Dynamic Chiropractic



Vitamin D Alert

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Generally, osteopenia seems to be increasing in my film stack. Nearly half of the women over the age of 50 seem to have some osteopenia, and it seems to be increasing even in younger women. I've even seen several cases of middle-aged men with osteopenia. This article serves as a brief alert regarding vitamin D deficiency and its link to many disease processes, some of which many of us do not recognize. After only a brief review of recent literature, I was surprised to discover that this is not a common problem. Partly due to poor nutrition and partly to the lack of sun exposure, vitamin D insufficiency is increasingly associated with a multitude of other disease processes I am guilty of not recognizing, too.

Conditions Associated With, Caused by and/or Contributing to Vitamin D Deficiency (partial list)

- Chronic pain
- Multiple sclerosis
- Type I diabetes
- Systemic lupus erathematosus
- Malabsorption syndromes
- Cancers: prostate, colon, breast
- Long-term: use of prednisone, anti-convulsant meds
- Use of medications with photosensitivity precautions
- Lack of sun exposure and sunscreen use
- Liver and kidney disease
- Congestive heart failure
- Peripheral arterial disease
- Rickets
- Osteoporosis
- Osteomalacia

Amazingly, more than 50 percent of women already receiving treatment for osteoporosis are

vitamin D deficient.¹ This fact alone is so astounding that I think we should re-evaluate how we assess this group of patients. Other patients at an extremely high risk for deficiency include those on long-term prednisone, certain anticonvulsant medications and any medication that has

photosensitivity precautions. Additionally, individuals with medical conditions that require sun avoidance, such as systemic lupus erythematosus or sickle cell disease, are at an extremely high risk. People with significant kidney or liver disease, congestive heart failure, or peripheral arterial disease are at risk for unrecognized, clinically significant deficiency. In fact, any patient whose medical condition significantly limits their sun exposure or fat absorption is at risk for significant deficiency.

Vitamin D insufficiency is common among people over 50 years of age, with reported prevalence between 25 percent and 50 percent of the population. Of course, with institutionalized elderly populations, the prevalence is much higher. Clinically significant vitamin D deficiency is common

worldwide. Numerous international medical authorities have emphasized its seriousness.²⁻⁷ Yet clinicians are still not evaluating patients at risk for vitamin D deficiencies. This is not just a problem in developing countries; it is prevalent in the U.S. and Europe. It often goes unrecognized.

This is unfortunate because vitamin D plays an important role in bone development and muscle function. It also facilitates absorption of calcium and phosphate from the gut and kidney, suppresses parathyroid hormone (PTH), and acts on osteoblasts to stimulate bone formation. We are well aware of the role of vitamin D deficiency as a risk factor for osteoporosis, osteomalacia and rickets, but new discoveries are finding its functions include a role in muscle strength and the prevention of autoimmune diseases and some cancers. Epidemiologic and *in vitro* data have generated provocative hypotheses regarding vitamin D deficiency and the increased risk of 13 types of cancer, including breast, colon and prostate. Along with cancer, the disease is being linked to type I diabetes, multiple sclerosis, hypertension, rheumatoid arthritis and chronic, unexplained

musculoskeletal pain.^{2,3,7,8}

An interesting statistic I was not aware of until recently is that chronic, nonspecific pain is estimated to occur in up to 20 percent of adults. Over the past 30 years, European studies have documented this type of pain syndrome in patients with severe hypovitaminosis D. A study reported by G.A. Poltnikoff and J.M. Quigley in the 2003 *Mayo Clinic Proceedings* evaluated patients with chronic pain, unresponsive to standard pharmaceutical agents and who did not have a diagnosis for their pain or any significant medical conditions which would interfere with production and

absorption of vitamin D.⁷ The patients were between the ages of 10 and 65; none was housebound or elderly. Ninety percent of the patients in the study were found to have deficient levels of vitamin D.⁹

Do you think we as chiropractors should assess blood levels of vitamin D? It should be noted that none of these patients had been diagnosed with vitamin D deficiency before the study. Vitamin D deficiency is not asymptomatic and results indicate that a clinical syndrome consisting of persistent musculoskeletal pain, later progressing to bone pain, is associated with the level of deficiency in these patients. In this study, a failure in development of peak bone mass in the younger patients and an excessive loss of skeletal integrity in older patients was documented.

The assessment of vitamin D status requires measurement of circulating 25-hydroxyvitamin D. The present opinion is that the optimal 25-hydroxyvitamin D concentration needs to be greater than 30 ng/mL. Several facts suggest that the current norms are too low. It has been documented that parathyroid hormone (PTH) is secreted by the body in response to insufficient calcium absorption. Elevated levels indicate long-standing inadequacy and a high bone-remodeling rate. PTH levels minimize with 25-OHD serum levels of at least 20 ng/mL. The fractional oral calcium absorption is

optimized at approximately 30 ng/mL.^{10,11} Several experts who attended the November 2004 NIH Vitamin D and Cancer Conference asserted that the physiologic lower end of normal should be set

at 32 ng/mL based on *in vitro* and randomized controlled trial data.¹⁰⁻¹⁴ We should keep this range in mind when assessing patients for vitamin D deficiencies.

