## Webtable 2 [as supplied by authors]: Quality assessment of included studies

| Study ID                       | Generation of<br>allocation<br>sequence<br>adequate <sup>a</sup>        | Concealed allocation | No differences<br>in treatment of<br>experimental<br>and control<br>group <sup>b</sup> | Outcome of<br>interest of the<br>systematic<br>review fully<br>defined <sup>c</sup> | Outcome<br>assessor<br>blinded <sup>d</sup> | Patients analysed in the group in which they were randomised | All<br>randomised<br>patients<br>analysed | Drop-out/<br>withdrawals<br>reported and<br>analysed<br>Number of<br>drop-outs <sup>e</sup> |
|--------------------------------|---|----------------------|--|---|---|--|---|---|
| Kissinger 2006 <sup>w1</sup>   | Yes   | Unclear              | Yes  | Yes <sup>f</sup>  | No  | Unclear  | No  | Partly (no<br>numbers per<br>group reported)<br>Overall: 87<br>(19%)                        |
| Golden 2005 <sup>w3</sup>      | Unclear   | Unclear              | No (condoms<br>and info on<br>STIs in<br>experimental<br>group only)                   | Yes   | No  | Unclear  | No  | Yes  Experimental: 446 (32%)  Control: 445 (32%)  |
| Kissinger 2005 <sup>w2</sup>   | No (allocation according to month of admission; months were randomised) | No                   | Yes  | Yes <sup>f</sup>  | No  | Unclear  | Unclear                                   | Partly (no<br>numbers per<br>group reported)<br>Overall: 207<br>(21%)                       |
| Schillinger 2003 <sup>w4</sup> | Unclear   | Yes                  | No (advice on sexual behaviour and info on STIs in experimental group only)            | Yes   | No  | Yes  | No  | Yes Experimental: 218 (23%) Control: 217 (23%)  |
| Ostergaard 2003 <sup>w8</sup>  | Unclear   | Unclear              | Yes  | Partly<br>(timeframe<br>unclear)  | Unclear                                     | Unclear  | Yes                                       | n/a   |
| Moyo 2002 <sup>w9</sup>        | Yes   | Yes                  | No (health care<br>voucher for<br>partners in<br>experimental<br>group only)           | Partly<br>(timeframe<br>unclear)  | No  | Yes  | No  | Yes Experimental: 70 (51%) Control: 68 (51%)  |

| Study ID                         | Generation of<br>allocation<br>sequence<br>adequate <sup>a</sup> | Concealed allocation | No differences<br>in treatment of<br>experimental<br>and control<br>group <sup>b</sup>                             | Outcome of<br>interest of the<br>systematic<br>review fully<br>defined <sup>c</sup> | Outcome<br>assessor<br>blinded <sup>d</sup> | Patients analysed in the group in which they were randomised | All<br>randomised<br>patients<br>analysed | Drop-out/<br>withdrawals<br>reported and<br>analysed<br>Number of<br>drop-outs <sup>e</sup> |
|----------------------------------|--|----------------------|--|---|---|--|---|---|
| Nuwaha 2001 <sup>w5</sup>        | Yes  | Unclear              | No (outcome<br>assessment<br>biased in favour<br>of experimental<br>group)   | Yes <sup>g</sup>  | No  | Unclear  | Yes                                       | No (analysis<br>based on<br>partners not on<br>index patients)                              |
| Kissinger 1998 <sup>w6</sup>     | Unclear <sup>h</sup>   | Unclear <sup>h</sup> | No (different providers)   | Partly<br>(timeframe<br>unclear)  | No  | Unclear  | No  | No (no information on all randomised patients)  |
| Andersen 1998 <sup>w7</sup>      | No (allocation according to date of birth)                       | No                   | Yes  | Partly<br>(timeframe<br>unclear)  | No  | Unclear  | Yes                                       | n/a   |
| Faxelid 1996 <sup>w10</sup>      | Yes  | Unclear              | No (contact cards in experimental group only; unclear if provider referral was offered in experimental group only) | Partly<br>(timeframe<br>unclear)  | No  | Unclear  | Yes                                       | Partly (no analysis reported)  Experimental: 8 (4%)  Control: 11 (6%)                       |
| Katz 1988 <sup>w11</sup>         | Unclear  | Unclear              | Yes  | Partly<br>(timeframe<br>unclear)  | No  | Yes  | Unclear                                   | n/a   |
| Solomon 1988 <sup>w12</sup>      | Unclear  | Unclear              | Yes  | Partly<br>(timeframe<br>unclear)  | No  | Unclear  | Unclear                                   | n/a   |
| Cleveland undated <sup>w13</sup> | Unclear  | Unclear              | Yes  | Yes   | No  | Unclear  | Yes                                       | n/a   |
| Ellison undatedw14i              | Unclear  | Unclear              | Unclear  | Unclear   | Yes   | Unclear  | Yes                                       | n/a   |

## Legend:

STI, sexually transmitted infection; superscripts w1-14 refer to the reference list available at www.bmj.com

- <sup>a</sup> Computer generated random numbers, tables of random numbers, drawing lots.
- If some of the intervention are viewed as complex interventions with several components assessment would change from "No" to "Yes" for four trials was was was was assessment would change from "No" to "Yes" for four trials was was was assessment would change from "No" to "Yes" for four trials was was was assessment would change from "No" to "Yes" for four trials was was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was was assessment would change from "No" to "Yes" for four trials was assessment would change from "No" to "Yes" for four trials was assessment would change from "No" to "Yes" for four trials was assessment was
- <sup>c</sup> Timeframe and measuring method defined.
- d Open assessment of outcomes assumed if blinding not explicitly mentioned.
- e Only assessed for the primary outcome of the individual study.
- Outcome assessment of the outcome 'partners treated' potentially biased in favor of experimental group (assessment based on index patients).
- Outcome assessment of the outcome 'partners treated' potentially biased (assessment based on index patients but only index patients in experimental group were refunded for return transport cost; outcome assessment differed between groups: interview of index patients + linking medical records of patients attending the clinic to index patients in control group versus interview of index patients only in experimental group).
- h Patients were randomised to providers. One provider worked fewer hours per week and used PDPT. The other providers in the center used patient referral.
- Unpublished study and study report not available for assessment. Quality assessment taken from Mathews et al.<sup>3</sup>