Does Arthrodesis in the Foot Need to be Delayed Due to Low Vitamin D Level: A Retrospective Study of Concomitant Treatment of Hypovitaminosis

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STATEMENT OF PURPOSE

The role of vitamin D has been demonstrated in osteoporosis and pathologic fractures, but there are limited studies on the impact of vitamin D regarding healing of arthrodesis procedures. Our routine is to identify and treat patients with hypovitaminosis D peri-operatively rather than waiting to order vitamin D levels after a nonunion is established. The purpose of this retrospective case control study is to assess fusion rates of the 1st metatarsal phalangeal joint (1st MTPJ), 1st tarsometatarsal joint (1st TMTJ), and subtalar joint (STJ) with stratified pre-operative vitamin D levels and concomitant peri-operative supplementation.

LITERATURE REVIEW

In conditions of chronic low vitamin D levels, excessive mobilization of bone calcium stores will lead to osteopenia and osteoporosis (1). Several studies have identified the high rate of vitamin D deficiency in the general population (2). In 2014, Smith evaluated 75 patients presenting with low-energy ankle fractures, fifth metatarsal fractures, or stress fractures, and 47% of patients had serum 25(OH)D levels below 30 ng/dl (3). Moore in a 2017 study reviewing 29 patients who had foot and ankle arthrodesis procedures found that patients with vitamin D deficiency or insufficiency were 8.1 times more likely to experience a nonunion (4).

The measurement of 25(OH)D is recommended to measure vitamin D levels (5). The Endocrine Society categorizes vitamin D status into sufficient (>30 ng/ml), insufficient (20-29 ng/ml), and deficient levels (<20 ng/ml) (Table 1) (6). Supplementation recommendations is based on vitamin D status (Table 1).

METHODOLOGY AND PROCEDURE

Lab Screening and Treatment Protocol: Patients undergoing 1st MTPJ, 1st TMTJ and STJ fusion completed a bone health assessment questionnaire (Figure 1). A serum 25-hydroxyvitamin D level was drawn within three months of surgery. The serum 25(OH)D level of each case was recorded in one of three categories: sufficient (> 30 ng/dl), insufficient (20-29 ng/dl), and deficient (10-19 ng/dl). Patients with sufficient levels are not given any supplementation perioperatively. Patients with insufficient levels are recommended over the counter supplementation of 2,000 IU D3 daily. Patients with deficient levels are prescribed 50,000 IU D2 weekly x 12 weeks plus 2,000 IU D3 daily. Surgery is only delayed with deficient levels < 10 ng/dl pending 12 weeks of 50,000 IU D2 weekly plus 2,000 IU D3 daily followed by a repeat course if levels have not improved (Figure 2).

Operative Technique: Joint preparation for fusion consisted of curette and bur for all three arthrodesis procedures. The 1st MTPJ was fixated with a compression screw and dorsal locking plate. The 1st TMTJ and STJ were fixated with two compression screws (Figure 3).

Study Methodology: A retrospective review was performed of 100 cases of foot arthrodesis from 2017-2019 including 1st MTPJ, 1 TMTJ, and STJ fusion. Cases were categorized as "insufficient" or "deficient" vitamin D levels when compared to cases with "sufficient" vitamin D levels regarding 10 week fusion rates.

