Is *Mycoplasma hominis* a vaginal pathogen?

O P Arya, C Y W Tong, C A Hart, B C Pratt, Stella Hughes, Pamela Roberts, Patricia Kirby, Jean Howel, Anne McCormick, A D Goddard

**Objective:** To evaluate the role of *Mycoplasma hominis* as a vaginal pathogen.

**Design:** Prospective study comprising detailed history, clinical examination, sexually transmitted infection (STI) and bacterial vaginosis screen, vaginal swabs for mycoplasmas and other organisms, follow up of bacterial vaginosis patients, and analysis of results using SPSS package.

**Setting:** Genitourinary medicine clinic, Royal Liverpool University Hospital.

**Participants:** 1200 consecutive unselected new patients who had not received an antimicrobial in the preceding 3 weeks, and seen by the principal author, between June 1987 and May 1995.

**Main outcome measures:** Relation of *M hominis* isolation rate and colony count to: (a) vaginal symptoms and with the number of polymorphonuclear leucocytes (PMN) per high power field in the Gram stained vaginal smear in patients with a single condition—that is, candidiasis, bacterial vaginosis, genital warts, chlamydial infection, or trichomoniasis, as well as in patients with no genital infection; (b) epidemiological characteristics of bacterial vaginosis.

**Results:** 1568 diagnoses were made (the numbers with single condition are in parenthesis). These included 291 (154) cases of candidiasis, 208 (123) cases of bacterial vaginosis, 240 (93) with genital warts, 140 (42) chlamydial infections, 54 (29) cases of trichomoniasis, and 249 women with no condition requiring treatment. *M hominis* was found in the vagina in 341 women, but its isolation rates and colony counts among those with symptoms were not significantly different from those without symptoms in the single condition categories. There was no association between *M hominis* and the number of PMN in Gram stained vaginal smears whether *M hominis* was present alone or in combination with another single condition. *M hominis* had no impact on epidemiological characteristics of bacterial vaginosis.

**Conclusion:** This study shows no evidence that *M hominis* is a vaginal pathogen in adults.

(Sex Transm Inf 2001;77:58–62)

Keywords: bacterial vaginosis; vaginal pathogens; *Mycoplasma hominis*

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**Introduction**

*Mycoplasma hominis* and *Ureaplasma urealyticum* are the most frequently isolated mycoplasmas from the human genital tract. They are ubiquitous resulting in colonisation of the genitalia by sexual contact. Taylor-Robinson and McCormack reviewed the genital mycoplasmas in 1980.

A more recent update described the putative roles of these and other species, notably *M fermentans*, *M genitalium*, and *M penetrans*.

*M hominis* has been linked to pelvic inflammatory disease and preterm labour. Although frequently isolated in association with bacterial vaginosis, chlamydial and gonorrhoeal infections, and trichomoniasis, its exact contribution in these conditions remains unclear. On its own it is thought to behave as a commensal. Nevertheless, *M hominis* has been isolated in pure culture from cases of vaginitis in children.

We sought to investigate its role by studying women with the above infections as well as women with genital warts and candidiasis comparing those with (including quantitative analysis) and without *M hominis*. Symptoms, notably vaginal discharge with or without irritation, and an inflammatory response indicated by the number of polymorphonuclear leucocytes (PMN) per high power field (hpf) were noted. To exclude any potential effect of concurrent infection(s), we analysed women with a single condition, as well as women not found to have any condition requiring treatment, termed “normal” in this paper. We also investigated the contribution of *M hominis*, if any, to the epidemiology and aetiology of bacterial vaginosis.

**Participants and methods**

The local hospital ethics committee’s approval was obtained. The study population comprised all new unselected female patients, who had not received any topical medication or a systemic antimicrobial in the preceding 3 weeks and were seen by one observer (OPA) at the department of genitourinary medicine, the Royal Liverpool University Hospital, between June 1987 and May 1995. After obtaining a detailed history the patients underwent clinical, including genital and pelvic examination. The appearance of the vulva, vagina, cervix, the quality and quantity of discharge, and other physical signs were noted.

**Laboratory methods**

**CANDIDIASIS**

Gram stained vaginal smears were examined for yeast cells and pseudohyphae in the clinic. Swabs were sent in Feinberg–Whittington medium to the laboratory for culture.