Vitamin D is found in numerous dietary sources, including fish, eggs, fortified milk and cod liver oil. The sun is also a significant contributor to our daily production of vitamin D, and as little as 10 minutes of daily exposure is thought to be enough to prevent deficiencies in young healthy individuals. However, it is rare to find true recommendations for patients who are compromised by health issues or degenerative diseases, or for elderly patients whose production of vitamin D_3 decreases with age.

The term "vitamin D" refers to several different forms. Two forms important in humans are: ergocalciferol (vitamin D_2) and cholecalciferol (vitamin D_3). Vitamin D_2 is synthesized by plants. Vitamin D_3 is synthesized by humans in the skin when it is exposed to ultraviolet B (UVB) rays from sunlight or diet. Vitamin D_3 is preferred to D_2 as a supplement because the half-life is longer. Also, D_3 is more potent and its bind to the vitamin D binding protein is stronger and ingestion does not result in unique biologically active metabolites.



Adequate levels of vitamin D, as recommended by the U.S. Institute of Medicine of the National Academy of Sciences, are presently at 200 IU daily for all individuals (male, female, or pregnant/lactating women) under the age of 50, 400 IU for all individuals between the ages 50 and 70, and 600 IU for those who are over the age of 70. The daily "upper limit" for vitamin D is 1,000 IU for infants up to 12 months of age and 2,000 IU for children, adults, and pregnant and lactating

women. The most effective way of ensuring adequate vitamin D levels is to take a daily multivitamin, which usually contains 400 IU. However, experts now believe that the current norms are too low. More study is needed to identify both prevalence and adequate daily intake. In addition, sun exposure guidelines that take into account geographic latitude, season, age, and factors such as race and clothing are needed.

Since I am not a biochemist or clinical nutritionist, I invite you to review the recent literature and assess this topic more with others who are more qualified. I recommend visiting www.vitamindcouncil.com and www.sunarc.org/index.htm. You might even consider assessing your own levels of serum 25-hydroxyvitamin D. I encourage you to inform me if you have information on this topic that you would like to share with our colleagues.

References

- Holick MF, Siris ES, Binkley N, et al. Prevalence of vitamin D inadequacy among postmenopausal North Amer-ican women receiving osteoporosis therapy. J Clin Endocrinol Metab, 2005; 90(6):3215-24.
- 2. Eugster EA, Sane KS, Brown DM. Minnesota rickets. Need for a policy change to support vitamin D supplementation. *Minn Med*, 1996;79(8):29-32.
- 3. Compston JE. Vitamin D deficiency: time for action. *BMJ*, 1998;317(7171):1466-7.
- 4. Utiger RD. The need for more vitamin D. N Engl J Med, 1998;338(12):828-9.
- 5. Stokstad E. The vitamin D deficit. Science, 2003;302:1886-8.
- 6. Meier C. Scientists probe role of vitamin D: deficiency a significant problem, experts say. *JAMA*, 2004;292(12):1416-8.
- 7. Holick MF. Vitamin D deficiency: what a pain it is. *Mayo Clin Proc*, 2003;78(12): 1457-9.
- 8. Nesby-O'Dell S, Scanlon KS, Cogswell ME, et al. Hypovitaminosis D prevalence and determinants among African American and white women of reproductive age: Third National Health and Nutrition Examination Survey, 1988-1994. *Am J Clin Nutr*, 2002;76(1):187-92.
- 9. Plotnikoff GA, Quigley JM. Prevalence of severe hypovitaminosis D in pa-tients with persistent, nonspecific musculoskeletal pain. *Mayo Clin Proc*, 2003;78(12):1463-70.
- 10. Malabanan A, Veronikis IE, Holick MF. Redefining vitamin D insufficiency. *Lancet*, 1998;351(9105):805-6.
- 11. Heaney RP. The vitamin D requirement in health and disease. *J Steroid Biochem Mol Biol*, 2005; July 15.
- 12. Vieth R. Vitamin D supplementation, 25-hydroxyvitamin D concentrations and safety. *Am J Clin Nutr*, 1999;69(5): 842-56.
- 13. Heaney RP, Davies KM, Chen TC, et al. Human serum 25-hydroxycholecalciferol response to extended oral dosing with cholecalciferol. *Am J Clin Nutr*, 2003;77(1):204-10.
- 14. Armas LA, Hollis BW, Heaney RP. Vitamin D_2 is much less effective than vitamin D_3 in humans. *J Clin Endocrinol Metab*, 2004;89(11):5387-91.

