PREVALENCE OF LOWER GENITAL TRACT INFECTION AMONG WOMEN ATTENDING MATERNAL AND CHILD HEALTH AND FAMILY PLANNING CLINICS IN HANOI, VIETNAM

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Abstract. To determine the prevalence of lower genital tract infection (LGTI) with Candida spp, Trichomonas vaginalis, Neisseria gonorrhoeae, Chlamydia trachomatis, and bacterial vaginosis among symptomatic and asymptomatic women attending maternal and child health and family planning (MCH/FP) clinics in Hanoi, Vietnam. A multi-centered, cross-sectional descriptive study stratified by reported symptoms of vaginal discharge was carried out in three MCH/FP clinics among 1,000 women aged 18-44 years in 1998. Of these, 89.1% lived in Hanoi, 97.6% were currently married, and 99.2% had only one sexual partner in the past 12 months. Regarding their contraceptive use, 28.2% did not use any contraception, 25.6% used an intrauterine device (IUD), 22.8% used condoms, and 23.4% used other methods. The overall prevalence of Candida spp was 11.1% (95% CI = 9.1-13.1%); T. vaginalis, 1.3% (95% CI = 0.6-2.0%); no gonococcal infection was found; the prevalence of C. trachomatis was 4.4% (95% CI = 3.1-5.7%); and of bacterial vaginosis, 3.5% (95% CI = 2.4-4.6%).

INTRODUCTION

The intrauterine device (IUD) is the most commonly used contraceptive in Vietnam (Allman et al, 1991; Johansson et al, 1996). Complications of IUD use are more likely in the presence of lower genital tract infection (LGTI) (Hatcher et al, 1998), but data on the prevalence of these infections in Vietnam are very limited.

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To fill this gap, the Institute for the Protection of Mother and Newborn (IPMN), supported by the World Health Organization (WHO), carried out this study to establish the prevalence of LGTI in women seeking care at maternal and child health and family planning (MCH/FP) clinics in Hanoi. This study aimed to determine the prevalence of LGTI so that prevention measures could be introduced to provide more effective reproductive health care. The objective of this research was to estimate the prevalence of important LGTI organisms in both self-reported symptomatic and asymptomatic women who attend the MCH/FP clinics in Hanoi, Vietnam. We also describe the clinical findings of these patients.
MATERIALS AND METHODS

Study design and data collection

The study was of multi-centered, cross-sectional, descriptive design (Kelsey et al, 1986) and was conducted in three MCH/FP clinics in Hanoi, Vietnam. The first clinic was the Obstetrics and Gynaecology Hospital of Hanoi, with 200 beds and 15,535 annual out-patient visits (March to May 1998). The second was the Hai Ba Trung Maternity Hospital, with 40 beds and 1,600 patients attending its polyclinic (May to September 1998). The third was the Obstetrics and Gynaecology Department, Thanh Nhan Hospital, Hanoi, with 38 beds and 7,151 out-patient visits (July-December 1998). Originally it was planned that only the first two clinics would be used in the study, but, because of insufficient number of women recruited into the study, the third clinic was added.

A total of 1,000 women, aged between 18-44 years old, were enrolled with sampling quotas of approximately one-half symptomatic and one-half asymptomatic, using a multi-staged stratification design (Kish, 1965; Moser and Kalton, 1972; Kelsey et al, 1986). Systematic sampling (Kelsey et al, 1986) was used at the Obstetrics and Gynaecology Hospital of Hanoi, while all potentially eligible patients were recruited at the other two sites.

Women were excluded if they had taken antibiotics, or had used vaginal antimicrobial agents or vaginal douches in the previous two weeks before the visit. Those who were menstruating, pregnant, physically incapacitated or mentally deprived or had undergone hysterecctomy, were also excluded.

The research protocol was reviewed and approved by the institutional ethics review committee of the IPMN on behalf of the other participating institutions, and by the Scientific and Ethical Review Group of the Special Programme of Research, Development and Research Training in Human Reproduction of the WHO.

All potentially eligible patients were interviewed by a member of the nursing staff and informed consent was obtained. Patients were asked whether they had genital symptoms. They were then stratified into two groups, ie, symptomatic (vaginal discharge) and asymptomatic. A pre-coded structured interview form was used to obtain information from each enrolled patient. The initial interview consisted of questions about age, education, profession, marital status, symptoms of genital tract infection and their current usage of contraceptives. Each patient then underwent general physical and pelvic examinations. A speculum was used to observe any vaginal discharge and its texture, color, and odor. The cervix was inspected for cervical lesions including inflammation, ectopy, dysplasia or any sign of ulceration or papillomata. The uterus and the adnexae were palpated for evidence of pelvic infection.

The gynecologist collected four specimens, two from the posterior vaginal fornix, and two from the endo-cervical canal. The results of the interviews were not revealed to the gynecologist, and the results of the clinical examinations were not available to the laboratory staff who entered their results on separate forms. All patients with detected abnormal conditions were treated according to the local standard of care.

Diagnostic procedures

Vaginal swabs. Material from the first high vaginal swab was examined for its appearance and pH estimation using narrow-range indicator papers (Whatman, Maidstone, UK). Gram stain was used for the identification of Candida spp, clue cells, and Neisseria gonorrhoeae (Barron and Finegold, 1990). The second high vaginal swab was used as a wet smear to examine for Trichomonas vaginalis, yeast pseudohyphae, clue cells, and for the 10% potassium hydroxide solution (KOH) ‘whiff’ test.

