.<sup>33</sup>

Overall, Creatine supplementation appears to be safe for healthy adults. It's a low molecular weight compound that is excreted in the kidneys by simple diffusion. In the maintenance phase, athletes consume only slightly more Creatine (3-5 gm per day) than is generally found in the diet, which is usually about 2 gm per day.<sup>10,11</sup>

### **Drug-Nutrient Interactions**

There are no well-known drug-nutrient interactions for Creatine at this time.<sup>38</sup>

### Pregnancy and Lactation

During pregnancy and lactation, the only supplements that are considered safe include standard prenatal vitamin and mineral supplements. All other supplements or dose alterations may pose a threat to the developing fetus and there is generally insufficient evidence at this time to determine an absolute level of safety for most dietary supplements other than a prenatal supplement. Any supplementation practices beyond a prenatal supplement should involve the cooperation of the attending physician (e.g., magnesium and the treatment of preeclampsia.)

### References: Pregnancy and Lactation

1. Encyclopedia of Nutritional Supplements. Murray M. Prima Publishing 1998.
2. Reavley NM. The New Encyclopedia of Vitamins, Minerals, Supplements, and Herbs. Evans and Company Inc. 1998.
3. The Healing Power of Herbs (2<sup>nd</sup> edition). Murray M. Prima Publishing 1995.
4. Boon H and Smith M. Health Care Professional Training Program in Complementary Medicine. Institute of Applied Complementary Medicine Inc. 1997.

43. Kreider RB. Creatine, the next ergogenic supplement? Sports Science Training and Technology. Internet Society for Sports Science. Available at: <http://www.sportsci.org/traintech/creatine/rbk.html>. Accessed May 5, 1998
44. Kreider RB. Creatine supplement: analysis of ergogenic value, medical safety, and concerns. Journal of Exercise Physiology Online 1998; 1(1). Available at: <http://www.css.edu/users/tboone2/asep/jan3.html>. Accessed May 5, 1998
45. Bramberger M. The magic potion. Sports Illus 1998;88(16):58-65
46. Bessman SP, Savabi F. The role of the phosphocreatine energy shuttle in exercise and muscle hypertrophy, in: Taylor AW, Gollnick PD, Green HJ (eds.), International Series on Sport Sciences: Biochemistry of Exercise VII. Champaign, IL Human Kinetics 1988;19:167-78
47. Ingwall JS. Creatine and the control of muscle-specific protein synthesis in cardiac and skeletal muscle. Circ. Res 1976;38(5 suppl 1):1115-23
48. Sipila I, Rapola J, Simell O et al. Supplementary creatine as a treatment for gyrate atrophy of the choroids and retina. N Engl J Med 1981;304(5):867-70
49. Almada a, Kreider R, Ferreira M et al. Effects of calcium-HMB supplementation with or without creatine during training on strength and sprint capacity, abstract. FASEB J 1997;11:A374
50. Earnest CP, Snell PG, Rodriguez R et al. The effect of creatine monohydrate ingestion on anaerobic power indices, muscular strength and body composition. Acta Physiol Scand 1995;153(2):207-9
51. Burke LM, Pyne DB, Telford RD. Effect of oral creatine supplementation on single-effort sprint performance in elite swimmers. Int J Sports Nutr 1996;6(3):222-3
52. Dawson B, Cutler M, Moody A et al. Effects of oral creatine loading on single and repeated maximal short sprints. Aust J Sci Med Sports 1995;27(3):56-61
53. Redondo DR, Dowling EA, Graham BL et al. The effect of oral creatine monohydrate supplementaiton on running velocity. Int J Sports Nutr 1996;6(3):213-21
54. Kreider RB, Ferreira M, Wilson M et al. Effects of creatine supplementation on body composition, strength, and sprint performance. Med Sci Sports Exerc 1998;30(1):73-82
55. Poortmans JR, Auquier H, Renaut V et al. Effect of short-term creatine supplementation on renal responses in men. Eur J appl Physiol 1997;76(6):566-7
56. Mazzini L, Balzarini C, Colombo R, Mora G, Pastore I, De Ambrogio R et al. Effects of creatine supplementation on exercise performance and muscular strength in amyotrophic lateral sclerosis: preliminary results. J Neurol Sci 2001 Oct 15;191(1-2):139-44
57. Persky AM, Brazeau GA. Clinical pharmacology of the dietary supplement creatine monohydrate. Pharmacol Rev 2001 Jun;53(2):161-76

