Applying Study Results to Patient Care



Updated May 2025

Also see our Glossary of Study Design and Statistical Terms.

Statistically Significant



 Results of a trial are statistically significant when p-values are less than a pre-specified number, usually 0.05.¹

Number Needed to Treat (NNT)



- Estimated number of patients that would need to be treated to prevent one outcome compared to a control group, over a specified time period.³
- Rounded up to a whole number.³
- Only appropriate for statistically significant differences.³
- NNT = 1/ ARR expressed as a decimal

Relative Risk (RR)



- The risk of an outcome in the treatment group divided by the risk of the outcome in the control group.⁴
- **RR = 1**: no association between treatment and outcome.
- RR >1: positive association between treatment and outcome (e.g., bleeding while taking an anticoagulant).
- RR <1: negative association between treatment and outcome (e.g., decreased risk of thrombosis while taking an anticoagulant).

p-value



- The probability that an outcome is due to chance.²
- A p-value less than 0.05 means there is less than a 5% chance that the result is due to chance.²

Number Needed to Harm (NNH)



- **Estimated** number of patients treated with a specified therapy/intervention in order for one to have a bad outcome, over a specified time period.⁴
- Rounded down to a whole number.⁴
- Only appropriate for statistically significant differences.
- NNH = 1/ Absolute risk increase expressed as a decimal

Clinically Significant



- Clinically significant results are meaningful or important enough to change clinical practice.¹
- Number needed to treat (NNT) and number needed to harm (NNH) are simple calculations that estimate the importance of study results to patients.

Confidence Interval (CI)



- A range of values which likely contains the true value.
- With a 95% CI, we're 95% sure that the true value lies in the reported range.⁵ If the same population is sampled repeatedly, approximately 95% of the calculated confidence intervals will contain the true value.

Relative Risk Reduction (RRR)



- The change in event rate between a treatment group and a control group.⁶
- RRR = 1 (treatment rate/control rate).⁶
- Often misinterpreted as an overestimation of treatment effect.

Absolute Risk Reduction (ARR)



- The absolute change in event rates between two groups.⁶
- ARR = control rate treatment rate
- Less commonly reported, and may need to be calculated to see true treatment effect.

Odds Ratio (OR)



- The odds of exposure in cases divided by the odds of exposure in controls.^{5,6}
- Analogous to relative risk; however, OR can be used in case-control studies.
- Can exaggerate risk when disease/outcome is common.^{5,6}

Composite Endpoints



 A combination of (similarly valued) endpoints that each have a low incidence but together may provide a single measure of effect.⁷

Non-inferiority Trial



- Designed to assess if the treatment being studied is not inferior to a control/currently used treatment.⁴
- Requires fewer patients compared to a superiority study.⁴



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