Detection of ureaplasma urealyticum by polymerase chain reaction examination in nonspecific genital infection patients

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Abstract

Non specific genital infection (NSGI) is a condition affecting females which causes inflammation of the endocervix or anterior urethra that is not caused by Neisseria gonorrhoeae. The causative sexually transmitted organisms include Chlamydia trachomatis (Groups D to K) and Ureaplasma urealyticum. Infection caused by Ureaplasma urealyticum is often asymptomatic even though many studies have pronounced that Ureaplasma urealyticum can contribute not only to lower genitourinary infection but also to infertility. Ureaplasma urealyticum cannot be stained by Gram stain due to the lack of a cell wall of the organism. This research aims to evaluate the prevalence of Ureaplasma urealyticum in NSGI patients by using the polymerase chain reaction (PCR) method targeted in the ureaplasma gene structure 429 bp area. The samples were extracted from eighteen DNA NSGI patients. Eleven out of eighteen (61.11%) DNA NSGI samples tested positive for Ureaplasma urealyticum. Most patients (44.44%) with Ureaplasma urealyticum were unemployed, and 27.78% were complaining of recurrent vaginal discharge. The high incidence of Ureaplasma urealyticum in this study needs further attention since doxycycline remains the drug of choice of NSGI. Moxifloxacin should be considered for patients who are making no clinical progress with doxycycline.

Introduction

Non specific genital infection (NSGI) is a condition affecting females which causes inflammation of the endocervix or anterior urethra that not caused by Neisseria gonorrhoeae (Hong Kong Social Hygiene Service, 2004). The term non specific is used if the causal organisms cannot be detected by conventional microscopy method. The causative sexually transmitted organisms include Chlamydia trachomatis (Groups D to K) and Ureaplasma urealyticum gonorrhoeae (Hong Kong Social Hygiene Service, 2004). The prevalence of NSGI at the Outpatient Clinic of Dr. Soetomo General Hospital in 2016 was 47 out of 3,753 new dermatology and venereology cases (1.25%). In the Sexually Transmitted Disease Division of the Outpatient Clinic, the percentage of NSGI was 17.22% (47 out of a total of 273 new cases). Determining the diagnosis of NSGI requires detailed anamnesis of patients and patient’s partner’s complaints, risk factors and obstetric history. On physical examination, hyperemia, erosion of the cervix and mucopurulent discharge may be found. In microscopy examination from the cervical smear with Gram staining is regarded as positive if 10 to 30 polymorphonuclear leucocytes were seen per high-power field and all other specific bacteria or fungal are not found such as diplococci Gram negative bacteria (Neisseria gonorrhoeae), Trichomonas vaginalis, candidiasis vulvovaginalis and bacterial vaginos.1

Based on the literature, Chlamydia trachomatis is the most common (30%-50%) cause of NSGI, which is inconsistent with the recent study conducted in the Dermatovenereology Outpatient Clinic of Dr. Soetomo General Hospital, Surabaya in 2017. This study found a low incidence (16.67%) of Chlamydia trachomatis in eighteen married NSGI patients.2 Ureaplasma urealyticum, the second most common cause of NSGI (10%-40%) is frequently found in the commensal flora of the lower genital tract.2 Nevertheless, Ureaplasma species are the most prevalent, potentially pathogenic bacteria isolated from the urogenital tract of both men and women. By evolving from Gram-positive bacteria by degenerative evolution, Ureaplasmas lose their peptidoglycan cell wall. The lack of a cell wall leaves these organisms insensitive to beta lactams, and also prevents the organisms from Gram staining. Ureaplasma has 14 known serotypes and is divided into two groups: Ureaplasma parvum (UPA, biovar 1, parvo) and Ureaplasma urealyticum (UUR, biovar 2, T960). In a study conducted by Dhawan B et al, Ureaplasma was found in 25.8% of patients with genital tract infections and in 20.8% of infertile women. Previous studies have shown that Ureaplasma urealyticum biovars were associated with pathogenicity. A study by Chua KB et al, stated that biovar 2 was more associated with the loss of lactobacilli in women than biovar 1. They verified that biovar 2 was associated with genitourinary tract infections (58.18%) compared to biovar 1, which was only a colonizer of the genitourinary tract. Several clinical reports stated that urogenital infections caused by Ureaplasma urealyticum may cause abnormal pregnancy outcomes by inducing bacterial vaginosis, cervicitis, chorioamnionitis, intrauterine infection, premature rupture of membranes, preterm delivery and neonatal pneumonia.3,4