**BACTERIAL VAGINOSIS**

The tests included vaginal pH, amine test, wet film, and Gram stained smears for clue cells and other morphotypes. Swabs were cultured aerobically and anaerobically on blood, choco-
late, and MacConkey agars, and on Gardnerella selective medium (Mobiluncus was not specifically sought). The wet films and Gram stained smears were examined in the clinic by technicians blinded to the clinical details. Bacterial vaginosis was diagnosed in the presence of vaginal pH > 4.5, a positive amine test, paucity or absence of lactobacilli, and the presence of clue cells. All these patients had a Nugent score^3 (applied retrospectively up to 1991 and thereafter prospectively) of >6.

**CHlamydial Infection**

Swabs were taken from the endocervix (and other sites as indicated) and sent in transport media to the hospital laboratory where they were processed using enzyme linked immunosorbent assay (ELISA, AntigEnz Chlamydia, Shiel diagnostic) and immunofluorescence techniques (DIF, Syva Microtrak, Syva).

**Gonococcal Infection**

Gram stained smears from the endocervix and other sites as indicated were examined in the clinic and swabs were inoculated onto modified Thayer–Martin medium (Oxoid, Basingstoke, UK) and cultured at 37°C in carbon dioxide in humidified air.

**Trichomoniasis**

Wet smears from the posterior vaginal fornix were examined in the clinic by microscopy. Swabs were sent in Feinberg–Whittington medium to the laboratory for culture.

**Mycoplasmas**

Vaginal and endocervical swabs were sent to the laboratory in A3×B transport medium.7 These were inoculated into PPLO broths (Difco) in serial 10-fold dilutions. *Urealyticum* broth contained 1% (w/v) urea at pH 6.0 and *M. hominis* broth contained 1% (w/v) arginine dihydrochloride at pH 7.4. Both contained 0.002% (w/v) phenol red indicator.8 Broths showing colour change after incubation at 37°C for 4–5 days were subcultured onto A7 mycoplasma agar^9 for confirmation. The amount of *M. hominis* in a sample was determined by end point titration in broth and expressed as number of colony forming units (CFU) per ml. A CFU of ≥ 5 ×10^5/ml indicated the presence of the organism in large numbers.

**Statistical Analysis**

The results were analysed using the SPSS package (Statistical Package for Social Sciences). The differences between the various categories were assessed using the χ^2 test or Fisher’s exact test (if cell values less than 5) using EPI-INFO version 6 (CDC, Atlanta, GA, USA).

**Results**

A total of 1200 patients were enrolled. A total of 1568 diagnoses were made (numbers with single condition in parentheses), including 291 (154) of candidiasis; 208 (123) of bacterial vaginosis (BV); 240 (93) of genital warts; 140 (42) chlamydial infections; 54 (29) with trichomoniasis (TV), and 249 “normal” women with no condition requiring treatment. *M. hominis* was found in the vagina in 341 women.

### M. hominis Isolation Rates

*M. hominis* isolation rates and of those with high numbers (≥ 5 ×10^5) among women with single conditions, and of the “normal” category, and all women are shown in table 1. *M. hominis* isolation rates were highest among those with BV (59%) followed by those in the chlamydia category (31%), warts (20%), the normal category (12%), and candidiasis (8%). The differences between TV and normal categories, and BV and normal categories were highly significant (p<0.0001). The difference between those with chlamydial infection and the normal category was also significant (p=0.0041). Women with TV who also had *M. hominis* had the highest rate (76%) of *M. hominis* high counts followed by those with BV (48%), compared with the normal category (23%). Using high colony counts alone, only the differences between TV and normal categories and between BV and normal categories remained significant (p, respectively, 0.0009 and 0.0283).

**Vaginal Discharge**

Table 2 shows the numbers and proportions of women who complained of, or admitted to experiencing, vaginal discharge with or without vaginal or vulval irritation in the various categories. In all categories, except candida, more of *M. hominis* positive women had experienced vaginal discharge than the *M. hominis* negative women. However, the difference was statistically significant only between the overall totals within the whole populations studied. Thus 36% (312 of 859) of *M. hominis* negative women had symptoms of vaginal discharge compared with 58% (199 of 341) of *M. hominis* positive (p<0.00001). The presence of *M. hominis* in large numbers (≥ 5 × 10^5) did not seem to have any additional effect.

