Table S1. GRADE table

| Quality assessment  |                         |                |                   |                  |                      |                       | № of patients       |                                  |                               | Effect  |                            | l<br>m                   |
|---------------------|-------------------------|----------------|-------------------|------------------|----------------------|-----------------------|---------------------|----------------------------------|-------------------------------|---|----------------------------|--------------------------|
| № of<br>studie<br>s | Study<br>design         | Risk of bias   | Inconsist<br>ency | Indirectne<br>ss | Impreci<br>sion      | Other conside rations | Women<br>self-admin | Healthcare<br>providers<br>admin | Relative<br>(95% CI)          | Absolute<br>(95% CI)                              | Certainty of the evidence  | p<br>or<br>ta<br>n<br>ce |
| continuat           | tion (non-interrup      | oted use at 1  | 2 months) RC      | T                |                      |                       |                     |                                  |                               |   |                            |                          |
| 1                   | randomised<br>trial     | not<br>serious | not<br>serious    | not serious      | very<br>serious<br>2 | none                  | 28/86<br>(32.6%)    | 14/46<br>(30.4%)                 | RR 1.07<br>(0.63 to<br>1.82)  | 21 more per 1000 (from 113 fewer to 250 more)     | $\bigoplus_{LOW} \bigcirc$ |                          |
| continuat           | tion (non-interrup      | oted use at '  | 2 months) no      | n-RCT            |                      |                       |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 51/58<br>(87.9%)    | 50/64<br>(78.1%)                 | RR 1.13<br>(0.96 to<br>1.32)  | 102 more per 1000 (from 31 fewer to 250 more)     | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| continua            | ation (non-interru      | pted use at    | 3 months) nor     | n-RCT            |                      |                       |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious<br>2 | none                  | 10/10<br>(100.0%)   | 10/10<br>(100.0%)                | RR 1<br>(0.83 to<br>1.2)      | 0 fewer per 1000 (from 170<br>fewer to 200 more)  | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| Safety: s           | erious adverse e        | events         |                   |                  |                      |                       |                     |                                  |                               |   |                            |                          |
| 0                   |                         |                |                   |                  |                      |                       |                     |                                  |                               |   |                            |                          |
| Safety: o           | ther complication       | ns             |                   |                  |                      |                       |                     |                                  |                               |   |                            |                          |
| 1                   |                         |                |                   |                  |                      |                       |                     |                                  | not<br>estimable              | not estimable                                     |                            |                          |
| Satisfacti          | ion with the conti      | raception m    | ethod (want to    | continue this r  | method) at 1         | 12 months             |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 55/61<br>(90.2%)    | 50/55<br>(90.9%)                 | RR 0.99<br>(0.88 to<br>1.12)  | 9 fewer per 1000 (from 109<br>more to 109 fewer)  | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| Satisfacti          | ion with the conti      | raception m    | ethod (prefer t   | o continue the   | method) at           | 3 months              |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 10/10<br>(100.0%)   | 8/10<br>(80.0%)                  | <b>RR 1.24</b> (0.87 to 1.75) | 192 more per 1000 (from<br>600 more to 104 fewer) | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| Satisfacti          | ion with the cont       | raception m    | ethod (recomr     | nend to a friend | d) at 12 mor         | nths                  |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 57/61<br>(93.4%)    | 53/53<br>(100.0%)                | <b>RR 0.94</b> (0.87 to 1.01) | 60 fewer per 1000 (from 130 fewer to 10 more)     | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| Satisfacti          | ion with the cont       | raception m    | ethod (recomr     | nend to a friend | d) at 3 mont         | hs                    |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 9/10<br>(90.0%)     | 8/10<br>(80.0%)                  | RR 1.13<br>(0.78 to<br>1.63)  | 104 more per 1000 (from<br>176 fewer to 504 more) | ⊕⊖⊖⊖<br>VERY LOW           |                          |
| Satisfacti          | ion in location (a      | t 3 months)    |                   |                  |                      |                       |                     |                                  |                               |   |                            |                          |
| 1                   | observationa<br>I study | serious<br>1   | not<br>serious    | not serious      | very<br>serious      | none                  | 9/10<br>(90.0%)     | 9/10<br>(90.0%)                  | <b>RR 1</b> (0.75 to 1.34)    | 0 fewer per 1000 (from 225 fewer to 306 more)     | ⊕⊖⊖⊖<br>VERY LOW           |                          |

MD - mean difference, RR - relative risk

High certainty evidence: Further research is very unlikely to change our certainty of the estimate of effect. Moderate certainty evidence: Further research is likely to have an important impact on the certainty of the estimate of effect and may change the estimate. Low certainty evidence: Further research is very likely to have an important impact on the certainty of the estimate of effect and is likely to change the estimate or any estimate of effect is very uncertain. Very low certainty evidence: Any estimate of effect is very

High risk of bias in included study
One study only with few participants