Hypovitaminosis D: Which Guidelines for Baseline Supplementation Should Be Followed?

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Background/Purpose: Hypovitaminosis D is prevalent among orthopaedic trauma patients and is a risk factor for fragility fractures as well as bone healing complications. There are two major sets of guidelines that address what level of baseline vitamin D supplementation is appropriate, but they differ significantly in their recommendations. The Institute of Medicine recommends 400 IU daily while the Endocrine Society recommends a higher dose (2000 IU daily). The objectives of this study were to prospectively evaluate risk factors for hypovitaminosis D in an orthopaedic trauma population and to determine the level of baseline supplementation associated with normal vitamin D levels at presentation.

Methods: A prospective observational study was performed in patients undergoing operative treatment for orthopaedic trauma at a Level I trauma center (January to December, 2014). Levels of 25-hydroxy vitamin D (25-OH D) were obtained for 259 patients. Patient and injury characteristics were recorded including age, sex, race, insurance, smoking, body mass index (BMI), comorbidities, preinjury supplementation, and low versus high-energy mechanism. Prevalence of insufficiency (25-OH D <30 ng/mL) and deficiency (25-OH D <20 ng/mL) were determined. Univariate analyses of patient and injury characteristics determined associations with hypovitaminosis D and multivariate logistic regression analysis assessed for independent associations.

Results: Among 259 patients, 191 (73.7%) were vitamin D insufficient and 109 (42.1%) were deficient. 52 patients (20.1%) were receiving preinjury supplementation (200 to 5000 IU daily). Supplementation was more common over age 70 (36 of 99, 36.6%) than below age 70 (17 of 159, 10.7%), P < 0.0001. Univariate predictors of hypovitaminosis D included lack of preinjury supplementation, non-white race, younger age, female sex, non-Medicare insurance, smoking, obesity, Charlson Comorbidity Index <2, and high-energy mechanism. On multivariate analysis only preinjury supplementation (odds ratio [OR] 0.33, 95% CI 0.16-0.71, P = 0.004) and non-white race (OR 4.58, 95% CI 1.94-10.79, P = 0.001) were independently associated with hypovitaminosis D. The 25-OH D level demonstrated a dose-dependent association with baseline vitamin D supplementation. Among those on supplementation, the prevalence of insufficiency was 9 of 11 (81.8%) for <500 IU daily, 17 of 31 (54.8%) for 500 to 1000 IU daily, 8 of 18 (44.4%) for 1000 to 2000 IU daily, and 4 of 16 (25%) for >2000 IU daily. Deficiency (25-OH D <20 ng/mL) was 4 of 11 (36.4%) for <500 IU daily, and 1 of 16 (6.3%) for >2000 IU daily.

Conclusion: Lack of preinjury supplementation and non-white race were independently associated with hypovitaminosis D, which was highly prevalent in the population. Although baseline vitamin D supplementation was infrequent, when present at a sufficient dose it was associated with a very low level of hypovitaminosis D. Given hypovitaminosis

The FDA has stated that it is the responsibility of the physician to determine the FDA clearance status of each drug or medical device he or she wishes to use in clinical practice.

D remained prevalent for supplementation less than 1000 IU daily, baseline supplementation consistent with recommendations from the Endocrine Society (2000 IU daily) appears most effective in this population.