**Inflammatory Response in the Vagina**

Table 2 shows the “inflammatory response” as indicated by the presence of more than 20 PMN per hpf in the Gram stained vaginal smears in the various diagnostic categories. No significant differences between *M. hominis* positive and *M. hominis* negative women were detected in all “single condition” categories. However, when all women were considered, 21% (179 of 859) of *M. hominis* negative women had inflammatory response compared

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**Table 1** *M. hominis* isolation rates in the various diagnostic categories

<table>
<thead>
<tr>
<th>Category</th>
<th>No with single conditions</th>
<th>No with <em>M. hominis</em> (%)</th>
<th>No with <em>M. hominis</em> count ≥ 5 ×10^5 (%) of those with <em>M. hominis</em> (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Candidiasis</td>
<td>154</td>
<td>12 (8%)</td>
<td>5 (42%)</td>
</tr>
<tr>
<td>Bacterial vaginosis</td>
<td>123</td>
<td>73 (59%)</td>
<td>35 (28%)</td>
</tr>
<tr>
<td>Genital warts</td>
<td>93</td>
<td>19 (20%)</td>
<td>8 (42%)</td>
</tr>
<tr>
<td>Chlamydial infection</td>
<td>42</td>
<td>13 (31%)</td>
<td>6 (46%)</td>
</tr>
<tr>
<td>Trichomoniasis</td>
<td>29</td>
<td>17 (59%)</td>
<td>13 (76%)</td>
</tr>
<tr>
<td>Normal</td>
<td>249</td>
<td>31 (12%)</td>
<td>7 (23%)</td>
</tr>
<tr>
<td>All women (1200)</td>
<td>249</td>
<td>341 (28%)</td>
<td>156 (46%)</td>
</tr>
</tbody>
</table>


with 27% (93 of 341) in M hominis positive women; this difference was statistically significant (p = 0.02).

We also looked at the vaginal inflammatory response as defined above in the various diagnostic categories in the presence of vaginal discharge as a symptom; the differences between those who were M hominis positive and those M hominis negative within the various categories were very small and not significant (table 2).

### M HOMINIS AND BACTERIAL VAGINOSIS

#### Epidemiology

Table 3 shows the epidemiological characteristics of the BV and the “normal women” categories (because of the aetiological association of Gardnerella vaginalis and strict anaerobes with BV, women with these organisms in vaginal swab cultures have been excluded from the normal women category). Only one characteristic emerged to be noteworthy—that is, the use of an intrauterine contraceptive device (IUCD); 14 (24%) of the 58 M hominis positive women were using IUCD compared with only two (5%) of the 40 M hominis negative women in the BV category. This difference was significant (p = 0.025). The significance level in this respect was higher when M hominis negative women (two of 40) were compared with those with high counts of M hominis (10 of 27) (p = 0.002). However, no such difference was observed in the normal women category. Indeed, none of the 12 M hominis positive women in the normal women category was using IUCD suggesting that the difference noted above in the BV category was not due to M hominis alone. The differences between M hominis negative and M hominis positive women with regard to all the other factors in the normal women category were also not significant.

#### Aetiology

Table 4 compares patients in the BV category with the normal women category of women with regard to the associations between the organisms (that is, M hominis, G vaginalis, and strict anaerobes) and BV. Although M hominis alone was isolated more often from women in the normal women category (14 of 249, 6%) than from those in the BV category (four of 123, 3%), this difference was not significant (p = 0.4559). Two of the four women with M hominis alone in the BV category, compared with two of the 14 women in the normal women category had M hominis colony counts of $\geq 5 \times 10^5$; this difference was not significant (p = 0.1970). Of the 86 women harbouring G vaginalis and/or strict anaerobes as well as M hominis, 69 (80%) had BV compared with 46 (73%) of 63 women with G vaginalis and/or strict anaerobes but without M hominis. The difference was, however, not significant (p = 0.4012), suggesting that the additional presence of M hominis with G vaginalis and strict anaerobes did not seem to increase the likelihood of the patient developing BV.

### Table 2: Relation of M hominis with vaginal discharge and vaginal inflammatory response in the various categories

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Candida (N=154)</th>
<th>M hominis</th>
<th>Bact vag (N=123)</th>
<th>M hominis</th>
<th>Warts (N=93)</th>
<th>M hominis</th>
<th>Chlamydia (N=42)</th>
<th>M hominis</th>
<th>Normal (N=249)</th>
<th>M hominis</th>
<th>All women (1200)</th>
<th>M hominis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Symptom: Vaginal discharge</td>
<td>45%</td>
<td>42%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
<td>45%</td>
</tr>
<tr>
<td>Vaginitis: &gt;20 PMN per high power field in Gram-stained vaginal smear</td>
<td>22%</td>
<td>20%</td>
<td>31%</td>
<td>24%</td>
<td>46%</td>
<td>19%</td>
<td>20%</td>
<td>20%</td>
<td>20%</td>
<td>20%</td>
<td>31%</td>
<td>20%</td>
</tr>
</tbody>
</table>

