Prevalence of Trichomoniasis by PCR in Women Attending Health Screening in Korea

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Abstract: Trichomoniasis is the most common curable sexually-transmitted infection (STI) worldwide. There are few reports on the prevalence of Trichomonas vaginalis in Korea. The purpose of this study was to examine the prevalence of trichomoniasis by PCR in Guri city, Korea. All adult women who visited Hanyang University Guri Hospital for health screening within the National Health Care Service were invited to participate in the study, and 424 women were enrolled between March and June 2011. PCR was used to detect Trichomonas vaginalis using primers based on a repetitive sequence cloned from T. vaginalis (TV-E650). Fourteen women (3.3%) were found to have T. vaginalis. All were over 50, and they were significantly older on average than the 410 Trichomonas-negative women (mean ages 63.4 vs 55.3 years). It seems that T. vaginalis infection is not rare in women receiving health screening, especially among those over 50.

Key words: Trichomonas vaginalis, PCR, prevalence, woman

Trichomoniasis is the most common curable sexually-transmitted infection (STI) in the world. Each year the pathogen causes an estimated 248 million new cases worldwide [1]. However, it has received much less consideration than other parasitic and sexually transmitted diseases. Increasing recognition of the sequelae of infection, including increased risk of infection with human immunodeficiency virus and adverse outcomes of pregnancy, has led to increased interest in T. vaginalis [2]. Although various methods such as wet mount, the Papanicolaou test, and culture are used to diagnose trichomoniasis, PCR is considered the most sensitive test. Following the introduction of PCR, more men were reported to be infected with T. vaginalis than considered when other diagnostic tests were used. Using PCR-ELISA of urine, Sena et al. [3] detected T. vaginalis infection in 177 (71.7%) of the male partners of 256 symptomatic women residing in Guri city, Korea. All adult women who visited Hanyang University Guri Hospital for health screening within the National Health Care Service were invited to participate in this study. All the subjects provided informed consent (IRB No. 2010-078). A total of 424 women were enrolled between March and June 2011. All the women had problem-directed histories taken and underwent a physical examination.

We used specimens of vaginal discharge for culture and
PCR. A sterile cotton-tipped applicator was used to swab the vaginal discharge, and the applicator was placed in a tube containing 5 ml of TYM medium and incubated at 37°C for 2-5 days. The presence of parasites was confirmed by a microscope. For PCR, 3-5 ml of PBS was injected into the posterior vaginal fornix using a Pasteur pipette, and after pipetting the PBS several times to mix it with the vaginal discharge and it was recollected into a 15 ml tube, centrifuged, and then the pellet was stored at -70°C.

For PCR, 5 µl of vaginal sediment was mixed with 20 µl of Gene Releaser® (BioVentures Inc., Murfreesboro, Tennessee, USA), and the mixture was boiled for 5 min in a microwave oven. It was then centrifuged, and 5 µl of the supernatant was added to the PCR reaction mixture [9]. The primers were based on the T. vaginalis-specific repetitive DNA sequence in clone TV-E650-1 [10]. The primer sequences were: primer 1: 5ʹCATCCCCAACATTTTTCAA 3ʹ and primer 2: 5ʹTCCCATTTTTAGACCCTTCA 3ʹ.

The PCR reaction mixture contained 1 µl each of the primers at 10 pmol/µl each, 2 µl dNTPs (2.5 mM each), 0.1 µl Taq polymerase (5 U/µl), 5 µl pre-treated vaginal discharge, 2 µl 10× PCR buffer, 5.2 µl 5 M betaine, and 3.7 µl distilled water. The DNA was denatured for 5 min at 94°C, followed by 40 cycles of 10 sec denaturation at 98°C, 30 sec annealing at 55°C/52°C, and 30 sec extension at 72°C. To avoid product carryover, the PCR reactions were set up in an area physically separated from all activities involving amplified target sequences, thermocycling, and running of gels. To assess the sensitivity of the PCR, a suspension of T. vaginalis was counted with a hemocytometer and the numbers of trophozoites were adjusted with PBS to 1, 5, 100, and 1,000 per PCR mixture. To confirm that the 318 bp band obtained by PCR originated from T. vaginalis, nested PCR was undertaken using primer 1: 5ʹATCCCCAACAATGACGAAG 3ʹ and primer 2: 5ʹAATGTGAGGGCATGGA 3ʹ, to confirm production of an 181 bp band.