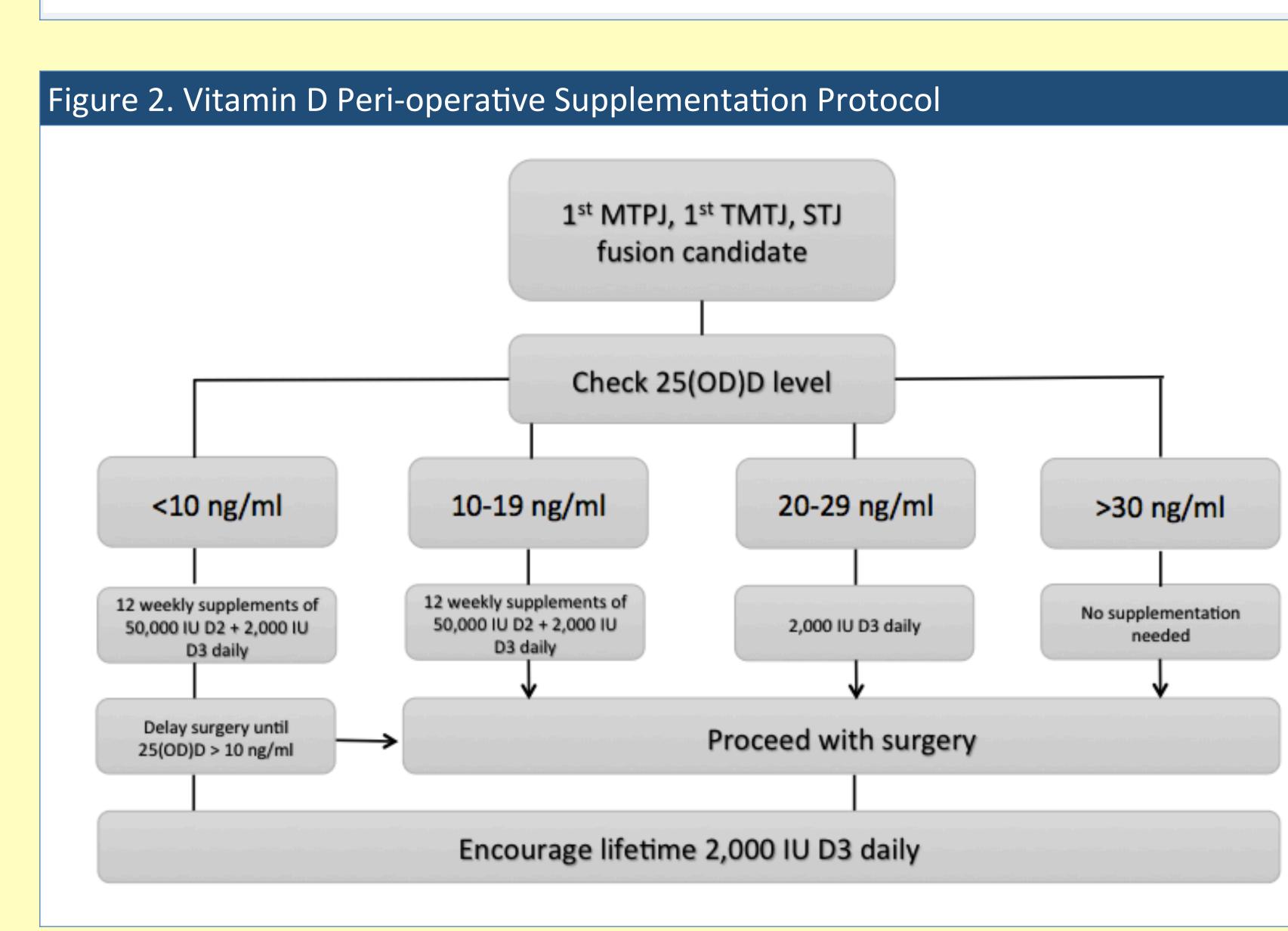
Table 1. Vitamin D Levels and Supplementation Proto						
Vitamin D Status	Serum 25(OH)D Value					
Sufficient	>30 ng/ml					

