Managing partners of people diagnosed with Chlamydia trachomatis: a comparison of two partner testing methods

L Østergaard, B Andersen, J K Møller, F Olesen, A-M Worm

Objective: To compare the effectiveness of “home sampling” with that of “office sampling” for testing partners to men and women infected with Chlamydia trachomatis.

Method: A randomised controlled effectiveness trial took place in the general community in Denmark. 1300 index women and 526 index men (>18 years) with a positive test result for C trachomatis were identified. Of these, 414 index women and 148 index men gave implied consent. Index patients were randomly assigned to provide their partner(s) through the past 12 months with either (1) a kit by which partner(s) could be tested by home sampling, or (2) a kit by which partner(s) could only be tested by seeing a healthcare professional (office sampling). The main outcome measure was the proportion of index patients who had at least one partner tested for C trachomatis.

Results: The proportion of index women with at least one partner tested was higher in the home sampling group (0.26) than in the office group (0.12) (difference 0.14; 95% CI 0.10 to 0.19; p<0.0001) and so it was for index men (0.15 vs 0.03; difference 0.12; 95% CI 0.07 to 0.16; p<0.0001). Also the proportion of index women for whom at least one partner was identified as infected was higher in the home sampling group compared with the office group (0.11 vs 0.07, p=0.01). The corresponding figures for index men were 0.06 vs 0.01, p=0.02.

Conclusion: The effectiveness of partner testing is higher when partners of C trachomatis infected patients are offered home sampling than when they are offered office sampling.

Key messages

- A partner testing strategy based on home sampling shows a twofold to fourfold higher effectiveness than a partner testing strategy based on samples obtained in a medical office
- The improved effectiveness of home sampling was relatively more pronounced for index men compared with index women
- A strategy of offering partners the opportunity of home sampling might contribute to improved public health control of the Chlamydia trachomatis infection

Recruitment of index patients

Whenever a swab sample was found to be positive for C trachomatis in the routine laboratory, individuals with a positive test result were thus randomised to give their partners the opportunity of “home sampling” or to give their partners the opportunity of “office sampling.”
The package to the index patients
The package to the index patients was mailed to his/her home address based on information provided by the healthcare provider. The package consisted of five specimen collection kits to be used by his/her sexual partners over the previous 12 months. This time period was decided based on advice from several European experts. The specimen collection kits for the two study groups were identical and the content of the specimen collection kit was thus blinded to the index patient.

The index patient was instructed to give the specimen collection kits to the partner(s) or mail them (post paid). In case the index patient chose not to forward the partner kits (dropout) they were asked to give the reason for this.

Specimen collection kits for partners
After receiving the specimen collection kits, partners in the home sampling group should mail their home obtained samples directly to the diagnostic laboratory in post paid and preaddressed envelopes. Partners in the office sampling group had to bring the specimen collection kit into the office of a healthcare provider to obtain a sample (office sampling). Partners in this group also brought a letter explaining the study and the importance that the health care provider used the provided specimen collection kit.

Content of partner kits
For male partners in the home sampling group the kit contained a 10 ml tube for a first void urine sample and for female partners the kit consisted of a vaginal pipette containing 5 ml sterile normal saline. Women introduced the tip of the pipette 3 cm into the vagina and flushed and aspirated the saline. We previously found a high patient acceptability of this vaginal pipette.

For the “office sampling” group the preaddressed envelope was replaced by a request to bring the specimen collection kit to a health care provider for testing. The healthcare provider then subsequently mailed the sample to the study centre.

Testing for C trachomatis
The initial swab sample, on which the identification of possible index patients was obtained, was analysed by the LCx Chlamydia trachomatis test (Abbott, Chicago, IL, USA). This also held for vaginal flush samples and urine samples and both home obtained and office obtained samples.

Follow up and treatment of the C trachomatis infection
All partners had their test result sent back to their own address or their physician’s address depending on their marked preference. If the test was positive, the patient was requested to seek a healthcare provider for treatment.

Study end point
The primary end point of the study was the proportion of index patients with at least one partner tested for C trachomatis. A secondary end point was the proportion of index patients with at least one partner having a C trachomatis infection diagnosed.

Definition of study populations
“Intention to test” population
This consisted of all patients who had a positive test for C trachomatis identified.

Included patients
The group of the “intention to test” population who accepted to receive information about the study.

RESULTS
Index women
A total of 1300 C trachomatis infected women were identified. A chart of patient flow is shown in figure 1. To compare the characteristics between the two groups of patients who actually participated (the recruited patients), key features such as previous sexually transmitted infections, symptoms, and sexual behaviour were collected. The characteristics of the 191 (28.8%; 191 of 663) recruited index women in the office group were similar to the characteristics of the 223 (35.0%; 223 of 637) recruited index women in the home sampling group (table 1).