Ureaplasmas are susceptible to antimicrobial agents that influence DNA, RNA, protein synthesis or the integrity of the cell membrane; these include tetracyclines, macrolides, chloramphenicol, aminoglycosides and fluoroquinolones. However, susceptibility to macrolides is moderate; a recent study indicated that the most common cause of recurrent or persistent urethritis is mixed infection with Mycoplasma genitalium or Ureaplasma urealyticum, particularly among those patients who have been treated with doxycycline, a drug to which Ureaplasma urealyticum or Mycoplasma genitalium may be resistant.1,5 Tetracycline resistance in Ureaplasma species has been detected and
resistance is mediated by the tetM determinant, which encodes a protein that binds to the ribosomes, protecting Ureaplasma spp. from the actions of these drugs. Abele Horn et al have studied the antibiotic susceptibility of the two Ureaplasma biovars and detected a major resistance to doxycycline and older fluoroquinolones. The resistance rate of the Ureaplasma urealyticum biovar to doxycycline was 55%, to ciprofloxacin it was 42% and to ofloxacin it was 61%. Moxifloxacin was the most active agent in vitro against Ureaplasma urealyticum with the narrowest difference between the minimal inhibitory concentration (MIC) and the minimal bactericidal concentration (MBC) values. Moxifloxacin should be considered for empirical therapy of non-gonococcal non-chlamydial infection, or if the symptoms still persist after gonococcal and/or chlamydial infection has been eradicated. Hundreds of publications have described various nucleic acid amplification tests (NAATs) and their applications in detecting mycoplasmas and ureaplasmas in clinical specimens since 1989. NAATs are also useful for the identification of organisms grown in culture to the species level, replacing older and less practical technologies. Molecular based methods, such as PCR, are able to detect and identify Ureaplasma urealyticum and Ureaplasma parvum separately, whereas culture cannot do this. PCR has also been adapted to detect antimicrobial resistance determinants and to analyze the genetic relatedness of clinical isolates. A study by Dhawan et al. established the prevalence of Ureaplasma urealyticum with genital discharge by both culture and PCR. The PCR targeted a 429 bp region in the urease structural gene of Ureaplasma urealyticum. The prevalence of Ureaplasma urealyticum as determined by culture was 32% while PCR was 45% with an agreement of 93.75%. PCR seems to be more sensitive for diagnostic purpose compared to culture; PCR detected twenty (15.2%) more positive samples among 132 clinical specimens compared to culture.

The aim of this study is to discover the incidence of Ureaplasma urealyticum in NSGI patients since it is important to understand the exact causative organism so that precise management can be given to the patients.

Materials and Methods

This research is a descriptive observational study, using a cross sectional method to evaluate the prevalence of Ureaplasma urealyticum among NSGI patients. This study used the extracted DNA of eighteen NSGI patients that were archived in the Tropical Disease Center (TDC), Universitas Airlangga Surabaya and performed PCR examinations in February 2018. Samples were collected from NSGI patients who came to the Dermatovenereology Outpatient Clinic (Sexual Transmitted Disease Division) of Dr. Soetomo General Hospital, Surabaya, and were taken consecutively for three months (June – August 2017). Inclusion criteria included women with IGNS who are married. Women who were menstruating, pregnant or diagnosed with mixed infection were excluded. Specimen samples were taken from endocervical swab which the diagnosis of NSGI had already established by detailed anamnesis and physical and light microscopy examinations. Informed consent was obtained from the patients prior to procedure. PCR examinations were performed on the DNA of eighteen NSGI patients, extracted from endocervical swabs. The PCR used in this study was MyQ-2 (Bio Rad). The PCR targeted a 429 bp region in the urease structural gene of Ureaplasma urealyticum that is forward strand: 5’-ACGAC GTCCA CTG TAAGC AACT-3’ and reverse strand: 5’-CAATC TGTC GTGAA GTATT AC-3’. Ethical clearance had been approved for this study by the Ethical Committee of RSUD Dr. Soetomo General Hospital.