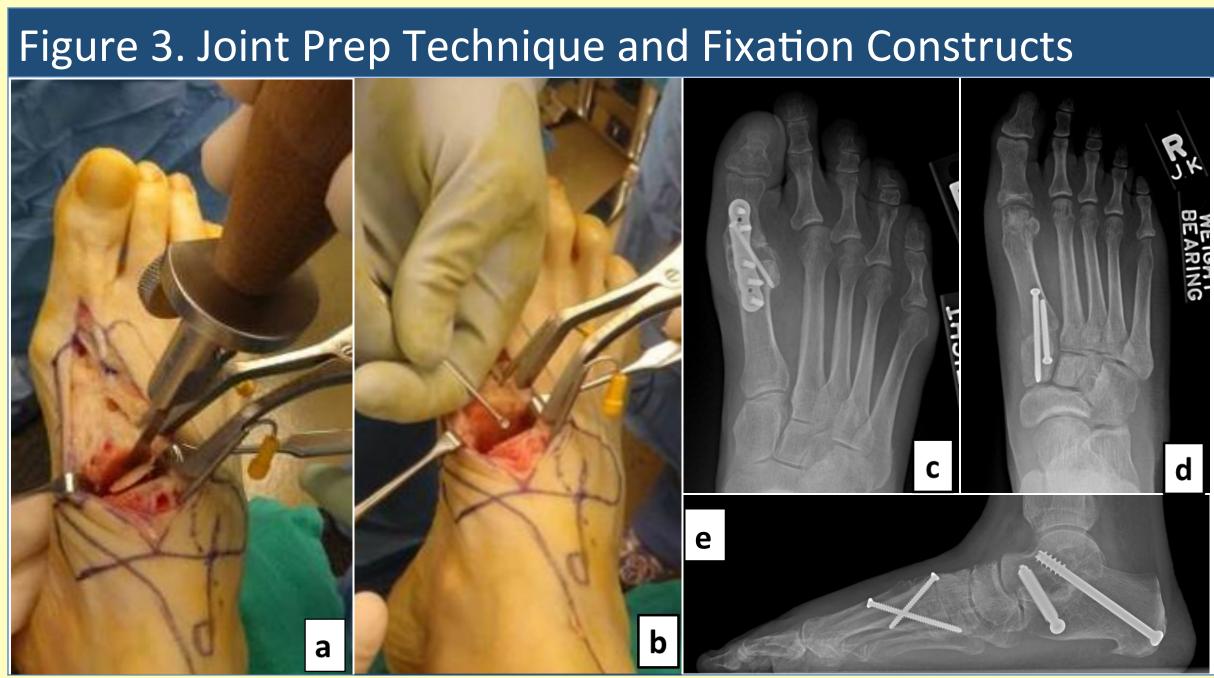
Sufficient Insufficient Deficient

20-29 ng/ml <20 ng/ml

Categories and supplementation recommendations were established by the Endocrine Society 2011 Clinical Practice Guideline

Figure 1. Bone Health Screening P	rotoco				
Have you ever been diagnosed with:					
Osteoporosis/Osteopenia	Yes	No			
Vitamin D Deficiency	Yes	No			
Celiac Disease	Yes	No			
Thyroid Disease	Yes	No			
Parathyroid Disease	Yes	No			
Kidney Disease	Yes	No			
Do you currently take.					
Do you currently take:					
Daily multivitamin	Yes	No			
Calcium supplements	Yes	No			
Vitamin D supplements	Yes	No			
Steroid medication	Yes	No			
Anti-seizure medication	Yes	No			





Supplementation Recommendation

No Supplementation needed

2000 IU D3 daily 50,000 IU D2 weekly x 12 weeks plus

1500-2000 IU D3 daily

Race (Please circle one):

American Indian or Alaskan Native Black or African American White Native Hawaiian or Pacific Islander

- Asian
- If you are a woman, are you post-menopausal? Yes No
- Have you ever had your vitamin D level checked? Yes No
 - If yes, what were the results?
- Have you ever had a DEXA scan?

