

primary studies - published RCT

Effectiveness and tolerability of high-dose salmeterol in cystic fibrosis.

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

Placebo and dual intervention cross-over trial, short-term arm 4 weeks; long-term arm 26 weeks

Participants

44 CF participants (16 males), age range 13 - 37 years. 36 participants finished short-term arm. Initial FEV1 69%

Interventions

Either nebulised albuterol 2.5 mg bd and placebo discus or salmeterol 100 mcg bd by discus and placebo nebuliser

Outcome measures

Changes in spirometry (FEV1, FVC, FEF25-75)measured at baseline, 4 weeks, 12 weeks and 24 weeks Side effects. Need for antibiotic interventions

Main results

Thirty-six out of 44 patients enrolled finished the short-term treatment period, and 19 out of 23 who continued the study also finished the long-term treatment period. There was no significant difference in the mean % change in FEV(1) from baseline to completion of 4 weeks with each drug in the short-term treatment period (0.1% vs. 0.06%, albuterol vs. salmeterol; respectively). In the long-term treatment period, there was a significant decrease from baseline in FEV(1) with albuterol vs. salmeterol, as measured after both 12 and 24 weeks of each treatment (-6.2% vs. 1.8%, P = 0.013 after 12 weeks, and -6.5% vs. 1.7%, P = 0.002, after 24 weeks, respectively). In both treatment periods, salmeterol was well-tolerated. While there were more rescue treatments per patient per week with albuterol than with salmeterol treatment in both the short- and long-term periods (0.67 vs. 0.40 and 1.76 vs. 0.74, respectively), rescue treatments were needed significantly more often for only the long-term period with albuterol compared to salmeterol (P = 0.022). Also, there were more antibiotic interventions with albuterol than with salmeterol treatment in both the short- and long-term period treatment in both the short salmeterol treatment in both the short salmeterol treatment in both the salmeterol treatment in both the short salmeterol treatment in both the short salmeterol (P = 0.022). Also, there were more antibiotic interventions with albuterol than with salmeterol treatment in both the short salmeterol treatment in both the short salmeterol (P = 0.011). In addition, there was a significantly higher symptom score with albuterol vs. salmeterol treatment during the second half of the long-term period (1.24 vs. 0.89, P = 0.001).

Authors' conclusions

long-term high-dose salmeterol was equally safe and was associated with better pulmonary function, fewer interventions, and fewer respiratory symptoms compared to standard therapy with albuterol in a population of outpatients with mild to moderate CF.

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See also

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Keywords

Adolescent; Adult; Albuterol; Anti-Bacterial Agents; Bronchodilator Agents; High-Dose; pharmacological_intervention; salmeterol; Adrenergic beta-Agonists; Respiratory System Agents;