58. Stout JR, Eckerson JM, May E, Coulter C, Bradley-Popovich GE. Effects of resistance exercise and creatine supplementation on myasthenia gravis: a case study. *Med Sci Sports Exerc* 2001 Jun;33(6):869-72
59. Witte KK, Clark AL, Cleland JG. Chronic heart failure and micronutrients. *J Am Coll Cardiol* 2001 Jun1;37(7):1765-74
60. A leg to stand on. *Better Nutrition* May 2002;64(5):p20
61. Chrusch MJ, Chilibeck PD, Chad KE, Davison KS, Burke DG. Creatine supplementation combined with resistance training in older men. *Med Sci Sports Exerc* 2001 Dec;33(12):2111-7
62. Gotshalk LA, Volek JS, Staron RS, Denegar CR, Hagerman FC, Kraemer WJ. Creatine supplementation improves muscular performance in older men. *Med Sci Sports Exerc* 2002 Mar;34(3):537-43
63. Rawson ES, Clarkson PM, Price TB, Miles MP. Differential response of muscle phosphocreatine to creatine supplementation in young and old subjects. *Acta Physiol Scand*, 2002 Jan;174(1):57-65
64. Ezquierdo M, Ibañez J, González-Badillo JJ, Gorostiaga EM. Effects of creatine supplementation on muscle power, endurance, and sprint performance. *Med Sci Sports Exerc*, 2002 Feb;34(2):332-43
65. Cottrell G.T, Coast JR, Herb RA. Effect of recovery interval on multiple-bout sprint cycling performance after acute creatine supplementation. *J Strength Cond Res*, 2002 Feb;16(1):109-16
66. Lawler JM, Barnes, WS, Wu G, Song W, Demaree S. Direct antioxidant properties of creatine. *Biochem Biophys Res Commun*, 2002 Jan 11;290(1):47-52
67. Green AL, Hultman, E, Macdonald IA, Sewell DA, Greenhaff PL. Carbohydrate ingestion augments skeletal muscle creatine accumulation during creatine supplementation in humans. *Am J Physiol*, 1996 Nov;271(5 Pt 1):821-6
68. Burke DG, Smith-Palmer T, Holt LE, Head B, Chilibeck PD. The effect of 7 days of creatine supplementation on 24-hour urinary creatine excretion. *J Strength Cond Res*, 2001 Feb;15(1):59-62
69. Nelson AG, Arnall DA, Kokkonen J, Day R, Evans J. Muscle glycogen supercompensation is enhanced by prior creatine supplementation. *Med Sci Sports Exerc*, 2001 Jul;33(7):1096-100
70. Culpepper R Michael. Creatine supplementation: Safe as steak? *Southern Medical Journal*, Sep98;91(9):890-3
71. Poortmans, JR, Francaux, M. Long-term oral creatine supplementation does not impair renal function in healthy athletes. *Med Sci Sports Exerc*, 1999 Aug;31(8):1108-10
72. Poortmans, JR, Francaux, M. Adverse effects of creatine supplementation: fact or fiction? *Sports Med*, 2000 Sep;30(3):155-70
73. Schilling, B.K., Stone M.H., Utter, A., Kearney, J.T., Johnson, M., Coglianese, R., Smith, L., O'Bryant, H.S., Fry, A.C., Starks, M., Keith, R., Stone, M.E. Creatine supplementation and health variables: a retrospective study. *Med Sci Sports Exerc*, 2001 Feb;33(2):183-8
74. Juhn MS, Tarnopolsky M. Potential side effects of oral creatine supplementation: a critical review. *Clin J Sport Med*, 1998 Oct;8(4):298-304
75. Dietary Supplement Information Bureau. [www.intramedicine.com](http://www.intramedicine.com)
76. Tarnopolsky M, Martin J. Creatine monohydrate increases strength in patients with neuromuscular disease. *Neurology*. Mar 1999;52(4):854-7
77. Vorgerd, M., Grehl, T., Jager, M., et al. Creatine therapy in myophosphorylase deficiency (McArdle disease): a placebo-controlled crossover trial. *Arch Neurol*. Jul 2000;57(7):956-63
78. Gordon A, Hultman E, Kaijeser L et al. Creatine supplementation in chronic heart failure increases skeletal muscle creatine phosphate and muscle performance. *Cardiovasc Res*. Sep1995;30(3):413-8
79. Andrews R, Greenhaff P, Curtis S et al. The effect of dietary creatine supplementation on skeletal muscle metabolism in congestive heart failure. *Eur Heart J*. Apr1998;19(4):617-22
80. Healthnotes, Inc 2001. [www.healthnotes.com](http://www.healthnotes.com)
81. Walter MC, Lochmuller H, Reilich P et al. Creatine monohydrate in muscular dystrophies: A double-blind, placebo-controlled clinical study. *Neurology* 2000;54:1848-50
82. Felber S, Skladal D, Wyss M et al. Oral creatine supplementation in Duchenne muscular dystrophy: a clinical and 31P magnetic resonance spectroscopy study. *Neurol Res* 2000;22:145-50