In women with symptom of vaginal discharge

<table>
<thead>
<tr>
<th>M hominis positive</th>
<th>18/64</th>
<th>1/5</th>
<th>0</th>
<th>10/44</th>
<th>16/67</th>
<th>5/32</th>
<th>1/4</th>
<th>0/2</th>
<th>0/1</th>
<th>5/9</th>
<th>3/6</th>
<th>0/2</th>
<th>6/8</th>
<th>10/13</th>
<th>9/9</th>
<th>15/79</th>
<th>3/14</th>
<th>0/3</th>
<th>89/312</th>
<th>64/199</th>
<th>26/89</th>
</tr>
</thead>
<tbody>
<tr>
<td>M hominis negative</td>
<td>28%</td>
<td>20%</td>
<td>23%</td>
<td>24%</td>
<td>16%</td>
<td>25%</td>
<td>56%</td>
<td>50%</td>
<td>75%</td>
<td>77%</td>
<td>100%</td>
<td>19%</td>
<td>21%</td>
<td>28%</td>
<td>32%</td>
<td>36%</td>
<td>58%</td>
<td>57%</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Arya, Tong, Hart, et al.


Table 3  Epidemiological characteristics of bacterial vaginosis and “normal women” categories

<table>
<thead>
<tr>
<th>Epidemiological characteristic</th>
<th>Bacterial vaginosis (123) M hominis</th>
<th>“Normal women” (215) * M hominis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Neg (50)</td>
<td>Pos (73)</td>
</tr>
<tr>
<td>Mean age</td>
<td>27.20</td>
<td>27.97</td>
</tr>
<tr>
<td>Mean age first sex</td>
<td>17.26</td>
<td>17.59</td>
</tr>
<tr>
<td>Marital status:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single</td>
<td>30 (60%)</td>
<td>46 (63%)</td>
</tr>
<tr>
<td>Currently married</td>
<td>10 (20%)</td>
<td>12 (16%)</td>
</tr>
<tr>
<td>Divorced/widowed/separated</td>
<td>10 (20%)</td>
<td>15 (20%)</td>
</tr>
<tr>
<td>Last sex in past 2 weeks</td>
<td>34 (68%)</td>
<td>45 (69%)</td>
</tr>
<tr>
<td>Positive history of cunnilingus†</td>
<td>27/38 (71%)</td>
<td>27/45 (60%)</td>
</tr>
<tr>
<td>Never pregnant</td>
<td>24 (48%)</td>
<td>24 (48%)</td>
</tr>
<tr>
<td>More than 1 sex partner past 12 months</td>
<td>14 (28%)</td>
<td>17 (23%)</td>
</tr>
<tr>
<td>Last menstrual period‡:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;9 days ago</td>
<td>7/47 (15%)</td>
<td>13/66 (20%)</td>
</tr>
<tr>
<td>10–19 days ago</td>
<td>22/47 (47%)</td>
<td>32/66 (48%)</td>
</tr>
<tr>
<td>&gt;19 days ago</td>
<td>18/47 (38%)</td>
<td>21/66 (32%)</td>
</tr>
<tr>
<td>Contraception§:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hormone</td>
<td>19/40 (47%)</td>
<td>21/58 (36%)</td>
</tr>
<tr>
<td>IUCD</td>
<td>2/50 (4%)</td>
<td>14/58 (24%)</td>
</tr>
<tr>
<td>None</td>
<td>7/40 (17%)</td>
<td>10/58 (17%)</td>
</tr>
<tr>
<td>Sometimes</td>
<td>4 (8%)</td>
<td>7 (10%)</td>
</tr>
<tr>
<td>Always</td>
<td>2 (4%)</td>
<td>5 (7%)</td>
</tr>
<tr>
<td>Condom</td>
<td>25 (50%)</td>
<td>42 (52%)</td>
</tr>
<tr>
<td>Partner circumcised‡:</td>
<td>3/7 (8%)</td>
<td>3/4 (7%)</td>
</tr>
</tbody>
</table>

*Women with G vaginalis/anaerobes in vaginal swab cultures have been excluded.
†Information not available for some patients; hence denominator shown.
‡Pregnant, hysterectomised, and menopausal women have been excluded.
§Pregnant, hysterectomised, and menopausal women and those with vasectomised husbands have been excluded.

