Urogenital Mycoplasmosis and Pregnancy

Põder A and Haldre M
Tartu University Clinics Foundation, Clinical Research Center, Estonia Tartu Sõdra-54, Estonia

Abstract

Urogenital mycoplasmas Ureaplasma urealyticum, Ureaplasma parvum and Mycoplasma hominis have long been considered commensals of human urogenital microflora. Several studies have recently clarified whether these bacteria could cause or be associated with various obstetric complications. Epidemiologic studies have found high prevalence of Ureaplasma spp. and M. hominis among otherwise healthy population. On the other hand, U. urealyticum, U. parvum and M. hominis have increasingly been linked to adverse pregnancy outcomes such as spontaneous pre-term labor, pre-term premature rupture of fetal membranes, miscarriage, stillbirth and low birth weight. However, no convincing causal relationship has been shown. Studies have shown that antibiotics resistance is on the rise and differs considerably by regions. Screening of asymptomatic patients for M. hominis, U. urealyticum and U. parvum is currently still not indicated.

Keywords: Mycoplasmas; Pregnancy; M. hominis; Ureaplasmas

Introduction

The name mycoplasma has been used as the general name for the class Mollicutes the smallest free-living microorganisms. In the urogenital tract, the relevant Mollicutes are Mycoplasma genitalium and Mycoplasma hominis and two species of ureaplasmas- Ureaplasma urealyticum and Ureaplasma parvum. This article aims to give a short overview of associations and causative roles of aforementioned mycoplasmas with various obstetric complications. Brief summary is provided about recent trends in treatment.

Epidemiology

Ureaplasmas and M. hominis are considered opportunistic pathogens because they can be isolated from the lower urogenital tract of healthy women as well as from individuals with disease. Table 1 summarizes the prevalence of urogenital mycoplasmas as found by different researchers.
Table 1: Prevalence of urogenital mycoplasmas.

**Ureaplasma spp.**

Ureaplasma genital tract colonization has been associated with adverse pregnancy outcomes such as spontaneous abortion [10], premature rupture of membranes (PROM) [4], premature delivery [11], neonatal morbidity and perinatal death [12]. The methodology and samples vary considerably between studies, which makes combining and synthesizing the result difficult. Therefore, the results are presented as stated by studies in a Table 2.

<table>
<thead>
<tr>
<th>Study</th>
<th>Sample size</th>
<th>Population</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ouzounova-Raykova et al.</td>
<td>15%</td>
<td>Women in Bulgaria</td>
<td></td>
</tr>
<tr>
<td>Tibaldi et al.</td>
<td>17%</td>
<td>Non-pregnant women</td>
<td></td>
</tr>
<tr>
<td>Zdrodowska-Stefanow et al.</td>
<td>23%</td>
<td>Women aged 18-55, gynaecological and STD outpatient clinics</td>
<td></td>
</tr>
<tr>
<td>Naaber et al.</td>
<td>7.7% Uu*</td>
<td>32%</td>
<td></td>
</tr>
<tr>
<td>Ye et al.</td>
<td>53%</td>
<td>Women in Hangzhou</td>
<td></td>
</tr>
<tr>
<td>Redelinghuys et al.</td>
<td>76%</td>
<td>Pregnant South-African women</td>
<td></td>
</tr>
<tr>
<td>Kataoka et al.</td>
<td>8.7% Uu*</td>
<td>52%</td>
<td></td>
</tr>
</tbody>
</table>

*Uu-Ureaplasma urealyticum

Table 2: Results of studies on association of ureaplasmas with obstetric complications.

Witt et al. [13] found that the amniotic cavity of 44% of patients with PROM and therapy-resistant preterm labor was colonized with *U. urealyticum* compared to only 2.7% in control group of pregnant women. Mitsunari et al. [11] findings support the association between cervical *U. urealyticum* and preterm delivery (controls colonisation 46% compared to 87% in preterm delivery patients).

