What is Missing in STD Screening in Hong Kong?

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Abstract: This retrospective analysis was to study the prevalence of Sexually Transmitted Diseases (STD) in a Reproductive Medical Center in Hong Kong. A total of 1190 patients were included in this study. Group 1A, couples had no symptoms but presented with subinfertility; Group 1B, the subfertile couples were positive for either Chlamydia trachomatis (CT) or Ureaplasma urealyticum (UU); Group 2, couples with symptoms were offered full STD screening including CT, UU, Mycoplasma hominis (MH), Neisseria gonorrhoea (NG), Syphilis, Herpes Simplex Virus 1 and 2. The methods of ELISA and quantitative real-time PCR were used for these analyses. Group 1A: UU detection rates in both male and female (13.84 v.s. 37.07%) were significantly higher than that of CT (5.65 v.s. 5.57%); Group 1B and Group 2: For those who had Full STD check, UU and MH detection rates were significantly higher than that of CT (35, 13.7 and 7.1% respectively). Over 47% of patients showed positive for one or more organism. In the subfertile couples, the UU and CT detection rates were much higher in females than those in males. Both semen and urine samples gave the same rates of infection among CT, UU, NG and MH. U. urealyticum infection rate rather than Chlamydis trachomatis infection is highest in Hong Kong. The infection rate in females is higher than in males. The detection rates in semen and urine samples in males are similar.

Keywords: STD, Reproduction, Ureaplasma urealyticum, Chlamydia trachomatis, Hong Kong

Introduction

The screening for Sexually Transmitted Diseases (STD) is a routine procedure for all patients receiving fertility treatment. With an increasing demand on sperm donors and egg donors, STD screening draws more attention to both patients and professionals. In U.S., STD prevalence was 15 million (Cates, 1999) and this number continues to rise every year. To date, there are not many publications on STD in the infertility field. Such information would shed light on making guidelines by the regulatory bodies.

According to the Hong Kong guidelines for screening of potential gamete/embryo donors and recipients against infectious diseases, the following serological tests should be performed:

- HIV
- Syphilis
- Hepatitis B surface antigen (HBsAg)
- Hepatitis B core antibody (anti-HBc) (IgG and IgM)
- Hepatitis C antibody
- Antibody tests (IgG and IgM) for CMV
- Semen, urine or urethral cultures for Neisseria gonorrhoea. Either urethral or urinary testing for Chlamydia trachomatis should be performed

CT is an obligate intracellular human pathogen. It is a gram-negative bacterium which appears as either cocoid or rod shape under light microscope. CT includes three human biovars: (1) Serovars Ab, B, Ba or C. They cause infection of the eyes (trachoma), which can lead to blindness. (2) Serovars D-K can cause urethritis, pelvic inflammatory disease, ectopic pregnancy, neonatal pneumonia and neonatal conjunctivitis. (3) Serovars L1, L2 and L3 which could cause lymphogranuloma venereum.

CT infections may cause chronic pelvic pain, ectopic pregnancy and infertility. Patients with CT infections are often not aware of their disease. In U.S., there are approximately 4 million people infected each year. Its...
prevalence is about 5% (CDCP, 2009). In South Africa, the reported infection rates were higher (5.9-33%) among general populations showing a big regional difference (Buve et al., 2001; Isibor et al., 2005; Nwankwo and Sadiq, 2014). Therefore CT has been the main target for general population screening.

Non-specific urethritis, proctitis (rectal disease with bleeding), trachoma and infertility are seen in both genders. The most commonly seen in men are prostatitis and epididymitis. Women, once infected, would suffer from cervicitis, pelvic inflammatory disease, ectopic pregnancy and acute/chronic pelvic pain. Trachoma and pulmonary infection are also seen in neonates. There are approximately 600 million people infected with Chlamydia worldwide, of which 20 million became blind. In Hong Kong, it is unclear whether there are other pathogens which need to be screened apart from the routinely screened STDs.

The aims of this study are to address three questions, namely, (1) Is CT the most prevalent organism in Hong Kong? (2) What is the relative prevalence of the various STD organisms? (3) For males, which sample is the more sensitive for STD screening, semen or urine?