Table 4  Association of M hominis, G vaginalis, and anaerobes with bacterial vaginosis

<table>
<thead>
<tr>
<th>Organisms</th>
<th>BV category</th>
<th>“Normal women” category</th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>4 (3%)</td>
<td>201 (81%)</td>
</tr>
<tr>
<td>M hominis only</td>
<td>4 (3%)</td>
<td>14 (6%)</td>
</tr>
<tr>
<td>G vaginalis only</td>
<td>4 (3%)</td>
<td>6 (2%)</td>
</tr>
<tr>
<td>Anaerobes only</td>
<td>4 (3%)</td>
<td>3 (1%)</td>
</tr>
<tr>
<td>G vaginalis and anaerobes</td>
<td>38 (31%)</td>
<td>8 (3%)</td>
</tr>
<tr>
<td>M hominis and G vaginalis</td>
<td>4 (3%)</td>
<td>6 (2%)</td>
</tr>
<tr>
<td>M hominis and anaerobes</td>
<td>2 (2%)</td>
<td>1 (1%)</td>
</tr>
<tr>
<td>G vaginalis, anaerobes, and M hominis</td>
<td>63 (51%)</td>
<td>9 (4%)</td>
</tr>
</tbody>
</table>

Discussion

To our knowledge this is the first large prospective study to investigate the role of M hominis as a vaginal pathogen in adults. One of us (OPA) undertook all the clinical examinations and sample collections thus avoiding the problems of interobserver variation. However, we acknowledge that this study has certain limitations. Firstly, the conventional diagnostic methods used in this study (as opposed to those increasingly being used now encompassing DNA technology, particularly for chlamydial infection) would not with certainty have excluded some of the concurrent infections. This may, possibly, be at least partly responsible for our finding of inflammatory response in a fifth of our patients with BV. Nevertheless, this potential bias would apply equally to all of the diagnostic categories. Secondly, having had to exclude the concurrent infections to make meaningful comparisons, the resulting number became rather small. This we deem, however, to be the correct approach in view of the significant differences found between M hominis positive and M hominis negative women among the total population, suggesting the potential role of concurrent infection(s). This reinforces the importance of taking steps to eliminate such potential influences when assessing the role of M hominis.

The above limitations notwithstanding, this study confirms the frequent isolation of M hominis in bacterial vaginosis, chlamydial infection, and trichomoniasis. The low rate of M hominis carriage in women with candidiasis has been observed previously, and is partly due to the low pH (<4.5) as well as low rates of anaerobes in women with candidiasis, both these features being unfavourable to M hominis. Previous, however, a study using a therapeutic approach to assess the cause of non-specific vaginitis (now termed bacterial vaginosis) had shown that, whereas eradication of G vaginalis (when patients treated with metronidazole) cleared vaginitis, eradication of M hominis alone (when patients treated with doxycycline) did not, raising doubts over the role of M hominis in BV.

In addition, M hominis did not seem to make any contribution to the epidemiology of bacterial vaginosis. In this context, the exclusion of those harbouring G vaginalis and strict anaerobes in the vagina from the normal women category (table 3) may be debatable. We have already given a reason for this to which may be added the possibility that vaginal colonisation with M hominis may in part be dependent on vaginal anaerobes. In a recent study M hominis was reported to be more strongly associated with bacterial vaginosis than G vaginalis. Our findings do not support this. That study, however, did not take the other anaer-
obes into account; nor is it clear as to whether
or not the concurrent infections were excluded.
Taylor-Robinson and McCormack1 surmised
*M. hominis* may act either in symbiosis with
other organisms or as a sole pathogen in bacte-
rrial vaginosis. In our study *M. hominis* did not
display an active role in either capacity.

Conflict of interest: None

Some preliminary results, while the study was still in progress,
were presented at the Spring Meeting of the Medical Society for
the Study of Venerable Diseases held in Liverpool in May 1994.
We are indebted to the staff of the department of
genitourinary medicine and medical microbiology, Royal Liver-
pool University Hospital for their kind cooperation at all times.
We thank M Blake who carried out the mycoplasma laboratory
work, Dr. L Cuevas of the Liverpool School of Tropical
Medicine for statistical advice, and Clare Kelly and Lynda Jones
for typing the manuscript.

Contributors: OPA had the original idea for the study and
designed the protocol, recruited and examined patients and
completed proformas, collected and interpreted data, carried
out and completed statistical analysis, and was the principal
author of the paper; BCP (deceased) participated in the proto-
col design and supervised the laboratory work on chlamydia and
mycoplasma, taken over later by CYWT, who, in addition, par-
ticipated in the interpretation of data and review of the
manuscript; CAH supervised the laboratory work on the
remaining aspects of bacteriology and participated in the
interpretation of data and manuscript reviews; SH, PR, PK, and
JH carried out all the laboratory work in the genitourinary
medicine department clinic; AMcC and ADG processed and
extracted the data in a form that could be easily analysed.

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*Sex Transm Infect* 2001 77: 58-62
doi: 10.1136/sti.77.1.58

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