Very few studies distinguish between *U. urealyticum* and *U. parvum* and describe the influence of *Ureaplasma spp.* combined. Even if an
article states that *U. ureaplasma* is studied, then careful examination of methods section reveals that the method used does not allow to differentiate between biovars. Nevertheless, some researchers have made the distinction. Kataoka et al. [14] found that *U. parvum* but not *U. urealyticum* is associated with late abortion or preterm birth. Kasprzykowska et al. [15] found that colonization of the lower urogenital tract with *U. parvum* can cause asymptomatic infection of the upper reproductive system. Kasper et al. [16] found that the increased amount of *U. parvum* was significantly linked to histological choioamnionitis, PROM together with preterm labor, early-onset sepsis, and bronchopulmonary dysplasia. Furthermore, no significant difference between the bacterial load of *U. urealyticum* and neonatal outcome was observed [16]. The importance of bacterial load was found by Abele-Horn et al. [17]. The study revealed that high colonization (>10^5 cfu/ml) level with *U. urealyticum* was associated with a significant increase of clinical choioamnionitis, PROM and preterm delivery. Low density vaginal colonization levels had no effect on clinical choioamnionitis and preterm delivery [17].

Greenow et al. [18] studied whether treatment of pregnant women heavily colonized with ureaplasma with erythromycin could have an impact on birth weight, but found no statistically significant difference [18]. However, in this study the antibiotics treatment was started only between 22-32th week of gestation, whereas obstetric adverse outcomes, such as abortion may arise much earlier.

**M. hominis**

Evidence has accumulated that *M. hominis* may be of significance in the condition of bacterial vaginosis. Bacterial vaginosis (BV) in turn has been associated with subsequent early pregnancy loss [19]. Donders et al. [19] studied BV and found that *M. hominis*, and *U. urealyticum* were associated with an increased risk of clinical choioamnionitis, PROM and preterm delivery. *M. hominis* may act symbiotically with other BV-associated bacteria or as the sole pathogen based on the observation that this mycoplasma can be found in large numbers in the vagina of most women with BV, but less often in healthy women [20].

**M. genitalium**

*Mycoplasma genitalium* is a globally important sexually transmitted pathogen known to cause urethritis [21]. However, there is no conclusive evidence supporting its role in adverse obstetric outcomes and tubal infertility. Data suggests an association between cervicitis and *M. genitalium*. Supporting evidence of causal relationship between pelvic inflammatory disease and *M. genitalium* is moderate to strong [21].

Existing data provide some support for the hypothesis that *M. genitalium* can cause female infertility, but are inconclusive [22]. A few studies have found an independent association between vaginal presence of *M. genitalium* and preterm delivery [23,24]. However, causative association of *M. genitalium* with adverse obstetric outcomes remains unestablished [25].

**Diagnostics**

Ureaplasmas and *M. hominis* are considered to be a part of natural microflora among healthy women and men. Consequently, the detection of these bacteria has a low positive predictive value in relation to diseases. Therefore, currently, most reviews do not recommend the inclusion of these pathogens into the routine sexually transmitted infections (STI) screening protocols and state that asymptomatic individuals should not be screened with culture or nucleic acid amplification tests (NAAT).

As previously discussed, detection of *M. genitalium* is strongly correlated to diseases. As it is detected only rarely in healthy individuals, the positive predictive value of the test for *M. genitalium* is high. It has been suggested that *M. genitalium* could be among routinely screened microbes as are *C. trachomatis* and *N. gonorrhoeae*.

**Laboratory Methods**

**Culture and microscopy**

Culture has been previously considered the gold standard in the detection of ureaplasmas but it is a difficult method since these fastidious organisms require the presence of serum, metabolic substrate and growth factors [26]. In addition, some commercial kits available for diagnosis of urogenital mollicute infections have demonstrated lack of diagnostic sensitivity [27]. Lack of a rigid cell wall makes it nearly impossible to directly visualize ureaplasma by light microscopy. The laboratory testing of *M. genitalium* has been particularly difficult as it takes several weeks or even months for each isolate to grow, making culture impossible to use for diagnostics in routine clinical practice [28].

Therefore, owing to the poor and extremely slow growth of the bacterium in culture, diagnosis of *M. genitalium* infection is performed exclusively using nucleic acid amplification tests (NAAT) [28].

**Polymerase chain reaction (PCR)**

PCR has been introduced in the practice because of the possibility to use different clinical materials, quick test results and possibility to detect different pathogens in a swab [7]. PCR is also more sensitive than culture for detection (<100 genome copies) of nonviable as well as viable ureaplasmas. The results of PCR are available in a day, whereas a culture takes 2-5 days. The most commonly used gel-based traditional and real-time PCR protocols target the common multiple-banded antigen (mba), urease or 16s RNA genes. Currently new sensitive and low-cost multiplex PCR methods are being developed to make diagnosis by nucleic acid amplification tests (NAATs) more cost-efficient [29].

