

primary studies - published RCT

Intravenous zoledronate improves bone density in adults with cystic fibrosis (CF).

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

Randomized, double-blind, placebo-controlled clinical trial.

Participants

Adult CF outpatient clinics at two hospitals. 22 non-transplanted CF patients aged > or = 18 years with a bone densitometry T-score of

Interventions

Participants were randomized to receive either 2 mg zoledronate i.v. (n = 10) or normal saline (placebo, n = 12) every 3 months for 2 years (8 infusions). All participants received calcium and vitamin D supplements twice daily.

Outcome measures

Percentage change in areal BMD from baseline.

Main results

Lumbar spine BMD increased from baseline more with zoledronate than placebo at 6 months (5.35 +/-0.76 +/-0.76 +/-0.10 +/-1.20%, P = 0.012), 12 months (6.6 +/-1.5 +/-0.15 +/-1.55%, P = 0.011) and 24 months (6.14 +/-1.86 +/-0.10 +/-0.10, P = 0.021). Femoral neck BMD increased more after zoledronate than placebo at 6 months (3.2 +/-1.6 +/-0.43%, P = 0.019), 12 months (4.12 +/-1.8 +/-0.15 +/-1.45 +/-0.43%, P = 0.024) and 24 months (4.23 +/-1.3 +/-1.45 +/-1.4

Authors' conclusions

Intravenous zoledronate was significantly more effective than placebo for increasing BMD in adults with CF and osteopaenia, but side effects limited its tolerability.

http://dx.doi.org/10.1111/j.1365-2265.2008.03434.x

See also

Clin Endocrinol (Oxf). 2009 Jun;70(6):838-46. Epub 2008 Sep 24.

Keywords

Adult; Bisphosphonates; Bone Density Conservation Agents; Bone Diseases; Intravenous; Osteoporosis; pharmacological_intervention; placebo; zoledronate;