Materials and Methods

This is a retrospective study encompassing over 1190 patients, approved by the local Ethic Committee, chaired by the first author, Clement Leung-Kwok Chan. The urine and semen samples were routinely collected from the patients who visited the Women’s Health and Reproductive Medical Center in Hong Kong during July 1, 2005 to June 30 2013. Patients were divided into two groups based on their presenting symptoms.

Group 1A: Female patients presented with subfertility or with vaginal discharge and male subfertility patients with discharges went through initial screening for DNA of CT (n = 1039) and UU (n = 1190) using the method of Real Time PCR.

Group 1B: If these patients were positive for the above initial testing, the couples were tested with full STD checking including CT, UU, NG, syphilis (VDRL or RPR), IgG of Herpes Simplex Virus (HSV) 1 and 2.

Group 2: Full STD check was also carried out for the patients who presented with pelvic or genital infection, or pelvic pain (n = 620).

Real-Time PCR

Detection of CT, NG, MH, UU DNA in endocervical scraping, semen and urine samples was conducted by the methods of real-time PCR using Applied Biosystem 7500 (based on 36% prevalence of infection, sensitivity, specificity, PPV and NPV were 87, 96, 94 and 93% respectively).

Syphilis Serology Detection

Rapid Plasma Reagin (RPR): RPR Card Test (Arlington Scientific, Inc., ASI) for Syphilis was conducted. It is a qualitative and semi-quantitative non-treponemal flocculation test for detection of reagin antibodies in human serum and plasma.

Enzyme-Linked Immunosorbent Assay (ELISA)

Antibodies against HSV type 1 and HSV type 2 were analyzed by ELISA with 100% sensitivity and specificity. The Chlamydia antigens were analyzed by ELISA. The antibodies against Chlamydia antigens were detected by Direct Fluorescent Antibody Test (DFA).

Chlamydia Cell Culture

Suspected Chlamydia samples were cultured in a vial of cells in the laboratory. The pathogen infected cells, after 48 h (up to 2 days), were stained and viewed under a fluorescent light microscope.

The data were analysed by Chi Square Test.

Results

Results of Group 1A were tabulated in the Table 1. The infection rate for CT was similar in both female and male patients (5.57 and 5.65%). Both female and male patients had higher detection rates of UU compared with those of CT (37.07 v.s. 5.57%, P<0.001 in females; 13.84 v.s. 5.65%, P = 0.006 in males). Also, detection rate for UU was significantly higher in females (37.07%) compared to males (13.84%, P<0.001).

Among the full STD check (Group 1B and Group 2, n = 620), 52% of the patients were negative, whereas UU detection rate was highest (27.9%) in all categories detected (Table 2).

Table 3 showed detection rates in those with full STD check in relation to the presenting symptoms. There is no difference among the diagnoses. However, the detection rate for HSV1 was the highest (74.5%, P = 0.015), followed by the second highest of UU infection (35.5% P<0.001) (Table 3).

Table 1. Initial screening of CT and UU for patient Group 1A

<table>
<thead>
<tr>
<th></th>
<th>Chlamydia trachomatis</th>
<th>Ureaplasma urealyticum</th>
<th>Pvalue</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female</td>
<td>48/862 (5.57%)</td>
<td>334/901 (37.07%)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Male</td>
<td>10/177 (5.65%)</td>
<td>40/289 (13.84%)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Pvalue</td>
<td>&gt;0.05</td>
<td>0.001</td>
<td></td>
</tr>
<tr>
<td>No. of cases</td>
<td>1039</td>
<td>1190</td>
<td></td>
</tr>
</tbody>
</table>
Table 2. Full STD screening

<table>
<thead>
<tr>
<th>Categories</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>CT</td>
<td>22/620 (3.55%)</td>
</tr>
<tr>
<td>NG</td>
<td>1/620 (0.16%)</td>
</tr>
<tr>
<td>UU</td>
<td>173/620 (27.90%)</td>
</tr>
<tr>
<td>MH</td>
<td>47/620 (7.58%)</td>
</tr>
<tr>
<td>CT+UU</td>
<td>13/620 (2.10%)</td>
</tr>
<tr>
<td>CT+</td>
<td>6/620 (0.97%)</td>
</tr>
<tr>
<td>NG+UU</td>
<td>4/620 (0.65%)</td>
</tr>
<tr>
<td>UU+MH</td>
<td>27/620 (4.35%)</td>
</tr>
<tr>
<td>CT+UU+MH</td>
<td>4/620 (0.65%)</td>
</tr>
<tr>
<td>All negative</td>
<td>323/620 (52.10%)</td>
</tr>
</tbody>
</table>

