

# Hematoma and abscess formation caused by *Mycoplasma hominis* following cesarean section

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**Abstract:** *Mycoplasma* species cannot be identified by routine bacteriological culture methods and are resistant to common antimicrobial agents. *Mycoplasma hominis* usually colonizes the lower urogenital tract and causes pyelonephritis, pelvic inflammatory disease, chorioamnionitis, rupture of fetal membranes, preterm labor, postpartum fever, postabortal fever, and neonatal infection. This organism is highly prevalent in cervicovaginal cultures of sexually active women. *M. hominis*, *M. genitalis*, *Ureaplasma urealyticum*, and *U. parvum* may invade and infect placental and fetal tissues, leading to adverse pregnancy outcomes. *M. hominis* occasionally causes nongenitourinary infection of the blood, wounds, central nervous system, joints, or respiratory tract. We present a case of a 27-year-old woman who developed abdominal wound hematoma and abscess after cesarean section. The wound was drained, but her high fever persisted, in spite of antibiotic treatment using flomoxef sodium and imipenem-cilastatin sodium. Because the exudate exhibited *M. hominis* growth in an anaerobic environment, we administered the quinolone ciprofloxacin. This therapy resolved her fever, and her white blood cell count and C-reactive protein level diminished to the normal ranges. To our knowledge, there are four published articles regarding the isolation of *M. hominis* from postcesarean incisions. Based on the current study and the literature, infection by this pathogen may cause hematoma formation with or without abscess after cesarean section or in immunosuppressed postoperative patients. In such cases, physicians may need to suspect *Mycoplasma* infection and initiate appropriate antibacterial treatment as soon as possible in order to avoid persistent fever.

**Keywords:** *Mycoplasma hominis*, cesarean section, hematoma, abscess

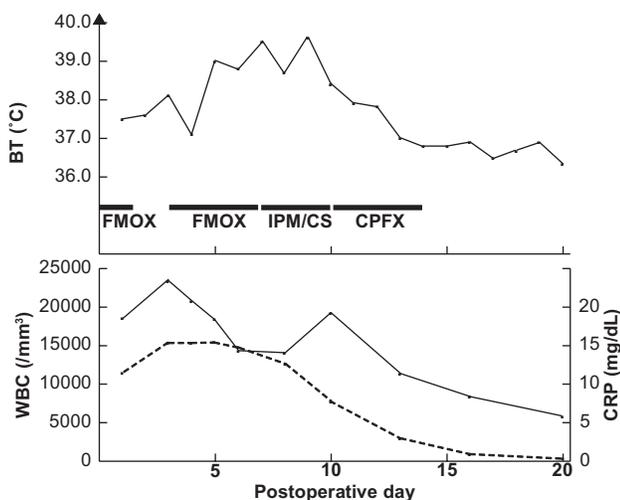
## Introduction

*Mycoplasma hominis* usually colonizes the lower urogenital tract and can cause pyelonephritis, pelvic inflammatory disease, chorioamnionitis, rupture of fetal membranes, preterm labor, postpartum fever, postabortal fever, and neonatal infection.<sup>1-9</sup> This organism is a marker for sexual activity, as demonstrated by its higher prevalence in cervicovaginal cultures of sexually active women than in those before sexual debut. *M. hominis*, *M. genitalis*, *Ureaplasma urealyticum*, and *U. parvum* may invade and infect placental and fetal tissues, leading to adverse pregnancy outcomes.<sup>1</sup> In particular, *M. hominis* and *U. urealyticum* have been documented to cause genital tract infectious diseases for several decades. *M. hominis* occasionally causes nongenitourinary infection of the blood, wounds, central nervous system, joints, or respiratory tract.<sup>10,11</sup> To our knowledge, there are four published articles regarding the isolation of this pathogen from postcesarean incisional infections.<sup>5,12-14</sup> We present a case of *M. hominis* infection after cesarean section. This case report and the literature provide the features of postoperative infection with this organism, of which physicians should remain mindful.

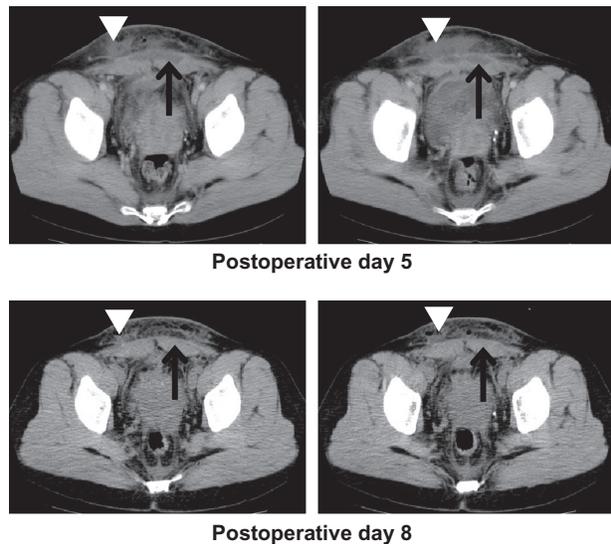
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## Case presentation

