

Clinical changes in periodontal subjects with the probiotic Lactobacillus reuteri Prodentis: A preliminary randomized clinical trial

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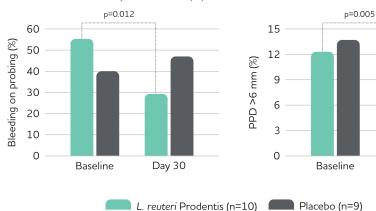
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Demonstrates that Lactobacillus reuteri Prodentis improved clinical periodontal parameters

Results

- Lactobacillus reuteri Prodentis significantly decreased plaque index, bleeding on probing and pocket probing depth of 5-6 mm and ≥6 mm. The placebo group did not show any statistically significant changes in periodontal parameters
- 100 % patient compliance
- No adverse events were reported in any of the groups

Variation of clinical parameters (%) before and after L. reuteri Prodentis treatment



Conclusion

 These data indicate that oral administration of Lactobacillus reuteri Prodentis improved the short-term clinical outcomes in non-smoking patients with initial-to-moderate chronic periodontitis

Day 30

Facts

- Study design: randomized, double-blind, placebo-controlled clinical trial. No professional cleaning was performed during or before the intervention period
- Subjects: 19 adults with initial-to-moderate chronic periodontitis
- Dosage: 1 lozenge daily (2x10⁸ CFU/day)
- Duration: 30 days
- Primary endpoints: plaque index, bleeding on probing, pocket probing depth
- Secondary endpoints: patient compliance and potential side-effects

Further reading

- Teughels W et al. Clinical and microbiological effects of Lactobacillus reuteri probiotics in the treatment of chronic periodontitis: a randomized placebo-controlled study. J Clin Periodontol. 2013;40:1025–1035
- Vivekanda MR et al. Effect of the probiotic Lactobacillus reuteri (Prodentis) in the management of periodontal disease: a preliminary randomized clinical study. J Oral Microbiol. 2010;2:5344-5352
- Twetman S et al. Short-term effect of chewing gums containing probiotic Lactobacillus reuteri on the levels of inflammatory mediators in gingival crevicular fluid. Acta Odontol Scand. 2009;67:19-24

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