

Multimedia Appendix 2: Description of items and scores used for the quality assessment of the included studies.

| Item Category | Item Code | Max Score | Description of quality assessment item |
|---|-----------|-----------|---|
| Reporting | A | 1 | Is the hypothesis/aim/objective of the study clearly described? |
| | B | 1 | Are the main outcomes to be measured clearly described in the Introduction or Methods section? |
| | C | 1 | Are the characteristics of the patients included in the study clearly described? |
| | D | 1 | Are the interventions of interest clearly described? |
| | E | 2 | Are the distributions of principal confounders in each group of subjects to be compared clearly described? |
| | F | 1 | Are the main findings of the study clearly described? |
| | G | 1 | Does the study provide estimates of the random variability in the data for the main outcomes? |
| | H | 1 | Have all important adverse events that may be a consequence of the intervention been reported? |
| | I | 1 | Have the characteristics of patients lost to follow-up been described? |
| | J | 1 | Have actual probability values been reported for the main outcomes except where the probability value is less than 0.001? |
| External validity | K | 1 | Were the subjects asked to participate in the study representative of the entire population from which they were recruited? |
| | L | 1 | Were those subjects who were prepared to participate representative of the entire population from which they were recruited? |
| | M | 1 | Were the staff, places, and facilities where the patients were treated, representative of the treatment the majority of patients received? |
| Internal validity-bias | N | 1 | Was an attempt made to blind study subjects to the intervention they have received? |
| | O | 1 | Was an attempt made to blind those measuring the main outcomes of the intervention? |
| | P | 1 | If any of the results of the study were based on "data dredging", was this made clear? |
| | Q | 1 | In trials and cohort studies, do the analyses adjust for different lengths of follow-up of patients, or in case-control studies, is the time period between the intervention and outcome the same for cases and controls? |
| | R | 1 | Were the statistical tests used to assess the main outcomes appropriately? |
| | S | 1 | Was compliance with the intervention/s reliable? |
| | T | 1 | Were the main outcome measures used accurate? |
| Internal validity-confounding (selection bias) | U | 1 | Were the patients in different intervention groups or were the cases and controls recruited from the same population? |
| | V | 1 | Were study subjects in different intervention groups or were the cases and controls recruited over the same period of time? |
| | W | 1 | Were the study subjects randomized to intervention groups? |
| | X | 1 | Was the randomized intervention assignment concealed from both patients and health care staff until recruitment was completed and irrevocable? |
| | Y | 1 | Was there adequate adjustment for confounding in the analyses from which the main findings were drawn? |
| | Z | 1 | Were losses of patients to follow-up taken into account? |
| Power | AA | 1 | Did the study have sufficient power to detect a clinically important effect where the probability value for a difference being due to chance is less than 5%? |