Resources

- 1. Looker AC, Dawson-Hughes B, Calvo MS, et al. Serum 25-hydroxyvitamin D status of adolescents and adults in two seasonal subpopulations from NHANES III. *Bone*, 2002;30(5):771-7.
- 2. Tangpricha V, Pearce EN, Chen TC, Holick MF. Vitamin D insufficiency among free-living healthy young adults. *Am J Med*, 2002;112(8):659-62.
- 3. Sullivan SS, Rosen CJ, Halteman WA, et al. Adolescent girls in Maine are at risk for vitamin D insufficiency. *J Am Diet Assoc*, 2005;105(6):971-4.
- 4. Weiler H, Fitzpatrick-Wong S, Veitch R, et al. Vitamin D deficiency and whole-body and femur bone mass relative to weight in healthy newborns. *CMAJ*, 2005;172(6):757-61.
- 5. Haney EM, Stadler D, Bliziotes MM. Vitamin D insufficiency in internal medicine residents. *Calcif Tissue Int*, 2005;76(1):11-6.

- 6. Moore CE, Murphy MM, Holick MF. Vitamin D intakes by children and adults in the United States differ among ethnic groups. *J Nutr*, 2005; 135(10):2478-85.
- 7. Thomas MK, Lloyd-Jones DM, Thadhani RI, et al. Hypovitaminosis D in medical inpatients. *N Engl J Med*, 1998;338(12):777-83.
- 8. Mylott BM, Kump T, Bolton ML, Greenbaum LA. Rickets in the dairy state. *WMJ*, 2004;103(5):84-7.
- 9. Dawson-Hughes B, Heaney RP, Holick MF, et al. Estimates of optimal vitamin D status. *Osteoporos Int*, 2005;16(7):713-6.
- Singh SK, Manjure S, Stott P, et al. Does routine blood bone biochemistry predict vitamin D insufficiency in elderly patients with low-velocity fractures? J Orthop Surg (Hong Kong), 2004;12(1):31-4.
- 11. Dusso AS, Brown AJ, Slatopolsky E. Vitamin D. *Am J Physiol Renal Physiol*, 2005;289(1):F8-28.
- 12. Lee S, Clark SA, Gill RK, Christakos S. 1,25 dihydroxyvitamin D_3 and pancreatic β -cell function: vitamin D receptors, gene expression, and insulin secretion. *Endocrinology*, 1994;134(4):1602-10.
- 13. Weishaar RE, Simpson RU. The involve-ment of the endocrine system in regulating cardiovascular function: emphasis on vitamin D_3 . *Endocr Rev*, 1989;10(3):1-15.
- 14. Li YC, Qiao G, Uskokovic M, et al. Vitamin D: a negative endocrine regulator of the reninangiotensin system and blood pressure. *J Steroid Biocehm Mol Biol*, 2004;89-90(1-5):387-92.
- 15. Sato Y, Iwamoto J, Kanoko T, Satoh K. Low-dose vitamin D prevents muscular atrophy and reduces falls and hip fractures in women after stroke: a randomized controlled trial. *Cerebrovascular Dis*, 2005;20(3):187-92.
- 16. Ustianowski A, Shaffer R, Collins S, et al. Prevalence and association of vitamin D deficiency in foreign-born persons with tuberculosis in London. *J Infect*, 2005;50(5):432-7.
- Bringhurst FR, Demay MB, Kronenberg HM. Mineral metabolism. In: Williams Textbook of Endocrinology. Larson PR, Kronenberg HM, Melmed S, Polonsky KS, Eds. Amsterdam: Elsevier; 2003, pp. 1317-20.
- 18. Venning G. Recent developments in vitamin D deficiency and muscle weakness among elderly people. *BMJ*, 2005;330(7490):524-6.
- 19. Visser M, Deeg DJ, Lips P. Low vitamin D and high parathyroid hormone levels as determinants of loss of muscle strength and muscle mass (sarcopenia): the Longitudinal Aging Study Amsterdam. *J Clin Endocrinol Metab*, 2003;88(12):5766-72.
- 20. Trivedi DP, Doll R, Khaw KT. Effect of four monthly oral vitamin D_3 (cholecalciferol) supplementation on fractures and mortality in men and women living in the community: randomized double-blind controlled trial. *BMJ*, 2003;326(7387): 469-74.
- 21. Dhesi JK, Bearne LM, Moniz C, et al. Neuromuscular and psychomotor function in elderly subjects who fall and the relationship with vitamin D status. *J Bone Miner Res*, 2002;17(5):891-7.
- 22. Bischoff-Ferrari HA, Dietrich T, Orav EJ, et al. Higher 25-hydroxyvitamin D concentrations are associated with better lower-extremity function in both active and inactive persons aged > or = 60 y. *Am J Clin Nutr*, 2004;80(3):752-8.
- 23. Peterlik M, Cross HS. Vitamin D and calcium deficits predispose for multiple chronic diseases. *Eur J Clin Invest*, 2005;35(5):290-304.
- 24. Holick MF, Siris ES, Binkley N, et al. Prevalence of vitamin D inadequacy among postmenopausal North American women receiving osteoporosis therapy. *J Clin Endocrinol Metab*, 2005;90(6):l3215-24.

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