The expected product of 318 bp was obtained from as little as 1 organism (Fig. 1A), and nested PCR confirmed the 181 bp band originated from T. vaginalis (Fig. 1C). Of the 424 outpatients, 14 yielded the 318 bp band, and the prevalence of T. vaginalis obtained was 3.3% (Fig. 1B). Our previous survey by PCR of 249 asymptomatic women who visited the same hospital for health care 15 years ago gave a positive rate of 2.4% [5]. Therefore, the rate of trichomoniasis infection in Guri city may be rising.

The age of the 424 outpatients ranged from 30 to 80. Every positive patient was over 50. There were 4 each in their 50s and 60s, and 6 in their 70s. When the 424 patients were divided into 2 groups, women under 60 (n=280) and over 60 (n=144) had infection rates of 1.4% and 6.9%, respectively, which were significantly different (Table 1). The mean age (63.4 ± 8.7 years) of the 14 trichomoniasis patients was also significantly higher than that of the 410 trichomonad-negative women (55.5 ± 9.9 years) (Table 2).

It is known that few T. vaginalis-infected persons have symptoms, especially, in the case of men. Among the 14 women with trichomoniasis, only 7 had symptoms such as dysuria, pruritus, burning sensation, dyspareunia, and postcoital bleeding. Therefore, significant numbers of asymptomatic T. vagina-
It is very interesting that the average age of the 14 infected women was over 60. In contrast, Goo et al. [6] reported that the median age of 19 infected subjects among 621 women who visited obstetrics and gynecology clinics in Daegu was 31.9 years old [6]. This age difference of the infected women probably may originate from different patient’s group; target of this study was women (generally old women) who visited for health screening, while Goo et al. [6]’s subjects were younger women with genital problems who came to visit the obstetrics and gynecology clinics.

However, the increase in the prevalence of the condition with age is remarkable as it is not seen in other STDs [11]. Studies in the United States and Papua New Guinea found similar increases with age [12,13]. Moreover, Bowden et al. [11] reported a statistically significant increase in the age-specific prevalence of *T. vaginalis* but a significant decrease for *Chlamydia trachomatis* and human papillomavirus infections.

The increase of trichomoniasis with age may be associated with a drop in the number of lactobacilli, which are commensals of the vagina. Murta et al. [14] reported a decreased frequency of *Lactobacillus* species in individuals above 50 years, and lactobacilli are reported to inhibit the adhesion of *T. vaginalis* to host ectocervical cells [15]. In addition, in post-menopausal women, the gradual loss of glycogen and lactobacilli due to the drop in circulating estrogen may cause an increase in vaginal pH [16,17]. Therefore, the reduction in lactobacilli and increased vaginal pH that occur above 50 years together may reduce the ability of women to defend against *T. vaginalis* infection. This suggestion would be helpful for understanding the higher prevalence in old women in this study.

In summary, the *T. vaginalis* infection rate in 424 women attending for health screening in Guri city was 3.3%, and the rate was higher in those above 50 years old than those below 50 years. The infection rate may have increased since 15 years ago when the rate was found to be 2.4% in the same area. The use of PCR for diagnosing *T. vaginalis* in women over 50 years is recommended even in asymptomatic women.

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**Ethical Standards:** This study complied with the current laws of this country.

**CONFLICT OF INTEREST**

The authors declare that they have no conflicts of interest.

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