

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

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Tool adapted from: Sterne JAC, Savović J, Page MJ, Elbers RG, Blencowe NS, Boutron I, Cates CJ, Cheng H-Y, Corbett MS, Eldridge SM, Hernán MA, Hopewell S, Hróbjartsson A, Junqueira DR, Jüni P, Kirkham JJ, Lasserson T, Li T, McAleenan A, Reeves BC, Shepperd S, Shrier I, Stewart LA, Tilling K, White IR, Whiting PF, Higgins JPT. RoB 2: a revised tool for assessing risk of bias in randomised trials. *BMJ* 2019; **366**: l4898.

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

Details of Study

| | |
|-------------------------------|---|
| Name of Study | Efficacy and safety of vaginally administered lyophilized Lactobacillus crispatus IP 174178 in the prevention of bacterial vaginosis recurrence |
| Study Design | Randomized Controlled Trial (Parallel Group) |
| Probiotic(s) Studied | Lactobacillus crispatus IP174178 |
| Conclusion on Efficacy | Efficacious |
| Reference | Bohbot JM, Darai E, Bretelle F, Bami G, Daniel C, Cardot JM. Efficacy and safety of vaginally administered lyophilized Lactobacillus crispatus IP 174178 in the prevention of bacterial vaginosis recurrence. Journal of gynecology obstetrics and human reproduction. 2018;47(2):81-6. (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> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI |
| 1.2 Was the allocation sequence concealed until participants were enrolled and assigned to interventions? | <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / 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 / <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 <u>Y/PY/NI</u> to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? | NA / Y / PY / <u>PN</u> / <u>N</u> / NI |
| 2.4 If <u>Y/PY</u> to 2.3: Were these deviations likely to have affected the outcome? | NA / Y / PY / <u>PN</u> / <u>N</u> / NI |
| 2.5. If <u>Y/PY/NI</u> to 2.4: Were these deviations from intended intervention balanced between groups? | NA / <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI |
| 2.6 Was an appropriate analysis used to estimate the effect of assignment to intervention? | <u>Y</u> / <u>PY</u> / <u>PN</u> / <u>N</u> / NI |
| 2.7 If <u>N/PN/NI</u> 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 / <u>PN</u> / <u>N</u> / 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 Y/PY/NI to 2.1 or 2.2: Were important non-protocol interventions balanced across intervention groups? | NA / <u>Y</u> / PY / <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> / PY / <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> / PY / <u>PN</u> / <u>N</u> / NI |
| 2.6. If N/PN/NI to 2.3, or Y/PY/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> / PY / <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> / PN / N / NI |
| 3.2 If N/PN/NI to 3.1: Is there evidence that the result was not biased by missing outcome data? | NA / <u>Y</u> / <u>PY</u> / PN / N |
| 3.3 If N/PN to 3.2: Could missingness in the outcome depend on its true value? | NA / Y / PY / <u>PN</u> / <u>N</u> / NI |
| 3.4 If Y/PY/NI to 3.3: Is it likely that missingness in the outcome depended on its true value? | NA / Y / PY / <u>PN</u> / <u>N</u> / NI |
| Risk-of-bias judgement | Low / 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 / <u>PN</u> / <u>N</u> / NI |
| 4.2 Could measurement or ascertainment of the outcome have differed between intervention groups? | Y / PY / <u>PN</u> / <u>N</u> / NI |
| 4.3 If <u>N</u> / <u>PN</u> /NI to 4.1 and 4.2: Were outcome assessors aware of the intervention received by study participants? | NA / Y / PY / <u>PN</u> / <u>N</u> / NI |
| 4.4 If <u>Y</u> / <u>PY</u> /NI to 4.3: Could assessment of the outcome have been influenced by knowledge of intervention received? | <u>NA</u> / Y / PY / <u>PN</u> / <u>N</u> / NI |
| 4.5 If <u>Y</u> / <u>PY</u> /NI to 4.4: Is it likely that assessment of the outcome was influenced by knowledge of intervention received? | <u>NA</u> / Y / PY / <u>PN</u> / <u>N</u> / NI |
| Risk-of-bias judgement | <u>Low</u> / 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? | Y / PY / <u>PN</u> / <u>N</u> / NI |
| 5.3 ... multiple eligible analyses of the data? | Y / PY / <u>PN</u> / <u>N</u> / NI |
| Risk-of-bias judgement | <u>Low</u> / High / Some concerns |

Overall risk of bias

| | | |
|-------------------------------|--|-----------------------------------|
| Risk-of-bias judgement | Comments: <ul style="list-style-type: none"> The minimum sample size was not met due to small sample size and dropout hence statistical power was not achieved. | Low / High / Some concerns |
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