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Calcium and vitamin D use among adults in periodontal-disease maintenance programs

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Abstract

Objectives—Determine the level of calcium and vitamin D oral supplementation in patients in periodontal disease maintenance programs.

Design—Convenience survey.

Setting—St. Louis Metropolitan region.

Subjects and Methods—Patients (n=228) in two university-based, periodontal-disease maintenance programs.

Main Outcome Measures—Reported amounts of oral calcium and vitamin D supplementation were tested for differences based on gender and race.

Results—The last published recommended daily intakes from the United States (U.S.) Food and Nutrition Board (FNB) for adults >50 years of age are 1200 mg calcium and 400 IU vitamin D (or

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600 IU if over 70). The mean age of the 228 patients (125 females and 103 males) was 63.6 ± 11.0 years (standard deviation). Of the 228 patients surveyed: (1) 204(89%) were >50 years of age, and of these, only 15(7%) met the U.S. FNB's recommended intakes of calcium and vitamin D from supplementation, (2) 138(66%) reported that they took no oral supplementation, with significantly more males (n=82) than females (n=56) not taking supplementation (p=0.03), (3) 88(39%) took calcium supplementation, with females (947±511mg/day) taking significantly (p<0.001) more than males (632±907mg/day), and (4) 66(29%) took vitamin D supplementation, with females(420±227 IU/day) taking approximately the same amount as males (443±317 IU/day, p>0.05). The amounts of oral supplementation did not vary with race (p>0.05).

Conclusion—The use of calcium and vitamin D supplementation has been promoted for years; yet the numbers of adults taking supplements remains low and the level of supplementation varies greatly. Knowledge of the benefits of supplementation needs to be better disseminated and research needs to be conducted to determine optimal levels of calcium and vitamin D supplementation.

Keywords

Calcium/therapeutic use; diet; nutrition; periodontal diseases/prevention and control; vitamin D

INTRODUCTION

Oral bone mass increases throughout infancy, childhood, and adolescence to reach a genetically determined peak bone mass in early adulthood. Thereafter, the oral bone, just as total skeletal bone, loses bone mass. The rate of increase and loss is dependent upon heredity and the availability of calcium and vitamin D. If calcium intakes are inadequate, skeletal calcium is resorbed to maintain the body's calcium homeostasis, which is essential for life-sustaining processes–such as blood clotting, muscle contraction, and nerve excitability. The resorption of the body's calcium is mediated by parathyroid hormone (PTH), which increases osteoclastic activity in bone to release stored calcium to the circulation and increases the synthesis of 1,25-dihydroxyvitamin D [1,25(OH)₂D] (the biologically active form of vitamin D), which in turn increases the absorption of calcium from the intestine. The best way to estimate vitamin D intakes is to measure the serum levels of 25-hydroxyvitamin D [25(OH)D].¹ When serum levels of 25(OH)D are low, there is decreased calcium absorption, increased PTH secretion, and increased osteoclastic activity.

In a September 2002 review of 180 articles on bone status and calcium intake, it was reported that 68 of 70 (97%) randomized clinical trials found that individuals who received calcium supplementation had greater gains in bone during growth, less loss of bone with age, and/or reduced fracture risk, relative to unsupplemented individuals, and that 85 of 110 (77%) observational studies found positive effects for calcium supplementation.² It is also well established that vitamin D is essential for bone growth and that it has its greatest effect when combined with calcium supplementation.³⁻⁵ *Dietary Reference Intakes* [the last published recommendations of the U.S. Institute of Medicine's Food and Nutrition Board (FNB)] suggests a total intake from diet and oral supplementation of 1200 mg/day calcium and 400 IU (10.0 μ g)/day vitamin D for ages 51-70 and for those >70 years of age, 1200 mg calcium/day and 600 IU vitamin D (15.0 μ g)/day.⁶ In the U.S., the median calcium intake for men 50 to 70 years of age is 708 mg/day and for women 571 mg/day and 702 and 517 mg/day respectively for those >70 years of age.⁷ Therefore, on average, adult men and women need to consume an additional 500-600 mg/day of calcium through food or supplementation, just to meet health needs.

