
primary studies - published RCT

A Prospective, Randomized, Double-Blind, Parallel-Group, Comparative Effectiveness Clinical Trial Comparing a Powder Vehicle Compound of Vitamin D With an Oil Vehicle Compound in Adults With Cystic Fibrosis.

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Study design (if review, criteria of inclusion for studies)

Double-blind, randomized controlled, Parallel-Group, Comparative Effectiveness trial

Participants

15 hospitalized adults with CF. The median (interquartile range) age, body mass index, and forced expiratory volume in 1 second were 23.7 (19.9-33.2) years, 19.9 (18.6-22.6) kg/m², and 63% (37%-80%), respectively.

Interventions

Patients were given a one-time bolus dose of 100,000 IU of cholecalciferol (D3) in a powder-based or oil-based vehicle.

Outcome measures

Serum D3, 25-hydroxyvitamin D, and parathyroid hormone concentrations were analyzed at 0, 12, 24, and 48 hours posttreatment. The area under the curve for serum D3 and the 12-hour time point were also assessed as indicators of D3 absorption.

Main results

The increase in serum D3 and the area under the curve was greater in the powder group ($P = .002$ and $P = .036$, respectively). Serum D3 was higher at 12 hours in the powder group compared with the oil group ($P = .002$), although levels were similar between groups by 48 hours.

Authors' conclusions

In adults with CF, cholecalciferol is more efficiently absorbed in a powder compared with an oil vehicle. Physicians should consider prescribing vitamin D in a powder vehicle in patients with CF to improve the absorption of vitamin D from supplements.

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See also

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Keywords

Bone Density Conservation Agents; Bone Diseases; Gastrointestinal Diseases; Pancreas insufficiency; Pancreatic Diseases; pharmacological_intervention; Supplementation; vitamins; Vitamin D; Vitamin D Deficiency; Vitamin deficiencies; Malabsorption; Powders;