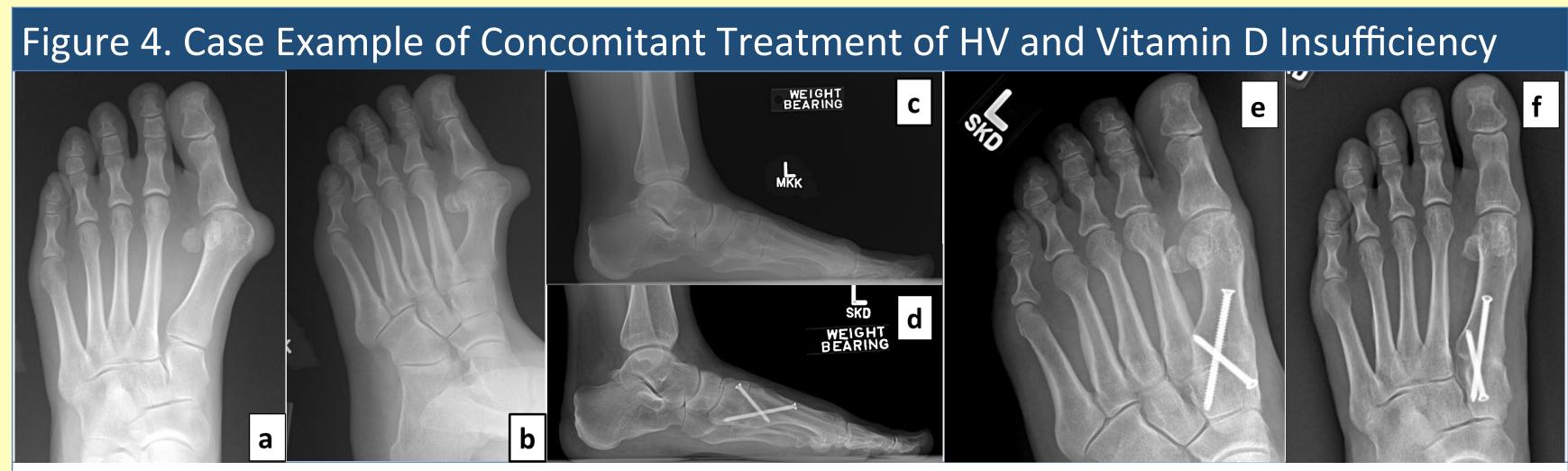
Yes No

If yes, what were the results?

Joint preparation for all fusions was performed with a flexible osteotome (a) and curettage (b) to remove cartilage followed by bur and 2.0 mm drill to penetrate the subchondral d bone plate. Fixation constructs for 1st MTPJ, 1st TMTJ, and STJ fusion are shown here (c-e).

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Table 2. Study Demographics							
Group	Ave. age (range)	Ave. BMI (range)	Gender		Fusion Procedure		
Sufficient	57.4 (19-81)	30.1 (24-35)	12 M	53 F	27 MTPJ	27 TMTJ	11 STJ
Insufficient	55.0 (28-72)	29.6 (27-38)	4 M	19 F	4 MTPJ	16 TMTJ	3 STJ
Deficient	60.8 (44-79)	29.4 (24-35)	2 M	10 F	4 MTPJ	7 TMTJ	1 STJ



44 year old female with a painful bunion deformity evident on x-ray (a-c). Pre-op serum 25(OH)D level was 16 ng/ml. Order placed for 12 weekly doses of 50,000 IU D2 plus long term recommendations for 2000 IU D3 daily. 1st TMTJ fusion was performed without delay with radiographic fusion at the 10 week visit (d-f). Serum 25(OH)D was not rechecked based on union.

RESULTS

There were a total of 100 cases of the selected foot arthrodesis procedures. 65 cases had sufficient vitamin D levels with a 94% (61/65) 10-week fusion rate. 23 cases were vitamin D insufficient and had a 96% (22/23) 10-week fusion rate. 12 cases were vitamin D deficient >10 ng/dl with a 92% (11/12) 10-week fusion. In the vitamin D sufficient group, there were three delayed unions (beyond 10 weeks) and one nonunion that required revision. There was one delayed union in each of the insufficient and deficient groups. By 14 weeks post-operatively all 35 cases in the insufficient and deficient groups achieved radiographic fusion.

Table 3. Fusion Rates Based on Vitamin D Category					
Group	n	10-week fusion rate	14-week fusion rate	P<0.05	
Sufficient (>30 ng/ml)	65	94% (61/65)	98% (64/65)	Control	
Insufficient (20-29 ng/ml)	23	96% (22/23)	100% (23/23)	No stat. sig. diff.	
Deficient (10-19 ng/dl)	12	92% (11/12)	100% (12/12)	No stat. sig. diff.	

ANALYSIS AND DISCUSSION

In this study, 35% of cases were found to have hypovitaminosis D. Surgery was not delayed while waiting for serum vitamin D levels >10 ng/dl to normalize. There was no statistically significant difference in healing rates, and all with hypovitaminosis progressed to osseous healing, suggesting that delaying surgery is not needed to maintain high fusion rates. One case of a nonunion occurred in the sufficient group, in a patient with hypothyroidism who healed successfully after revision surgery. Post treatment vitamin D levels were not routinely obtained; however, this should be considered for patients who demonstrate delayed healing as some patients need a second course at prescription dosage. It remains unknown if no supplementation in these groups would produce poor healing rates. We also selected fusion procedures that have a high fusion rate and these results may not be applicable to ankle or talonavicular fusion.

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