There is also growing concern that recent studies have identified high prevalence's of vitamin D insufficiency in otherwise healthy adults and that current FNB recommendations of vitamin D intake are inadequate. A number of investigators convincingly argue that serum 25(OH)D levels < 70-80 nmol/liter are deficient.⁸ Another source for information, the National Osteoporosis Foundation (NOF) recommends 1,000 mg calcium and 400-800 IU Vitamin D daily for adults <50 years of age, and for those >50 years of age, 1,200 mg calcium and 800-1,000 IU Vitamin D daily.⁹ A periodontal-disease, vitamin-D study of the Third National Health and Nutrition Examination Survey (NHANES III) data found that the majority of subjects had serum 25(OH)D levels far below 80 nmol/liter.¹⁰ As pointed out in recent studies, the older recommendations of 400 or 600 IU vitamin D/day were intended to maintain health and are inadequate for correcting vitamin D insufficiencies.¹¹

It is reasonable to think that calcium and vitamin D affect oral bone in the same way that they affect other bones of the body, and various studies, including analyses of data from NHANES III, have found associations between periodontal health and intakes of calcium and vitamin D.^{10, 12}Researchers have found that (1)optimal intakes of calcium and vitamin D slow bone resorption and speed bone formation, thereby reducing bone loss¹³, (2)vitamin D stimulates the expression of antimicrobial peptides (α - and β -defensins and cathelicidins),^{14, 15} and (3)vitamin D has an immunosuppressive effect on periodontal disease.¹⁰ As the levels of calcium and vitamin D oral supplementation for patients with periodontal disease are unknown, the purpose of this study was to survey adult patients in two university-based, periodontal-disease maintenance programs to determine their levels of calcium and vitamin D oral supplementation.

METHODS

Study Population

We conducted our survey at Southern Illinois University School of Dental Medicine (SIU-SDM) in Alton, Illinois and the Saint Louis University Center for Advanced Dental Education (SLU-CADE) in St. Louis, Missouri. Study subjects were recruited from periodontal-disease recall programs based at these two institutions, whose patient populations are from the Greater St. Louis region, an area comprising 12 counties in Eastern Missouri and Southern Illinois. According to the latest U.S. census, this area has a population of approximately 2.3 million, which makes it the 18th largest market area in the United States. There are approximately 920 patients in the periodontal-disease maintenance program at Southern Illinois University and 200 at Saint Louis University.

Patient Questionnaire

We obtained Institutional Review Board (IRB) approvals for our protocol and our use of a brief patient questionnaire at SIU-SDM, SLU-CADE, and Washington University in St. Louis (principal investigator's home institution). Signs were posted in the periodontal maintenance program dental operatories at SIU-SDM and SLU-CADE, with the sign's informing patients of a potential research study involving calcium and vitamin D. Interested patients were invited to complete our brief questionnaire, which requested information on age, gender, number of years postmenopausal, use of vitamin D and/or calcium supplements (including multivitamins and antacids), and use of bone-active prescription drugs (such as, risedronate,^{*} alendronate,[†] raloxifene,[‡] or estrogen).

^{*}Actonel, Procter & Gamble Pharmaceuticals, Inc.

[†]Fosamax, Merck Co., West Point, VA

[‡]Evista [raloxifene hydrochloride, a selective estrogen receptor modulator (SERM), Eli Lilly and Company, Indianapolis, IN)

Questionnaires were checked for completeness, and if additional information on amounts of calcium and vitamin D was needed after the patients had gone home, they were contacted by telephone.