However, at the moment there is no gold standard PCR test which to compare other new tests with. When choosing the PCR test for use, it must be accounted for that PCR of some MgPa-related sequences demonstrates lack of diagnostic sensitivity [27]. It has been suggested that *M. genitalium* could be among routinely screened microbes as are *C. trachomatis* and *N. gonorrhoeae*.

**Quantitative-PCR (qPCR)**

Conventional PCR is restricted in the accurate quantification of microorganisms. In contrast, quantitative real-time PCR (qPCR) using fluorescence dyes or probes facilitate the quantification of amplified viral, bacterial, and parasitic products. As it was previously discussed, in some obstetric complications the amount of bacteria is more important than the mere presence of it [16].

PCR is an excellent alternative to culture, but culture allows antibiotic susceptibility testing. However, molecular testing for macrolide and fluoroquinolone resistance mediating mutations is also possible.
Specimen collection

There is still no consensus as to which specimens have the best sensitivity in detection of ureaplasmas and *M. genitalium*. For *M. genitalium* the use of more than one specimen may significantly improve the diagnostic sensitivity. Lillis et al. [30] found that the single best specimen for the detection of *M. genitalium* infection was vaginal swab specimen (74.3%), followed by urine specimen (61.4%), and rectal swab specimen (24.3%).

Vaginal swab combined with endocervical swab specimen provided a sensitivity of 95.7% [30]. Research has also shown that self-collected vaginal swabs are equal to clinician-collected vaginal swabs for diagnosis of *C. trachomatis* and *N. gonorrhoeae*, and there is no reason to doubt that the same would be true for *M. genitalium* [30].

**General Treatment**

Mycoplasmas and ureaplasmas lack a cell wall, the target of beta-lactam antibiotics and vancomycin, which makes them resistant to these antibiotics. Additionally, ureaplasma species have natural resistance to lincosamides (e.g. clindamycin) and *M. hominis* possesses inherent resistance to macrolides, except josamycin [31].

Historically, tetracyclines, macrolides, and quinolones have been the major antibiotics used in the treatment of urogenital infections caused by mycoplasmas. However, their therapeutic efficacy may be unpredictable due to increasing resistance. The extent of resistance varies regionally according to different antimicrobial therapy policies and the history of prior antimicrobial exposure in different populations. Recently, several articles have been published that study the resistance of *M. hominis* and ureaplasma to antibiotics. Table 3 summarizes the results of these studies. De Francesco et al. [32] studied a sample of 9,956 patients in Italy. Both *M. hominis* and *Ureaplasma spp.* were most sensitive to doxycycline and tetracycline, as well as to josamycin.

Clarithromycin and josamycin were the most potent macrolides against ureaplasmas. The only macrolide effective against *M. hominis* was josamycin. Another antibiotic effective against both microbes was pristinamycin. Other two macrolides that were studied-azithromycin and erythromycin - were only moderately effective [32]. Resistance to erythromycin has been reported additionally by Ponyai et al. [31] and Krausse et al. [33] who determined that the resistance of ureaplasmas to erythromycin is 81% and 21%, respectively.