Index men
A total of 526 C trachomatis infected men were identified (fig 2). A chart of patient flow is shown in figure 2. The characteristics of the 67 (24.9%; 67 of 269) recruited index men who actually participated (the recruited patients), key features such as previous sexually transmitted infections, symptoms, and sexual behaviour were collected. The characteristics of the 81 (30.2%; 81 of 269) recruited index men in the office group were similar to the characteristics of the 47 (17.6%; 47 of 269) recruited index men in the home sampling group (table 2).
men in the office group were similar to the characteristics of the 81 (31.5%; 81 of 257) recruited index men in the home sampling group (table 1).

Reasons for withdrawals (dropout)
The major reported reason for index patients dropping out of the study was that his/her partner(s) were already aware of having a C trachomatis infection. This reason was stated by 30.5% of the index women and by 32.8% of the index men (table 2).

Effectiveness of testing male partners to infected index women
The proportion of index women with one or more partner tested was 26.1% in the home sampling group and 11.8% in the office group (p<0.001) (table 3). The relative effectiveness of offering home sampling to male partners compared with office sampling was thus 2.2 (26.1%/11.8%) (95% CI: 1.7 to 2.8). The proportion of all index women with at least one partner identified as infected with C trachomatis was 10.8% in the home sampling group and 6.6% in the office group (p = 0.01), a relative effectiveness of 1.63 (10.8%/ 6.6%) (95% CI: 1.1 to 2.3).

Effectiveness of testing female partners to infected index men
Among randomised index men, the proportion of one or more partners tested was 14.8% in the home sampling group and 3.3% in the office group (p<0.001) (table 3). The relative effectiveness of offering home sampling to female partners compared to office sampling was thus 4.4 (14.8%/3.3%) (95% CI: 2.2 to 9.0). The proportion of index men with at least one partner identified as infected with C trachomatis was 6.2% in the home sampling group and 1.9% in the “in office” group (p = 0.02), a relative effectiveness of 3.3 (6.2%/1.9%) (95% CI: 1.2 to 9.0).

Efficacy of partner testing
Table 4 shows the proportion who had at least one partner tested when the population was restricted to only index men and women who were actually recruited to the study.

DISCUSSION
More partners exposed to C trachomatis infection are tested and more infections are detected when partners are given the opportunity of home sampling compared with office sampling. Based on data from a previous study, 95% of the infected partners received treatment. The major limitation of our study is the rather high proportion of randomised individuals who did not submit questionnaires to the study centre and kits to their partners. We consider that many general practitioners did not provide information about the study as they were not paid or given any incentives for this extra work with the index case. The proportion of recruited men and women were higher in the home sampling group. Although the partner kits were blinded, it could, however, not be excluded that the index patients opened the partner package before they decided to forward it to their partners. Furthermore, partners might have been tested outside the study. We know that contact slips are often not brought by partners when they come for testing but partners in the office sampling group were requested to bring the specimen collection kit that allowed non-invasive testing, which otherwise would not have been the case.

The prevalence of C trachomatis infection in partners to infected patients varies between 27–68%.17 20 21 Treating the partners through patient delivered therapy rather than testing the partners first could therefore seem reasonable.22 23 However, if the partner has a positive test, he could serve as a new index case from where a new partner tracing activity can begin. In particular, this is important when treating patients in sexual networks with no regular partner. According to this, patient delivered therapy has been shown to be of highest efficacy in index patients with regular partners.24 Schillinger et al25 conducted a comparative study assessing reinfection rates after 4 months in index women assigned to patient referral or patient delivered therapy. The two groups had a reinfection rate of 15% and 12%, respectively. The modest absolute reduction of 3% might be explained by the fact that although patient delivered therapy is well accepted22 sexual activity may be more complex and not restricted to one partner only. In fact multi-partnership has increased in the United Kingdom over the past 10 years.26 From a public health point of view, in which a reduction of disease prevalence is warranted, we therefore argue that testing first is important in order to identify new index cases. In this study we have shown that the test rate can be increased twofold to fourfold by offering partners home sampling in stead of office sampling. Assessing only index patients who actually took part in the study (recruited index patients), 41% of index women and 13% of index men had at least one partner tested. This corresponds well with the partner notification rates found in other studies.13 The partner notification rates were increased by home sampling, but the rates might have been even higher if we had used provider referral instead of self referral. This is
so because Katz et al.\(^2\) found that provider referral instead of self-referral increased the effectiveness from 0.18 to 0.72 notifications per index patient. It might therefore be reasonable to speculate that a combination of home sampling and provider referral would be beneficial. This needs to be addressed in subsequent studies.

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**CONTRIBUTORS**

LØ contributed to study design, study logistics, data analyses and wrote the paper; BA and FO contributed to study design, clinical coordination, data analyses and edited the paper; JKM contributed to study design, clinical coordination, data collection, data storage, data analyses and edited the paper; AW contributed to study design, clinical coordination, data collection, data analyses and edited the paper.