Data Analysis

Data are reported as means \pm standard deviation. For categorical variables, frequencies were calculated for patients who were or were not taking calcium/vitamin D and/or bone-active prescription drugs. Distributions of age, mg of calcium taken per day, and international units $(IUs-40 IUs = 1 \mu g)$ of vitamin D taken per day were tested for normality with the Shapiro-Wilk W test, and equality of variances were tested with O'Brien, Brown-Forsythe, Levene, and Bartlett tests. Differences in responses based on gender, menopausal status, and race were tested for statistical significance with the chi-square test, Fisher's exact test, analysis of variance (ANOVA) and Student's t test. If data were non-normally distributed and/or variances were not equal, the Welch ANOVA and/or Wilcoxon tests were used. For those who took neither calcium nor vitamin D oral supplementation, the binomial test was used to determine if there were more males than females. This test is used when there are only two possible outcomes, with fixed rates of occurrence-in this case 0.5 for males and 0.5 for females.¹⁶ The chi-square, goodness-of-fit test was used to test for differences in racial proportions between respondents and the St. Louis region. If for the chi-square test, 20% of the cells had expected counts of less than 5, exact p values were calculated by means of data permutation. If default time limits were exceeded for calculations of exact p values, Monte Carlo estimates were used. Monte Carlo estimates and exact calculations of p values were performed with StatXact Statistical Software for Exact Nonparametric Inference (Version 6 with Cytel Studio[™], Cytel Software Corporation, Cambridge, MA). All other statistical testing was performed with JMP Statistical Software (SAS Institute, Inc., Cary, NC).

RESULTS

From May 2003 through September 2004, 228 questionnaires were completed and returned. The racial make-up of the pooled sample was not significantly different (p = 0.15) from that for the St. Louis region. The mean age of the 228 patients surveyed was 63.6 ± 11.0 years, with 204 (89%) being over the age of 50 years. Males were significantly older (64.2 ± 9.5 years) than females (58.8 ± 11.9 years, Table 1). Postmenopausal women were significantly older than pre-menopausal women (p < 0.0001), and African Americans were significantly younger than Whites (p < 0.05). Eighty-eight patients reported taking oral calcium supplementation. Females took significantly more calcium than did men (p<0.001). The percentage of African Americans, who we surveyed, who took calcium (13.6%, n = 12, Table 1) was higher than the percentage of African Americans living in the St. Louis region (9.6%). Conversely, the percentage of Whites, who we surveyed, who took calcium (84.1%, n = 74, Table 1) was lower than the percentage of Whites living in the St. Louis region (87.9%).

Sixty-six patients reported taking oral vitamin D supplementation. Unlike what was observed with calcium, there was no significant difference between men and women in the amounts of vitamin D supplementation taken. There was also no variation in vitamin D oral supplementation by race when compared to the makeup of the St. Louis region (p>0.05) (Table 1).

Among all subjects studied, 64 of the 228 (28%) took both oral calcium and vitamin D supplementation, and 138 of the 228 (61%) took no oral supplementation. Insufficient calcium and vitamin D intake, as assessed by supplementation amounts, was seen in 93% of the study population (Table 2).

Thirty-three patients reported use of bone-active prescription drugs used for osteoporosis, specifically, risedronate (n = 5), alendronate (n = 13), estrogen (n = 4), raloxifene (n = 10), 1 unknown drug (unable to follow up with patient). Ninety-one percent of these patients were females, 97% were postmenopausal; and 91% were White (Table 1). Fifteen (45%) of these 33 patients indicated use of calcium supplements (1252 ± 456 mg/day); and 11 (33%) of the 33 indicated use of vitamin D supplements (473 ± 185 IU/day).

DISCUSSION

With regard to periodontal disease, the many benefits of calcium and/or vitamin D include strengthening the bone¹³, stimulating an antimicrobial effect,^{14, 15} and providing an immunosuppressive effect.¹⁰

The self-reported calcium and vitamin D intake levels would be insufficient by themselves to prevent deficiencies in our older adult population. Periodontal maintenance patients receive no advice concerning calcium and vitamin D intake at either institution. Accurate estimates of total intakes of calcium and vitamin D for our study subjects would have required performing dietary analyses, assessing sunlight exposures, or (in the case for vitamin D) assessing serum levels of 25(OH)D-none of which were done for our study. Nevertheless, of the patients we surveyed, 204 (89%) of 228 were over the age of 50, and based on the recommendations given in *U. S. Dietary Reference Intakes*,⁶ of these, only 15 (7%) obtained adequate intakes of calcium and vitamin D from oral supplementation, and only 6 (.03%) met the higher recommendations of the National Osteoporosis Foundation.⁹ Other studies of St. Louis women also indicate inadequate intakes (food and oral supplementation) of calcium and vitamin D.^{17, 18}