A 27-year-old woman, para 0, underwent emergency cesarean section because of labor arrest and cephalopelvic disproportion at 39 weeks of gestation. A transverse skin incision was made at the lower abdomen, and the patient was delivered of an infant with meconium-stained amniotic fluid. Four 1 g doses of flomoxef sodium prophylaxis were administered intravenously. The first dose was infused at the start of the operation, followed by the other doses 12, 24, and 36 hours later. On postoperative day 5, the patient developed a fever of 39.0°C and erythema with swelling and tenderness around the abdominal wound. Blood examination revealed a white blood cell count of 18,400/mm<sup>3</sup> and a C-reactive protein (CRP) level of 15.4 mg/dL (Figure 1). A lower abdominal computed tomography (CT) scan revealed subcutaneous hematoma and rectus abdominis muscle abscess (Figure 2). These findings prompted us to open the central portion of her abdominal wound (approximately 2 cm width) under local anesthesia, for the purpose of drainage. A bloody purulent exudate was obtained from the incision, which was subjected to bacteriological culture. The Gram stain indicated numerous altered polymorphonuclear leukocytes without any pathogens (Figure 3). We irrigated the incision with 100 mL of saline daily and intravenously administered 1 g flomoxef sodium twice daily. On day 8, an abdominal CT scan suggested that the subcutaneous hematoma and rectus abdominis muscle abscess had subsided (Figure 2), indicating that our drainage was effective. Nevertheless, her fever persisted, and the white blood cell count and CRP level

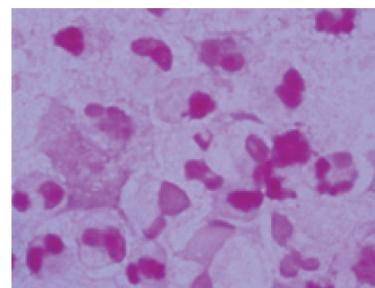


**Figure 1** Body temperature (BT), white blood cell count (WBC), and C-reactive protein (CRP) level after cesarean section. One gram of flomoxef sodium (FMOX) twice daily, 500 mg of imipenem-cilastatin sodium (1:1; IPM/CS) twice daily, and 300 mg of ciprofloxacin (CPFXX) twice daily were administered intravenously. Solid line: WBC; broken line: CRP. CPFXX chemotherapy resolved the patient's fever when WBC and CRP diminished to their normal ranges.

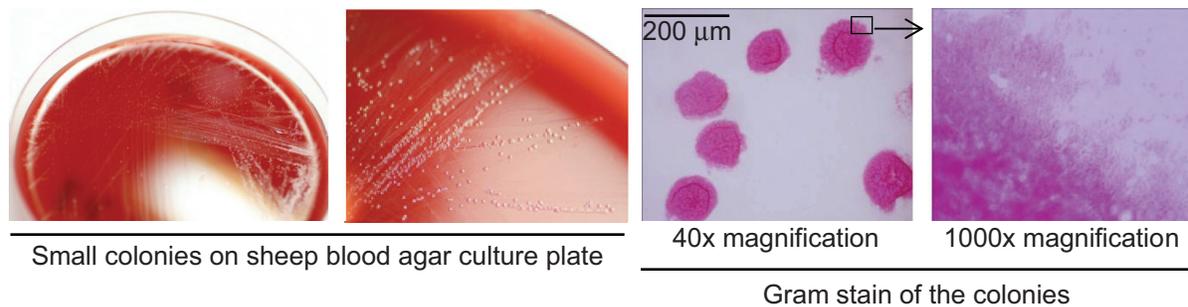


**Figure 2** Computed tomography scans. Rectus abdominis muscle abscess (arrow) and subcutaneous hematoma (arrowhead) were revealed on postoperative day 5 and had subsided on day 8.

remained high. On day 7, we began an intravenous infusion of imipenem-cilastatin sodium (1:1; 500 mg) twice daily, instead of flomoxef sodium. On day 10, the abdominal incisional exudative fluid specimen obtained on day 5 revealed the growth of tiny colonies on a sheep blood agar culture plate under anaerobic conditions. The colonies were pinpoint sized, translucent, measuring 50–300 μm in diameter, and grew on blood agar in an anaerobic or 5% carbon dioxide atmosphere in 2–3 days but showed no growth in an aerobic environment in subculture. On Gram stain of the colonies, the cells were so small that their cellular morphology was invisible under a routine light microscope (Figure 4). The isolate was identified as *M. hominis* based on morphology of colonies, size of cells, and culture condition, being compatible with unsuccessful chemotherapy using a β-lactam agent. Therapy with a quinolone, ciprofloxacin (300 mg twice daily), was started intravenously, and imipenem-cilastatin sodium therapy was discontinued. This antibacterial treatment, during which breastfeeding was avoided, was continued until day 14 when



**Figure 3** Gram stain of the abdominal incisional exudate. This indicates numerous altered polymorphonuclear leukocytes without any pathogens.