Table 3. Association of different infections and various diagnoses

<table>
<thead>
<tr>
<th>CC NG MH UU VDRL/RPR HSV type 2 IgG HSV type 1 IgG</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frequency</td>
</tr>
<tr>
<td>Pelvic discomfort</td>
</tr>
<tr>
<td>Subfertility</td>
</tr>
<tr>
<td>Vaginal discharge</td>
</tr>
<tr>
<td>Unclassified</td>
</tr>
<tr>
<td>Total</td>
</tr>
</tbody>
</table>

Table 4. Association of different infections and gender

<table>
<thead>
<tr>
<th>Male Female</th>
<th>CC NG MH UU VDRL/RPR HSV type 2 IgG HSV type 1 IgG</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frequency</td>
<td>10/81 (12.35%)</td>
</tr>
<tr>
<td>Pelvic discomfort</td>
<td>1/92 (1.09%)</td>
</tr>
<tr>
<td>Subfertility</td>
<td>13/92 (14.13%)</td>
</tr>
<tr>
<td>Vaginal discharge</td>
<td>26/92 (22.83%)</td>
</tr>
<tr>
<td>Unclassified</td>
<td>21/92 (22.83%)</td>
</tr>
<tr>
<td>Total</td>
<td>78/92 (84.78%)</td>
</tr>
</tbody>
</table>

Discussion

This study, for the first time in Hong Kong, revealed that UU infection rates in both male and female subfertile patients were higher than those of CT infection; Among the full STD screening, UU still represented the highest rate of infection; Among CT, NG, MH and UU, MH showed significantly higher detection rates compared to that of CT. There was no significant difference among patients with different presenting symptoms; in the subset of 127 subfertility couples, the UU and CT detection rates in female were significantly higher than the those of the male partners; lastly, both semen and urine samples showed the same rate of infections.

Many studies in the past two decades have shown great regional difference worldwide. A study reported that in Israel when the patients showed U. Urealyticum positive, the chance of Condyloma Acuminatum was higher (Zvulunov et al., 2000). Our results further demonstrated U. Urealyticum and HSV type 1 should be included in the routine STD screening in Hong Kong, especially for the subfertile patients. As early as in 1997, a large scale study including 41,980 patients in Vienna, Austria, who suffered from infectious Venere-Dermatological Diseases, Mycoplasma Hominis and Ureaplasma urealyticum, were detected more often in the vaginal fluid that in the male urethra (Koch et al., 1997).

Ureaplasma have two species-urealyticum and parvum. Ureaplasma urealyticum is a bacterium belonging to the family of Mycoplasmataceae. U. parvum serovar 3/14 and T960 biovar were found to be significantly associated with symptomatic patients, while on the contrary, U. parvum serov 6 was significantly correlated with asymptomatic women and normal vaginal flora (De Francesco et al., 2009). Ureaplasmaparvum is more prevalent than Ureaplasma urealyticum in specimens from infertile men (Abusarah et al., 2013). U. urealyticum is part of the normal genital flora of both men and women. Indeed its infection was reported not affecting semen quality (Wang et al., 2006; Gdoura et al., 2007). However, U. Urealyticum infection is closely related to reduction in the pregnancy rate of IVF patients, possibly because of its infection at the endometrial level owing to the fact that fertilization rate and the total number of eggs retrieved was not affected (Montagut et al., 1991).
normal flora of sexually active women. However, it is not known that both may cause chorioamnionitis, salpingitis, acetalvaginositis and postpartum endometritis and therefore they should no longer be considered as commensals (Larsen and Hwang, 2010). In addition, it was reported that Ureaplasma was detected positive in 50% preterm babies in one or more compartments (i.e., Respiratory, blood and/or cerebrospinal fluid) (Viscardi, 2014).