The median calcium intakes (food plus oral supplementation) in the U.S. for men 50 to 70 years of age is 708 mg/day and for women 571 mg/day. Over the age of 70, the respective median intakes are 702 and 517 mg/day.⁷ *U. S. Dietary Reference Intakes* contains no table for vitamin D intakes such as it does for calcium; however, there is growing concern among researchers that recent studies have identified an alarming prevalence of vitamin D insufficiency in otherwise healthy adults and a re-emergence of vitamin D-deficiency-induced rickets.¹⁹ Many researchers now believe that for adults, daily supplementation with vitamin D at levels of, at least, 800 to1,000 IU (20 and 25 μ g)/day would be safe and beneficial to health, and that the U. S. Food and Nutrition Board (FNB) of the Institute of Medicine needs to raise the recommended levels of intake for vitamin D.⁸

Dental Studies of the Effects of Calcium and/vitamin D on Alveolar Bone, Periodontal Health, and Tooth Retention

It has been found that increased periodontal attachment loss is related to decreased intakes of both calcium and vitamin D.^{10, 12} The investigators suggested that inadequate calcium intake increased risk of periodontal disease and could be related to decreased alveolar bone density. Inadequate intake of vitamin D could increase levels of periodontal disease either through an effect on bone mineral density or a suppression of vitamin D (particularly the suppression of cytokine production) and vitamin D's inducing the production of antimicrobial cathelicidin have been noted by various investigators.^{15, 20-24} There have been a number of studies that indicate that calcium/vitamin D intakes affect alveolar bone, periodontal health, and tooth retention, and even though there were limitations associated with these studies, there is strong evidence that calcium and vitamin D intake are positively associated with periodontal/alveolar bone health.²⁵

Anti-resorptive agents—Fourteen percent (33/228) of the patients took bone-active prescription drugs, with the majority being female. Some patients in our study took risedronate or alendronate, which are bisphosphonates, which may improve periodontal health but also may cause osteonecrosis and painful refractory bone exposures of the jaws.²⁶⁻³¹ The incidence of osteonecrosis of the jaw in patients taking oral bisphosphonate therapy for osteoporosis is low, with a reported risk of less than 1 in 100,000 patients.³² Some of the patients in our study took estrogen, which may benefit periodontal health.³³⁻³⁶ Other patients in our study took raloxifene [a selective estrogen receptor modulator, SERM]. No study has been published on the effect of this anti-resorptive agent on periodontal health.

Study limitations—The results of our study were drawn from self-reported questionnaires. Although the data were unconfirmed (except by telephone in a few cases), we gained knowledge about the use of oral calcium and vitamin D supplementation in a group of adults with known periodontal disease.

Conclusion

Calcium and vitamin D supplements are reasonably priced and widely available; yet many older adults in the U. S. take neither calcium nor vitamin D supplementation, and of those who do, intake levels are often inadequate.

Randomized clinical trials and observational studies have consistently demonstrated positive effects of calcium and/or vitamin D oral supplementation on postcranial (below the head) and alveolar bone.²⁵ There are few health concerns associated with calcium and vitamin D use, but perhaps more important, there is considerable potential for health benefits,³⁷ including periodontal health. Dentists and dental health professionals should advise all of their adult patients to follow the National Osteoporosis Foundation's daily intake recommendations of 1,000 mg calcium and 400-800 IU vitamin D for adults <50 years of age, and 1200 mg calcium and 800-1,000 IU vitamin D for adults >50 years of age. Controlled studies should be conducted to establish the extent to which adequate intakes of calcium and vitamin D can stabilize/improve periodontal health.

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REFERENCES

- Food and Nutrition Board. Dietary Reference Intakes (DRI) for Calcium, Phosphorus, Magnesium, Vitamin D, and Fluoride.: Food and Nutrition Board (FNB), Institute of Medicine. National Academy Press; Washington, DC: 1997. p. 253
- 2. Heaney RP. The importance of calcium intake for lifelong skeletal health. Calcif Tissue Int. 2002a; 70(2):70–3. [PubMed: 11870410]
- Lips P. Vitamin D deficiency and secondary hyperparathyroidism in the elderly: consequences for bone loss and fractures and therapeutic implications. Endocr Rev. 2001; 22(4):477–501. [PubMed: 11493580]
- Papadimitropoulos E, Wells G, Shea B, et al. Meta-analyses of therapies for postmenopausal osteoporosis. VIII: Meta-analysis of the efficacy of vitamin D treatment in preventing osteoporosis in postmenopausal women. Endocr Rev. 2002; 23(4):560–9. [PubMed: 12202471]

