

## Submission and reviewer's abstract checklist for intervention studies<sup>a</sup> and outbreak reports

<sup>a</sup> carried out either to reduce infection or to improve compliance with infection control measures such as hand hygiene, antibiotic prescription or care bundle implementation

|   |  |
|---|--|
| <b>Title</b>  | <b>1.</b> Clear statement that this is an intervention study or outbreak report.   |
| <b>Background</b>   | <b>2.</b> Rationale for study with clear hypothesis for intervention studies or objective for outbreak reports.  |
| <b>Methods</b>  | <b>3.</b> Clear statement of intervention study design <sup>1</sup> or case definition for outbreak report.  |
|   | <b>4.</b> Brief description of setting, participants, and intervention or outbreak control measures with start and stop dates.   |
|   | <b>5.</b> Clearly defined outcomes & denominators at regular time intervals <sup>2</sup> , not as totals for each phase (can be in results).   |
|   | <b>6.</b> Statistical analysis accounts for any dependencies in the data (can be in results instead).<br>(Statistical analysis may not be appropriate for outbreak reports).   |
|   | <b>7.</b> Which potential confounders or biases were considered, recorded or adjusted for <sup>3</sup> (can be in results instead)   |
| <b>8.</b> Where relevant: details of culture, typing, environmental sampling, and risk factors for acquisition, root cause analysis or organisational risk assessment |  |
| <b>Results</b>  | <b>9.</b> For the main outcomes: estimated effect size & its precision (usually using 95% C.I.)<br>(A graphical summary is often appropriate for dependent data -such as most time series).  |
| <b>Conclusions</b>  | <b>10.</b> For intervention studies: consider in relation to original hypothesis, accounting for potential confounders & biases.<br>For outbreak reports: consider clinical significance of observations and hypothesis to explain them. |

<sup>1</sup> e.g. Interrupted Time Series, Cluster or other Randomised Controlled Trial, Cross over, Controlled Before and After intervention, Uncontrolled Before and After Intervention (see explanatory document and [www.ccg.cochrane.org/en/newPage1.html](http://www.ccg.cochrane.org/en/newPage1.html) for standard terminology).

<sup>2</sup> at least 3 time points per phase and for many two phase studies 12 or more monthly data points

<sup>3</sup> e.g. changes in length of stay, case mix, bed occupancy, staffing levels, hand-hygiene compliance, antibiotic use, strain type, processing of isolates, seasonality, other interventions, incomplete blinding, fidelity to intervention, non standardised outcome assessment