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## Dehydroepiandrosterone (DHEA)

### General Features

DHEA is an intermediate steroid hormone produced mostly by the adrenal glands. All steroid hormones are derived from cholesterol. In the synthesis of adrenal androgen hormones cholesterol is converted to pregnenolone and then to DHEA. From DHEA the adrenal glands can synthesize androstenedione, which is further converted to testosterone. In fat tissue androstenedione can be converted to estrone hormone by the aromatase enzyme, which is also known as estrogen synthase enzyme. Thus, DHEA supplementation can affect the increased production of androstenedione as well as testosterone and estrogen.

DHEA is the most abundant hormone made by the adrenal glands. Some DHEA is secreted by the adrenal glands and circulates in the bloodstream, where it is picked up by other tissues (i.e. adipose, testis, ovaries) and further converted into other androgens or estrogens.

The serum concentration of DHEA (really DHEA - sulfate), is used as a measure of adrenal androgen production, when monitoring various conditions.<sup>1</sup>

DHEA supplements can be made in the laboratory from diosgenin a steroid compound found in wild yams. However, the body is unable to convert diosgenin into DHEA or any other hormone. Thus, supplementing with wild yam as a means to affect hormone levels is unsubstantiated.<sup>2</sup>

In humans, DHEA blood levels peak in early adulthood and then starts a lifelong descent. By the age of 70 DHEA levels have declined by up to 75 percent compared with young adult levels. By age 90, we make 90 percent less DHEA than a young adult.<sup>3,4</sup>

These findings have led some researchers to investigate whether returning DHEA levels to those of a young adult (through supplementation) can serve as an anti-aging, and degenerative disease prevention strategy. Preliminary reports in this regard are conflicting. Some evidence suggests that DHEA supplementation (25-200 mg per day) can reverse some parameters of aging and improve wellbeing. Other reports correlate higher blood DHEA levels (and supplementation in some cases) with increased risk of prostate cancer, postmenopausal breast cancer, and ovarian cancer.<sup>5-13</sup>

As a result many health authorities are cautious about recommending DHEA supplementation as an anti-aging intervention. Individuals with a history or family history of breast, ovarian or prostate cancer should not supplement with DHEA indiscriminately until further studies are completed.<sup>14</sup>

The average male produces 31 mg of DHEA per day, while women make about 19 mg.<sup>15</sup>

### Supplementation Studies and Clinical Applications

#### 1. Systemic Lupus Erythematosus (SLE)

In a Stanford Medical Center study, DHEA supplementation (200 mg per day) decreased the SLE Disease Activity Index by nearly two points, while the placebo group increased by almost a full point. DHEA patients had significantly fewer flare ups and their required dosage of corticosteroid drug used to control symptoms decreased by 35 percent, whereas the placebo group increased their dose of corticosteroids by forty percent. This was a three-month study only. Long-term benefits are yet unknown and the major side effects in this study was mild to severe acne in women in the DHEA group.<sup>16,17</sup>

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