- Shea B, Wells G, Cranney A, et al. Meta-analyses of therapies for postmenopausal osteoporosis. VII. Meta-analysis of calcium supplementation for the prevention of postmenopausal osteoporosis. Endocr Rev. 2002; 23(4):552–9. [PubMed: 12202470]
- Food and Nutrition Board. Dietary Reference Intakes (DRI) for Calcium, Phosphorus, Magnesium, Vitamin D, and Fluoride.: Food and Nutrition Board (FNB), Institute of Medicine. National Academy Press; Washington, DC: 1997. p. 115-7.p. 273-5.
- Food and Nutrition Board. Dietary Reference Intakes (DRI) for Calcium, Phosphorus, Magnesium, Vitamin D, and Fluoride.: Food and Nutrition Board (FNB), Institute of Medicine. National Academy Press; Washington, DC: 1997. p. 388-9.
- Dawson-Hughes B, Heaney RP, Holick MF, et al. Estimates of optimal vitamin D status. Osteoporosis International. 2005; 16(7):713–6. [PubMed: 15776217]
- NOF. National Osteoporosis Foundation's Updated Recommendations for Calcium and Vitamin D intake. National Osteoporosis Foundation; 2007. http://www.nof.org/prevention/ calcium_and_VitaminD.htm
- Dietrich T, Joshipura KJ, Dawson-Hughes B, Bischoff-Ferrari HA. Association between serum concentrations of 25-hydroxyvitamin D3 and periodontal disease in the US population. Am J Clin Nutr. 2004; 80(1):108–13. [PubMed: 15213036]
- Heaney RP. Barriers to optimizing vitamin D3 intake for the elderly. Journal of Nutrition. 2006; 136(4):1123–5. [PubMed: 16549492]
- Nishida M, Grossi SG, Dunford RG, et al. Calcium and the risk for periodontal disease. J Periodontol. 2000; 71(7):1057–66. [PubMed: 10960010]
- 13. Frost, HM. Bone Remodeling and Its Relationship to Metabolic Bone Disease. Charles C. Thomas; Springfield, Ill: 1973. Remodeling as a determinant of envelope physiology; p. 28-53.
- Cannell JJ, Zasloff M, Garland CF, Scragg R, Giovannucci E. On the epidemiology of influenza. Virol J. 2008; 5:29. [PubMed: 18298852]
- 15. Gombart AF, Borregaard N, Koeffler HP. Human cathelicidin antimicrobial peptide (CAMP) gene is a direct target of the vitamin D receptor and is strongly up-regulated in myeloid cells by 1,25dihydroxyvitamin D3. Faseb J. 2005; 19(9):1067–77. [PubMed: 15985530]
- Levin, RI.; Rubin, DS. Applied Elementary Statistics. Prentice-Hall, Inc.; Englewood Cliffs, New Jersey: 1980.
- Hildebolt CF, Pilgram TK, Dotson M, et al. Attachment loss with postmenopausal age and smoking. J Periodontal Res. 1997; 32(7):619–25. [PubMed: 9401935]
- Villareal DT, Civitelli R, Chines A, Avioli LV. Subclinical vitamin D deficiency in postmenopausal women with low vertebral bone mass. J Clin Endocrinol Metab. 1991; 72(3):628– 34. [PubMed: 1997517]
- Holick MF. Resurrection of vitamin D deficiency and rickets. Journal of Clinical Investigation. 2006; 116(8):2062–72. [PubMed: 16886050]
- 20. Murakami M, Ohtake T, Dorschner RA, Gallo RL. Cathelicidin antimicrobial peptides are expressed in salivary glands and saliva. J Dent Res. 2002; 81(12):845–50. [PubMed: 12454100]
- Wang TT, Nestel FP, Bourdeau V, et al. Cutting edge: 1,25-dihydroxyvitamin D3 is a direct inducer of antimicrobial peptide gene expression. J Immunol. 2004; 173(5):2909–12. [PubMed: 15322146]
- 22. Cannell JJ, Vieth R, Umhau JC, et al. Epidemic influenza and vitamin D. Epidemiology & Infection. 2006; 134(6):1129–40. [PubMed: 16959053]
- Liu PT, Stenger S, Li H, et al. Toll-like receptor triggering of a vitamin D-mediated human antimicrobial response.[see comment]. Science. 2006; 311(5768):1770–3. [PubMed: 16497887]
- 24. Zittermann A. Vitamin D in preventive medicine: are we ignoring the evidence? Br J Nutr. 2003; 89(5):552–72. [PubMed: 12720576]
- Hildebolt C. Effect of Vitamin D and Calcium on Periodontitis. Journal of Periodontology. 2005; 76:1576–1587. [PubMed: 16171451]
- Takaishi Y, Miki T, Nishizawa Y, Morii H. Clinical effect of etidronate on alveolar pyorrhoea associated with chronic marginal periodontitis: report of four cases. J Int Med Res. 2001; 29(4): 355–65. [PubMed: 11675910]