**Figure 4** *Mycoplasma hominis* colonies grown in an anaerobic environment. They were pinpoint sized, translucent, and measuring 50–300  $\mu\text{m}$  in diameter. Subsequently, the colonies were Gram stained to be visualized more distinctly.

her fever was resolved. The white blood cell count and CRP level diminished to their normal ranges, and the patient was discharged home 21 days after the cesarean section.

## Discussion

*Mycoplasma* species lack a cell wall. They appear as an amorphous mass on Gram staining and are therefore unidentifiable by this method.<sup>9,13</sup> In addition, *M. hominis* is a slow-growing bacterium, and its identification and antimicrobial sensitivity testing with routine culture methods are difficult. Therefore, appropriate antibacterial therapy is generally started late. Incubation in an anaerobic environment may enhance the growth of this pathogen,<sup>9</sup> although 48–96 hours are usually required for its growth on nonselective routine bacteriological media. Furthermore, this organism is cultured most successfully on designed media, such as Shepard's 10B urea broth and A8 solid medium, after 24–48 hours of incubation.<sup>15</sup> Media designed for detection of *Mycoplasma* are commercially available (Remel Inc., Lenexa, KS, USA). *M. hominis* colonies are pinpoint sized, flat, measuring 20–300  $\mu\text{m}$  in diameter, and have a characteristic fried-egg appearance when observed under 10 $\times$  magnification. In our case, the pathogen grew under anaerobic conditions.

*Mycoplasma* species do not have a cell wall, which is the target of  $\beta$ -lactam antimicrobials that are commonly used as prophylactic agents for postsurgical infections. Erythromycins are also less effective against *M. hominis*,<sup>10,13</sup> whereas *M. pneumoniae* is susceptible to these drugs. Tetracycline has historically been considered the antibiotic of choice for treating *M. hominis* infections,<sup>9,16</sup> and clindamycin and lincomycin chloride are often used to treat infection with tetracycline-resistant strains.<sup>9,10,17</sup> Quinolones, including ciprofloxacin, ofloxacin, temafloxacin, sparfloxacin, trovafloxacin, gemifloxacin, gatifloxacin, moxifloxacin, and garenoxacin, are equally active against tetracycline-susceptible as well as tetracycline-resistant strains of *M. hominis*.<sup>18–24</sup> Because *M. hominis* is typically resistant

to macrolides,  $\beta$ -lactams, aminoglycosides, vancomycin, trimethoprim, and sulfonamides,<sup>9,10,13,17,23,25</sup> administration of effective antimicrobial agents is often initiated late.

Due to the aforementioned reasons, high fever and inflammation persist for a long time in many cases of *M. hominis* infection. Polymerase chain reaction (PCR) is a simpler, more rapid, and more sensitive method than culturing for the identification of this organism; therefore, a PCR assay should be performed if *M. hominis* infection is suspected.<sup>26</sup>

In our patient, abdominal wound hematoma and abscess due to *M. hominis* were formed following cesarean section, and drainage and the administration of a quinolone, ciprofloxacin, improved the infection. To our knowledge, there are four published articles on the isolation of this organism from postcesarean incisions in which the patients developed hematomas.<sup>5,12–14</sup> In one patient with *M. hominis* wound infection after cesarean section and in three patients with *M. hominis* and *U. urealyticum* infection, aspiration of the incision yielded a cloudy serosanguinous fluid.<sup>12,13</sup> The serosanguinous exudates suggest the presence of abdominal incisional purulent hematoma. In a case of hematoma and abscess formation due to *M. hominis* infection at the site of a uterine incision after cesarean section, drainage and administration of a quinolone, pazufloxacin mesilate, promptly resolved the patient's fever.<sup>14</sup> A perihepatic hematoma with *M. hominis* was observed after a liver transplant,<sup>27</sup> and a perinephric hematoma was infected with this pathogen after a renal transplant.<sup>28</sup> Moreover, hematoma infection with this bacterium developed following transplant nephrectomy.<sup>29</sup> These indicate that *M. hominis* infections after surgery may cause hematoma formation in immunosuppressed patients. When a hematoma with bacterial infection or an abscess presents, drainage, aspiration, and debridement are promising treatments. Physicians may need to subject the specimens to bacteriological culture in an anaerobic environment, on designed media, or with PCR assay, to initiate appropriate antibacterial therapy as soon as possible if the causative organism is *Mycoplasma*.

The route of wound infection by *M. hominis* in this case is unknown. One possible speculation is that the pathogen invaded and infected fetal membranes and amniotic fluid from the vagina or uterine cervix during pregnancy, resulting in postcesarean infection.

## Conclusion

We present a case of abdominal wound infection by *M. hominis* following cesarean section. Based on our observation, this pathogen may cause hematoma formation with or without abscess after cesarean section or in immunosuppressed postoperative patients. In such cases, physicians may need to suspect *Mycoplasma* infection and start effective antimicrobial treatment as soon as possible in order to avoid persistent fever.

## Disclosure

The authors report no conflicts of interest in this work.

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