Discussion

Although Ureaplasma urealyticum is known to be frequently found in the non-pathogenic flora of the lower genital tract, this organism is potentially pathogenic isolated from the urogenital tract of both men and women and causes 10% - 40% of NSGI. Some clinical reports declared that urogenital infection caused by Ureaplasma urealyticum can cause chorioamnionitis, spontaneous abortion, stillbirth and preterm abortion during pregnancy. Ureaplasma has been isolated from patients with pelvic inflammatory disease (PID) affecting fallopian tube; it was also detected in 25.8% of patients with genitourinary tract infections and 20.8% of infertile women. Detection of the exact organisms is important, but the lack of rigid cells makes...
it nearly impossible to directly visualize Ureaplasma by light microscopy, and culture is difficult since these fastidious organisms require the presence of serum, metabolic substrate and growth factors like yeast extract for isolation. In addition, routine bacterial culture may give negative results for commercial sexual workers or asymptomatic people who have recently experienced unprotected sexual contact and have acquired sexual transmitted infections (STI). These neglected asymptomatic patients in the community can be the reservoir of STIs. The identification of the specific organism leads the clinician to make precise management of the infections so that no complication will occur. Currently, the focus is on syndromic management, which is not a very sensitive or specific method for establishing diagnosis of upper genital infection. Furthermore, because of ever increasing drug resistance, it is better to diagnose and treat the specific causative organism as far as this is possible. In a study involving patients with infertility and genital discharge, Ureaplasma spp. 91% were susceptible to doxycycline, 77% to ofloxacin, and 71% to azithromycin. Govender and Chalkley mentioned nine tetracycline-resistant strains concurrently resistant to doxycycline. Another study by Kenny and Cartwright stated Ureaplasmas were susceptible to quinolones with the highest activities being shown by moxifloxacin and sparfloxacin. Most pathogens causing STI as well as commensal microorganisms are difficult to cultivate by routine microbiological diagnosis. NAATs, such as PCR, are useful for the identification of microorganisms that are difficult to cultivate and for those that grow slowly. The present study used PCR and targeted a 429 bp region in the urease structural gene of Ureaplasma urealyticum that is forward strand: 5’-AGGAC GTCCA CTG TAAGC AACT-3’ and reverse strand: 5’-CAATC TGCTC GTGAA GTTT AC-3’. The samples used in this study were the extracted DNA of eighteen NSGI patients. The results of this study performed in TDC Universitas Airlangga Surabaya and the sample specimens collected in Dr. Soetomo General Hospital, Surabaya, determined that eleven out of eighteen NSGI patients (61.11%) were detected Ureaplasma urealyticum, using the PCR method. This prevalence was in keeping with the study by Peerayeh et al on infertile women with endocervical specimens, which found that 51.7% tested positive for Ureaplasma urealyticum by the PCR method. Our study also found four patients (22.22%) were aged between 28 – 37 years, which is comparable to the study by Peerayeh et al, which found that 20.7% of patients with Ureaplasma urealyticum were aged 28 – 37. There is no previous study that has been performed in similar populations.

Conclusions
To determine the specific causative organism of NSGI is important, due to the severe complications that may be caused. The increasing resistance of tetracycline in Ureaplasma strains requires careful antimicrobial preferences. This study may prove useful in providing information on the prevalence of Ureaplasma urealyticum and the management of NSGI caused by non chlamydial infection. The high incidence of Ureaplasma urealyticum in this study needs further attention since doxycycline remains the drug of choice of NSGI. Moxifloxacin should be considered for patients who are making no clinical progress with doxycycline. Research concerning the susceptibilities and resistances of Ureaplasma urealyticum to antimicrobial classes such as macrolides, tetracyclines and quinolones may also need to be undertaken.

References