

Studies on gastrointestinal and immune health in older children and adults supplemented with the probiotic *Lactobacillus reuteri* DSM 17938 (*Lactobacillus reuteri*) showed several positive effects. There were no clinical safety or tolerance problems.

Immune system effects

REFERENCE	STUDY OBJECTIVES	STUDY DESIGN*	SUBJECTS AND (DAILY DOSE)	RESULTS
Mangalat N, 2012	Primary objective was to investigate the safety of the <i>L. reuteri</i> Protectis drops in healthy adults. Secondary aim was to study changes in some immune factors.	R, DB, PC 2 months with follow-up after 1 and 4 months	<i>L. reuteri</i> : 30 (5 drops/d = 5×10^8 CFU) Placebo: 10	2 months of <i>L. reuteri</i> intake had no significant effect on: <ul style="list-style-type: none"> subclasses of PBMC (peripheral blood mononuclear cells) regulatory T cells (Tregs) TLRs (toll like receptors) 2 and 4 expression cytokine expression by stimulated PBMCs There was a small, significant increase in the faecal calprotectin level, although within the normal clinical range
Böttcher MF, 2008	To evaluate effect on the immunological composition of breast milk (as part of a study on allergy prevention in the offspring). Pregnant women ingested <i>L. reuteri</i> before giving birth.	R, DB, PC 4 weeks before delivery, follow-up after 1 month	<i>L. reuteri</i> : 54 10^8 CFU Placebo: 55	<ul style="list-style-type: none"> Colostrum content of the cytokine TGF-beta2 was significantly reduced while its content of the anti-inflammatory cytokine IL-10 increased The effect was not retained at follow-up
Tubelius P, 2005	To study prevention of short-term illness in healthy adults.	R, DB, PC 80 days	<i>L. reuteri</i> : 94 (1×10^8 CFU) Placebo: 87	<i>L. reuteri</i> significantly reduced short-term sick leave due to cold or GI infection.
Valeur N, 2004	To evaluate effect on immune cells in the gut epithelium in healthy adults.	Open 28 days + 28d follow-up	<i>L. reuteri</i> : 19 (4×10^8 CFU)	<i>L. reuteri</i> significantly increased/stimulated CD4+ T-lymphocytes in the small intestine (ileum).

H. pylori infection

REFERENCE	STUDY OBJECTIVES	STUDY DESIGN*	SUBJECTS AND (DAILY DOSE)	RESULTS
Ojetti V, 2012	Increase the eradication rate of <i>H. pylori</i> and reduce side-effects of 7 days of second line eradication treatment in adults	R, open 14 days + 6w follow-up	<i>L. reuteri</i> : 45 (3×10^8 CFU for 14d) Control: 45	<i>L. reuteri</i> supplementation significantly increased the eradication rate of <i>H. pylori</i> to 80% compared to 60% in the control group.
Francavilla R, 2008	Reduce GI symptoms and bacterial load, and increase eradication rate in <i>H. pylori</i> -infected dyspeptic adults.	R, DB, PC 28 days followed by 10d Hp eradication therapy.	<i>L. reuteri</i> : 20 (1×10^8 CFU) Placebo: 20	<ul style="list-style-type: none"> <i>L. reuteri</i> for 4 weeks significantly: <ul style="list-style-type: none"> Reduced the load of <i>H. pylori</i> Improved GI health scores There was no additional effect on eradication rate.
Scaccianoce G, 2008	Evaluate if <i>L. reuteri</i> (Lr) or a multi-strain probiotic (MSP) could increase the effect of eradication therapy (ET) of <i>H. pylori</i> .	Open, 4 arms: Lr + 7d ET MSP + 7d ET MSP + 14d ET 7d ET (control) 4-6w follow-up.	<i>L. reuteri</i> : 17 (2×10^8 CFU) Probiotic blend: 32 ($4,2 \times 10^{10}$ CFU) for 7 and 14 days Control: 16	No probiotic showed any additional effect on the eradication rate of <i>H. pylori</i> .
Imase K, 2007	Evaluate the effect of <i>L. reuteri</i> on infection load in non-symptomatic <i>H. pylori</i> -infected adults.	R, DB, crossover 4 + 4 weeks.	<i>L. reuteri</i> → Placebo: 15 (4×10^8 CFU) Placebo → <i>L. reuteri</i> : 15 (4×10^8 CFU) <i>L. reuteri</i> → <i>L. reuteri</i> : 5 (4×10^8 CFU) Placebo → placebo: 5	<ul style="list-style-type: none"> <i>L. reuteri</i> significantly reduced <i>H. pylori</i> bacterial load measured by urea breath test The suppressive effect was sustained another 4 weeks in the group testing <i>L. reuteri</i> first and then placebo
Saggiaro A, 2005	Evaluation of the combination of <i>L. reuteri</i> + omeprazole for eradication of <i>H. pylori</i> in symptomatic adults.	R, B 30 days + 4w follow-up.	<i>L. reuteri</i> + omeprazole: 15 (2×10^8 CFU) omeprazole: 15	<ul style="list-style-type: none"> 9/15 (60%) given <i>L. reuteri</i> + omeprazole were completely eradicated of <i>H. pylori</i> Omeprazole alone without effect

* R= randomized, DB= double blind, PC= placebo controlled.  A video presentation of this study is available on www.biogaia.com

Dosage: 1 to 2 tablets a day