<table>
<thead>
<tr>
<th>Specimen collection</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Francesco et al. (9,956)</td>
<td>Pinyai et al., (2,309)</td>
<td>Krausse et al., (469)</td>
<td>Ye et al. (37,055)</td>
<td>Pignanelly et al. (2,480)</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Ureap.</strong></td>
<td><strong>M. hominis</strong></td>
<td><strong>Ureap.</strong></td>
<td><strong>M. hominis</strong></td>
<td><strong>Ureap.</strong></td>
<td><strong>M. hominis</strong></td>
<td><strong>Ureap.</strong></td>
</tr>
<tr>
<td><strong>Doxycycline</strong></td>
<td>2%</td>
<td>1%</td>
<td>2%</td>
<td>2%</td>
<td>1%</td>
<td>10%</td>
</tr>
<tr>
<td><strong>Tetracycline</strong></td>
<td>2%</td>
<td>1%</td>
<td>4%</td>
<td>12%</td>
<td>3%</td>
<td>11%</td>
</tr>
<tr>
<td><strong>Erythromycin</strong></td>
<td>10%</td>
<td>99%</td>
<td>81%</td>
<td>-</td>
<td>21%</td>
<td>96%</td>
</tr>
<tr>
<td><strong>Clarithromycin</strong></td>
<td>5%</td>
<td>99%</td>
<td>-</td>
<td>-</td>
<td>5%</td>
<td>99%</td>
</tr>
<tr>
<td><strong>Josamycin</strong></td>
<td>1%</td>
<td>1%</td>
<td>-</td>
<td>-</td>
<td>2%</td>
<td>3%</td>
</tr>
<tr>
<td><strong>Azithromycin</strong></td>
<td>5%</td>
<td>99%</td>
<td>10%</td>
<td>-</td>
<td>7%</td>
<td>99%</td>
</tr>
<tr>
<td><strong>Ciprofloxacin</strong></td>
<td>60%</td>
<td>100%</td>
<td>-</td>
<td>-</td>
<td>16%</td>
<td>8%</td>
</tr>
<tr>
<td><strong>Oxofloxacin</strong></td>
<td>5%</td>
<td>100%</td>
<td>25%</td>
<td>5%</td>
<td>2%</td>
<td>2%</td>
</tr>
<tr>
<td><strong>Pristinamycin</strong></td>
<td>1%</td>
<td>1%</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td><strong>Clindamycin</strong></td>
<td>-</td>
<td>-</td>
<td>75%</td>
<td>5%</td>
<td>43%</td>
<td>0%</td>
</tr>
</tbody>
</table>

Table 3: Percentage of resistant strains, sample size in brackets.

Pignanelly et al. [5] studied *M. hominis* and found that the isolates were sensitive to tetracycline and doxycycline but resistant to fluoroquinolones and macrolides except josamycin. Krausse et al. [33] found that doxycycline was the most active tetracycline against ureaplasmas and *M. hominis*. However, 10-13% of *M. hominis* was resistant to tetracyclines [33].

Nevertheless, doxycycline is still the drug of choice for the treatment of urogenital mycoplasma infections and may also be used for coinfection with *M. hominis* [33]. However, resistant strains are increasingly prevalent. In accordance with current guidelines, the International Union against Sexually Transmitted Infections (IUSTI) recommends use of macrolides in case of uncomplicated *M. genitalium* in the absence of macrolide resistancy mediating mutations: azithromycin 500 mg on day one, then 250 mg once daily for days 2-5 orally or josamycin 500 mg 3 times daily for 10 days.

If case of macrolide-resistancy, moxifloxacin 400 mg once daily for 7-10 days may be considered as a treatment option, however it is contraindicated in pregnancy [25,34]. Josamycin has been shown to be *in vitro* effective against *M. genitalium* [35]. This 16-membered macroline is widely used in Italy, Russia, France, Spain, etc.

In recently published study treatment with josamycin was associated with 93.5% eradication rate in male patients with *M. genitalium* infection with josamycin 500 mg three times daily for 10 days [36]. In countries where josamycin is available, it could be considered for treatment in case of *M. genitalium* infection.
Treatment during pregnancy

During pregnancy, the list of antibiotics that are not contraindicated but effective against urogenital mycoplasmas is much shorter. Macrolides and clindamycin are allowed. Erythromycin, the antibiotic pathogen and needs treatment. In accordance with current guidelines, whether to treat the colonization to prevent obstetrics complications question of who should be screened for urogenital mycoplasmas and flora may possess [39,40]. M. genitalium infections can be treated with azithromycin and josamycin, however, resistance may pose a problem.

Conclusion

M. hominis and U. urealyticum may be part of the normal vaginal flora but both are associated with obstetrics complications. The question of who should be screened for urogenital mycoplasmas and whether to treat the colonization to prevent obstetrics complications remains currently unsettled. M. genitalium is a definite urogenital pathogen and needs treatment. In accordance with current guidelines, josamycin and azithromycin are treatment of choice in case of the absence of macrolide resistance associated mutations in M. genitalium. Josamycin is safe to use in pregnancy in case there is a need to treat urogenital mycoplasmosis.

References