**REFERENCES**


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**Table 2** Reasons for dropping out among index patients

<table>
<thead>
<tr>
<th>Reasons</th>
<th>Women</th>
<th></th>
<th></th>
<th>Men</th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No</td>
<td>%</td>
<td></td>
<td>No</td>
<td>%</td>
<td></td>
</tr>
<tr>
<td>My partner(s) is/are already aware of a C. trachomatis infection and has/have taken action regarding diagnosis and treatment outside this study</td>
<td>36</td>
<td>30.5</td>
<td>20</td>
<td>32.8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>I found it unpleasant to contact my partner(s)</td>
<td>19</td>
<td>16.1</td>
<td>9</td>
<td>14.8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unknown identity of partner(s)</td>
<td>12</td>
<td>10.2</td>
<td>6</td>
<td>9.8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>The study interferes too much with privacy/personal integrity</td>
<td>11</td>
<td>9.3</td>
<td>5</td>
<td>8.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Participation too complicated and time consuming</td>
<td>7</td>
<td>5.9</td>
<td>6</td>
<td>9.8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No partner(s) through the past 12 months</td>
<td>3</td>
<td>2.5</td>
<td>1</td>
<td>1.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No particular reason stated</td>
<td>30</td>
<td>25.4</td>
<td>14</td>
<td>23.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>118</td>
<td>100</td>
<td>61</td>
<td>100</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 3** Effectiveness of office sampling and home sampling for testing of partners. Calculations based on the ‘‘intention to test’’ population

<table>
<thead>
<tr>
<th></th>
<th>Index women</th>
<th></th>
<th></th>
<th>Index men</th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Office sampling (n = 663)</td>
<td>Home sampling (n = 637)</td>
<td>p Value</td>
<td>Office sampling (n = 269)</td>
<td>Home sampling (n = 257)</td>
<td>p Value</td>
</tr>
<tr>
<td>No of index cases with one or more partner(s) tested</td>
<td>78</td>
<td>166</td>
<td>&lt;0.0001</td>
<td>9</td>
<td>38</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Proportion of identified cases</td>
<td>11.8%</td>
<td>26.1%</td>
<td></td>
<td>14.3% (10.1 to 18.5)</td>
<td>3.3%</td>
<td></td>
</tr>
<tr>
<td>No of index cases with one or more partner(s) infected</td>
<td>44</td>
<td>69</td>
<td></td>
<td>14.8%</td>
<td>16</td>
<td></td>
</tr>
<tr>
<td>Proportion of identified cases</td>
<td>6.6%</td>
<td>10.8%</td>
<td></td>
<td>4.2% (1.1 to 7.3)</td>
<td>1.9%</td>
<td></td>
</tr>
<tr>
<td>Average number of tested partner(s) per index patient</td>
<td>0.14</td>
<td>0.31</td>
<td></td>
<td>0.04</td>
<td>0.16</td>
<td></td>
</tr>
</tbody>
</table>

**Table 4** The proportion of recruited index men and index women who had at least one partner tested

<table>
<thead>
<tr>
<th></th>
<th>Office sampling of partners</th>
<th>Home sampling of partners</th>
</tr>
</thead>
<tbody>
<tr>
<td>Participating index women</td>
<td>41% (78/191)</td>
<td>74% (166/223)</td>
</tr>
<tr>
<td>Participating index men</td>
<td>13% (9/67)</td>
<td>47% (38/81)</td>
</tr>
</tbody>
</table>
Young injecting drug users have a high incidence of hepatitis C

The detected incidence of hepatitis C among injecting drug users in Dublin, Ireland is higher than corresponding figures from surveys in similar settings elsewhere. It contrasts with a comparatively low incidence of HIV.

Between 1992 and 1998, attenders at a drug treatment centre were encouraged to consent to testing for evidence of hepatitis C infection. The incidence of infection was calculated using the person years method. The date of a first negative test represented the starting point for all patients when calculating their person years at risk. The end point was the date of the last negative test if they remained seronegative or the midpoint of the last negative and first positive test for those who seroconverted.

Sixty seven of 100 patients retested seroconverted during the study period—an overall incidence of 66/100 person years (CI 51–84). A history of imprisonment was associated with a higher incidence while the group who usually smoked heroin and only occasionally injected had a lower incidence. Repeat testing for HIV in 164 patients uncovered two who seroconverted. The incidence of HIV was 0.7/100 person years (CI 0.1–2.5).

The authors considered the high hepatitis C incidence related to the fact that the cohort was young (median age 20.5y) and were mostly recent injectors (median six months). The reason for the extra risk posed by imprisonment is unclear: it predated entry to the study so could not have been causal but might reflect a lifestyle that involves generally increased risk taking behaviour.

Strategies to halt the spread of hepatitis C need to target drug users who have recently started injecting as well as to deflect heroin smokers from moving on to injecting.

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