- 27. Jeffcoat, MK.; Reddy, MS. Alveolar bone loss and osteoporosis: Evidence for a common mode of thearpy using the bisphoshonate alendronate. In: Davidovitch, Z.; Norton, L., editors. The Biologic Mechanism of Tooth Resorption and Replacement by Implants. Harvard Society for the Advancement of Orthodontics; Boston: 1996. p. 365-373.
- Rocha M, Nava LE, Vazquez de la Torre C, et al. Clinical and radiological improvement of periodontal disease in patients with type 2 diabetes mellitus treated with alendronate: a randomized, placebo-controlled trial. J Periodontol. 2001; 72(2):204–9. [PubMed: 11288794]
- Lane N, Armitage GC, Loomer P, et al. Bisphosphonate therapy improves the outcome of conventional periodontal treatment: results of a 12-month, randomized, placebo-controlled study. J Periodontol. 2005; 76(7):1113–22. [PubMed: 16018754]
- Marx RE, Sawatari Y, Fortin M, Broumand V. Bisphosphonate-induced exposed bone (osteonecrosis/osteopetrosis) of the jaws: risk factors, recognition, prevention, and treatment. J Oral Maxillofac Surg. 2005; 63(11):1567–75. [PubMed: 16243172]
- Migliorati CA, Schubert MM, Peterson DE, Seneda LM. Bisphosphonate-associated osteonecrosis of mandibular and maxillary bone: an emerging oral complication of supportive cancer therapy. Cancer. 2005; 104(1):83–93. [PubMed: 15929121]
- 32. Grbic JT, Landesberg R, Lin SQ, et al. Incidence of osteonecrosis of the jaw in women with postmenopausal osteoporosis in the health outcomes and reduced incidence with zoledronic acid once yearly pivotal fracture trial. J Am Dent Assoc. 2008; 139(1):32–40. [PubMed: 18167382]
- 33. Payne JB, Zachs NR, Reinhardt RA, Nummikoski PV, Patil K. The association between estrogen status and alveolar bone density changes in postmenopausal women with a history of periodontitis. J Periodontol. 1997; 68(1):24–31. [PubMed: 9029448]
- Krall EA, Dawson-Hughes B, Hannan MT, Wilson PW, Kiel DP. Postmenopausal estrogen replacement and tooth retention. Am J Med. 1997; 102(6):536–42. [PubMed: 9217668]
- Paganini-Hill A. The benefits of estrogen replacement therapy on oral health. The Leisure World cohort. Arch Intern Med. 1995; 155(21):2325–9. [PubMed: 7487257]
- 36. Civitelli R, Pilgram TK, Dotson M, et al. Hormone/estrogen replacement therapy improves alveolar and postcranial bone density in postmenopausal women. Archives of Internal Medicine. 2002
- 37. Food and Nutrition Board. Dietary Reference Intakes (DRI) for Calcium, Phosphorus, Magnesium, Vitamin D, and Fluoride.: Food and Nutrition Board (FNB), Institute of Medicine. National Academy Press; Washington, DC: 1997.

