

Evaluation of Probiotic Studies Based On The Revised Cochrane Risk-Of-Bias Tool For Randomized Trials (RoB 2)

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Website: <https://sites.google.com/site/riskofbiastool/welcome/rob-2-0-tool?authuser=0>

Details of Study

Name of Study	Lactobacilli reduce recurrences of vaginal candidiasis in pregnant women: a randomized, double-blind, placebo-controlled study
Study Design	Randomized Controlled Trial (Parallel Group)
Probiotic(s) Studied	Lactobacillus plantarum LP115, Lactobacillus helveticus LA25, Lactobacillus rhamnosus LRH10, Lactobacillus paracasei LPC12, Lactobacillus fermentum LF26, Lactobacillus delbrueckii subsp. lactis LDL114
Conclusion on Efficacy	Efficacious
Reference	Ang XY, Chung FY, Lee BK, et al. Lactobacilli reduce recurrences of vaginal candidiasis in pregnant women: a randomized, double-blind, placebo-controlled study. J Appl Microbiol. 2022;132(4):3168-3180. doi:10.1111/jam.15158 (Link to Article)

Risk of bias assessment

Responses underlined in green are potential markers for low risk of bias, and responses in red are potential markers for a risk of bias. Where questions relate only to sign posts to other questions, no formatting is used.

Domain 1: Risk of bias arising from the randomization process

Signalling questions	Response options
1.1 Was the allocation sequence random?	<u>Y</u> / PY / PN / N / NI
1.2 Was the allocation sequence concealed until participants were enrolled and assigned to interventions?	<u>Y</u> / PY / PN / N / NI
1.3 Did baseline differences between intervention groups suggest a problem with the randomization process?	Y / PY / <u>PN</u> / <u>N</u> / NI
Risk-of-bias judgement	<u>Low</u> / High / Some concerns

Domain 2: Risk of bias due to deviations from the intended interventions (*effect of assignment to intervention*)

Signalling questions	Response options
2.1. Were participants aware of their assigned intervention during the trial?	Y / PY / PN / N / NI
2.2. Were carers and people delivering the interventions aware of participants' assigned intervention during the trial?	Y / PY / PN / N / NI
2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context?	NA / Y / PY / PN / N / NI
2.4 If Y/PY to 2.3: Were these deviations likely to have affected the outcome?	NA / Y / PY / PN / N / NI
2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups?	NA / Y / PY / PN / N / NI
2.6 Was an appropriate analysis used to estimate the effect of assignment to intervention?	Y / PY / PN / N / NI
2.7 If N/PN/NI to 2.6: Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?	NA / Y / PY / PN / N / NI
Risk-of-bias judgement	Low / High / Some concerns

Domain 2: Risk of bias due to deviations from the intended interventions (*effect of adhering to intervention*)

Signalling questions	Response options
2.1. Were participants aware of their assigned intervention during the trial?	Y / PY / <u>PN</u> / <u>N</u> / NI
2.2. Were carers and people delivering the interventions aware of participants' assigned intervention during the trial?	Y / PY / <u>PN</u> / <u>N</u> / NI
2.3. [If applicable:] If <u>Y</u> / <u>PY</u> /NI to 2.1 or 2.2: Were important non-protocol interventions balanced across intervention groups?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
2.4. [If applicable:] Were there failures in implementing the intervention that could have affected the outcome?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
2.5. [If applicable:] Was there non-adherence to the assigned intervention regimen that could have affected participants' outcomes?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
2.6. If <u>N</u> / <u>PN</u> /NI to 2.3, or <u>Y</u> / <u>PY</u> /NI to 2.4 or 2.5: Was an appropriate analysis used to estimate the effect of adhering to the intervention?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
Risk-of-bias judgement	Low / High / Some concerns

Domain 3: Missing outcome data

Signalling questions	Response options
3.1 Were data for this outcome available for all, or nearly all, participants randomized?	<u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
3.2 If <u>N/PN/NI</u> to 3.1: Is there evidence that the result was not biased by missing outcome data?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u>
3.3 If <u>N/PN</u> to 3.2: Could missingness in the outcome depend on its true value?	NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
3.4 If <u>Y/PY/NI</u> to 3.3: Is it likely that missingness in the outcome depended on its true value?	<u>NA</u> / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
Risk-of-bias judgement	<u>Low</u> / High / Some concerns

Domain 4: Risk of bias in measurement of the outcome

Signalling questions	Response options
4.1 Was the method of measuring the outcome inappropriate?	Y / PY / PN / N / NI
4.2 Could measurement or ascertainment of the outcome have differed between intervention groups?	Y / PY / PN / N / NI
4.3 If N/PN/NI to 4.1 and 4.2: Were outcome assessors aware of the intervention received by study participants?	NA / Y / PY / PN / N / NI
4.4 If Y/PY/NI to 4.3: Could assessment of the outcome have been influenced by knowledge of intervention received?	NA / Y / PY / PN / N / NI
4.5 If Y/PY/NI to 4.4: Is it likely that assessment of the outcome was influenced by knowledge of intervention received?	NA / Y / PY / PN / N / NI
Risk-of-bias judgement	Low / High / Some concerns

Domain 5: Risk of bias in selection of the reported result

Signalling questions	Response options
5.1 Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?	<u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
Is the numerical result being assessed likely to have been selected, on the basis of the results, from...	
5.2. ... multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?	<u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
5.3 ... multiple eligible analyses of the data?	<u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI
Risk-of-bias judgement	Low / <u>High</u> / Some concerns

Overall risk of bias

Risk-of-bias judgement	Comments:	Low / High / Some concerns
	<ul style="list-style-type: none"> In assessing reduction in VC symptoms using the seven questions in the VSQ, there were only 3 significant differences observed in the probiotic group compared to the placebo group. The probiotic group did not show any improvement in terms of four other symptoms (itching, hurting, dry and foul-smelling vulva) compared to the placebo group. 	