Moreover, Potts et al. (2000) reported seminal Reactive Oxygen Species (ROS) levels were elevated among patients infected with U. urealyticum. These authors suggested that ROS induces lipid peroxidation, which reduces membrane fluidity and sperm fertilization capability, possibly explaining how U. urealyticum impairs sperm function. The level of Zinc and Selenium in the seminal fluid is decreased in Ureaplasma infected patients (Han et al., 2003). Based on the characteristics, U. urealyticum has two genotypes-biovar1 (serovars 1, 3, 6 and 14) and biovar II (serovars 2, 4, 5 and 7-13) (Povlsen et al., 1998). Biovar II is more likely related to male infertility (Zhang et al., 2014a). This work was conducted in Southern China. Whether it is also the same in Hong Kong needs to be further studied.

The tissue sources and methods of detection for STD pathogens resulted in a variety of outcomes. The tests of using menstrual tissues were proven to be sensitive, allowing early detection and early treatment of a condition that can otherwise lead to serious consequences such as tubal obstruction, pelvic inflammatory disease, ectopic pregnancy, spontaneous abortions and unexplained infertility (Michou et al., 2014). Detection of Neisseria Gonorrhoeae in pregnant women in Iran showed that endocervical swab specimens should be detected by both Nucleic Acid Amplification Tests and culture method to limit false negative samples (Hassanzadeh et al., 2013). Recently a novel method of using bead-based multiplex Sexually Transmitted Infection Profiling was reported (Schmitt et al., 2014). This method specifically detects Chlamydia trachomatis, HSV type 1 and 2, Treponema pallidum, Trichomonas vaginalis, Neisseria gonorrhoeae, Mycoplasma, M. genitalium, M. hominis, M. pneumonia, M. spermatophilum, Ureaplasma urealyticum and U. Parvum and quantifies bacterial vaginosis-associated Atopobium vaginae and Gardnerella vaginalis as well as three Candida species and normal genital flora-associated Lactobacillus species. Moreover, a Multilocus sequence typing scheme based on four housekeeping genes (fisH, rplL22, valS, thrS) is more adequate for investigations of molecular epidemiology and population structure with highly discriminating capacity (Zhang et al., 2014b).

Our results concluded both semen and urine samples presented similar detection rates for C. trachomatis, N. gonorrhoeae, M. hominis, U. urealyticum, syphilis and HSV type 1 and 2. A similar finding was reported that there is no difference between semen samples and First Void Urine (FVU) samples in detecting C. trachomatis, as well as U. urealyticum in infertile patients in Jordan (Abusarahet et al., 2013) whereas N. Gonorrhoeae was found to be present in semen rather than FVU which was different from our result.

As far as treatments are concerned, the Ureaplasma are tested against azithromycin, josamycin, ofloxicin and doxycycline. Resistance to macrolides, tetracyclines and fluoroquinolones has been reported (Kokkayil and Dhawan, 2015). Patients with infection of U. urealyticum have been associated with low birth weight and premature delivery (Kafetzis et al., 2004). Therefore appropriate treatments prior to conception are essential in achieving healthy pregnancies.

### Conclusion

1. Ureaplasma urealyticum infection rates in both male and female were significantly higher than those of Chlamydia trachomatis infection; 2. Among the full STD check U. urealyticum still represents the highest rate of infection; 3. U. urealyticum exhibited the second highest infection among all diagnoses including pelvic discomfort, subfertility, vaginal discharge and unclassified diagnoses; 4. In female patients, U. urealyticum infection rate is much higher than that in the males; 5. Both semen and urine samples gave the same rate of infection among C. trachomatis, N. gonorrhoeae, M. hominis, U. urealyticum, Syphilis and HSV type 1 and 2.

### Acknowledgement

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### Author’s Contributions

**Clement Leung-Kwok Chan:** Initiated project; Recruiting patients; Data collection; Experimental Design and contribution to the key ideas.

**Wai Yee Chan:** Manuscript revision; Experimental design and Critical Review for significant intellectual content.

**Christopher Hon Ki Cheng:** Manuscript revision; Data analysis and Critical Review for significant intellectual content.
Ping Xia: Interpretation of the data; manuscript draft; Critical Review for significant intellectual content and Manuscript submission and revisions.

Ethics

No ethical issues should arise from this publication.

References