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Frequency and amounts of oral calcium and vitamin D supplementation used by the study population.

					1											
	No c supp n = 1	alcium nor vit dements 138	amin		sup] n = n	cium plements 88					Vit: sup	umin D plements 66				
	Age		<u>Rx I</u>	Drug ^{§§}	Age		Calc	ium	Rx]	Drug	Age		Vita	min D	Rx])rug
	u	<u>mean±SD</u>	ū	<u>%</u>	ū	<u>mean±SD</u>	ū	<u>mg/day</u>	ū	<u>%</u>	ū	mean±SD	ū	IU/day	ū	<u>%</u>
Gender																
Male	82	$64.2{\pm}9.5$	ю	17	19	68.3±13.2	19	632±907	0	0	17	63.9 ± 12.1	17	443±317	0	0
Female	56	58.8 ± 11.9	15	$83^{ m t}$	69	65.6 ± 10.1	69	947±511 <i>‡</i>	15	$100^{\$}$	49	65.9 ± 10.7	49	420±227	11	100
Female																
Menopausal Status																
Pre	6	43.4±7.3	0	0	3	40.3 ± 12.3	3	687±458	-	٢	3	40.3 ± 12.3	3	400 ± 200	-	6
Post	46	$62.1{\pm}10^{**}$	15	100	66	66.7 ± 8.4	99	959±513	14	93	46	67.6 ± 8.3	46	421±231	10	91
Race																
White African	117	$64.0\pm9.8^{\div\uparrow}$	17	94	74	68.2±9.2 <i>††</i>	74	884±658	13	87	59	$67.1{\pm}9.1\%$	59	430±260	10	91
American Indian	15	50.2 ± 9.1	-	9	12	53.9±13	12	819±422	5	13	٢	51.3±16.2	٢	393±164	-	6
Alaska	1	44	0	0	1	67	1	600	0	0						
Asian	4	56.3 ± 16.2	0	0	0	0	0	0	0	0						
Hispanic			0	0	-	60	-	1500	0	0						
* Male versus fem	ale,															
$_{p < 0.01}^{*};$																
$\stackrel{ au}{p}$ < 0.01;																
$t_{\rm p}^{t} < 0.001;$																
$\hat{s}_{p < 0.05}$																
** Pre versus post	, p < 0.	0001														
$^{ au \uparrow \uparrow}$ White versus A	frican .	American, p <1	0.05													

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Table 2

Use of combined calcium and/or vitamin D supplementation in patients with periodontal disease.

		Age		Ca	Λ	itamin D	Rx	Drug	Meet inta	s FNB ikes [§]	Meet inta	s NOF kes ^{**}
	ū	<u>mean±StD</u>	u	<u>mg/day</u>	ū	IU/day	u	<u>%</u>	u	<u>%</u>	u	<u>%</u>
Gender												
Male	15	64.9 ± 12.5	15	670 ± 1010	15	448 ± 338	0	0	7	13	1	17
Female	49	65.9 ± 10.7	49	934 ± 548	49	420 ± 227	Ξ	100	13	87	5	83
<u>Female Menopausal</u>												
Status												
Pre	3	40.3 ± 12.3	З	687 ± 458	3	400 ± 200	-	6	13	100	0	0
Post	46	$67.6\pm8.3^{*}$	46	950 ± 554	46	421 ± 231	10	91	0	0.0	5	100
Race												
White	57	$67.5\pm9.0^{\div}$	57	886 ± 707	57	431 ± 265	10	91	14	93	9	100
African American	٢	51.3 ± 16.2	٢	761 ± 487	٢	393 ± 164	-	6	-	9	0	0
Indian/Alaska	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0	0	0
Asian	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0	0	0
Hispanic	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0	0	0
* Pre versus nost: $n < 0.000$												
1 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	111-11	.005										
ALITCALI ALITERICALI VEISUS	N III	e, p < u.u.;										
T Male versus female, p <	0.01											
$\hat{s}_{\text{Last published recomment}}$	ndatio	ns of the U. S.	Food	and Nutrition I	3oard	(FNB) of the	Institu	te of N	Aedicin	e		

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** National Osteoporosis Foundation Recommendations