The Gram-stained specimen was also examined for yeast or pseudo-hyphae under microscopy for 10-15 oil immersion fields. The gradings obtained were negative if no yeast cells were found, border-line if one or two yeast cells were found, and positive if more than two yeast cells were found (Odds, 1979). The diagnosis of bacterial vaginosis was made by using the Amsel criteria (Moser and Kalton, 1972; Amsel et al, 1983).
**Endocervical swabs.** Dacron-tipped swabs were inserted into the endocervical canal and rotated two to three times to be sure of obtaining endocervical cells. The first swab was introduced into the external cervical os and gently rotated. It was then plated directly onto Thayer-Martin medium (Difco, MI, USA) and incubated in a 5-10% CO₂ candle jar at 37°C for 48 hours and examined for *N. gonorrhoeae* (Koneman et al, 1988). The second swab was used to diagnose *C. trachomatis* using an enzyme immuno-assay kit (Chlamydiazyme, Abbott Laboratories, North Chicago, IL, USA) (Mazara et al, 1989). The direct detection of the chlamydial antigen was used and the positive tests were confirmed by the Chlamydiazyme blocking reagent kit. The order of use of the swabs were alternately allocated to either of these two tests.

**Data management and sata analysis**

All coded data were entered into computer twice and validated by using Epi Info Version 6.04 (Atlanta, GA, USA: Centers for Disease Control and Prevention, 1997) to minimize key-punch errors. Data were then tabulated and cross-checked for out-of-range errors and inconsistencies until they were considered clean. For bi-variable analysis, Pearson’s $\chi^2$ corrected for survey designs was used for categorical data analysis. Statistical inference on prevalence data was done by using the 95% confidence intervals, and odds ratios (OR) were used to measure the strength of association (Fleiss, 1981). We used Intercooled Stata version 6.0 for Windows 98/95/NT (College Station, TX, USA: Stata Corporation, 1999) statistical package. Kappa statistics ($\kappa$-statistics) were used to measure agreements between observers (Fleiss, 1981).

**RESULTS**

**Study population**

One thousand women were enrolled into this study out of 1,620 potential participants. One hundred and seventy women were excluded because of prior recent use of antibiotics or vaginal antimicrobial medication. Five hundred women (50.0%) were enrolled at the Obstetrics and Gynaecology Hospital of Hanoi, 286 (28.6%) were from the Hai Ba Trung Maternity Hospital and 214 (21.4%) were from Thanh Nhan Hospital. The mean age (± SD) of these 1,000 women was 34.1 ± 6.8 years; their mean age at their first marriage was 23.5 ± 3.5 years; and their age at first sexual intercourse was 23.4 ± 3.5 years. Of these women, 99.2% had had only one sexual partner in the past 12 months; 97.6% were currently married; 89.1% were living in Hanoi; 35.3% were laborers or farmers, 25.5% were office workers, and 39.2% were housewives or had other occupations (such as students); 35.6% attained primary school education or lower, 43.6% had secondary school education, and 20.7% attained above secondary school education. Regarding their contraceptive use, 28.2% did not use any contraception, 25.6% used an IUD, 22.8% used condoms, and 23.4% used other methods, such as periodic abstinence or coitus interruptus. A total of 504 women (50.4%) were symptomatic with vaginal discharge.

When the above characteristics between the two strata of symptomatic and asymptomatic were compared, the asymptomatic group was 1.3 years older than the asymptomatic group (p < 0.01). There were, however, no differences between strata in any of the other above-mentioned variables (p > 0.07 for any variables, data not shown).

**Clinical histories and findings**

Many women reported multiple complaints; 50.4% complained of vaginal discharge, 24.3% complained of vulva itching, 8.6% reported having had dyspareunia, 4.8% had a chief complaint of dysuria, 3.7% complained of genital ulcer, and 0.8% complained of non-menstrual vaginal bleeding. These complaints were not mutually exclusive. Of women who complained of having vaginal discharge on the current visit, 46.3% described the discharge as creamy, 9.2% as colorless, 9.0% as greenish-yellow, 4.2% as blood-stained, and 31.3% with different variable characteristics. More than half (55.1%) complained of having at least one episode of vaginal discharge during the past six months, not including the current episode; 39.9% had it treated. Of those who sought treatment, 81.7% were treated by medical doctors, 14.6% had treated themselves, and 3.7% had used other sources, such as traditional medicine.
On pelvic examination, 15.5% of women had abnormal vulva findings such as vulvitis or leukoplakia-like lesions; 50.2% with vaginal discharge were clinically classified as abnormal, 52.7% had abnormal vaginal discharge color, 45.1% had abnormal vaginal mucosal appearance, and 57.8% had abnormal cervical appearance, such as cervicitis or ectopy. Most (91.3%) had normal uterine size and 85% had a normal adnexal examination. Again, these pelvic examination results were not mutually exclusive. The bi-variable analyses of characteristics between the reported asymptomatic, versus the symptomatic, women showed six clinical findings associating with symptomatic conditions (p < 0.05), ie, presence of vulvitis, texture of vaginal discharge and its color, vaginitis, cervicitis, and cervical ulceration (Table 1). Of note, the major complaints of the patients did not agree well with the clinical findings (κ = 0.22, p < 0.001). Only 84.8% of those reported as having any symptoms had positive clinical findings, and 63.4% reported as having no symptoms had some positive clinical findings.

### Table 1

Comparison of pelvic examination results between asymptomatic and symptomatic Vietnamese women.

<table>
<thead>
<tr>
<th>Findings</th>
<th>Asymptomatic (N = 496)</th>
<th>Symptomatic (N = 504)</th>
<th>Odds ratio (95% CI)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>number</td>
<td>number</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Vulva</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vulvitis</td>
<td>No</td>
<td>455</td>
<td>408</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
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<tr>
<td>Vulvitis</td>
<td>Yes</td>
<td>41</td>
<td>96</td>
<td>2.6 (1.8-3.9)</td>
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<tr>
<td><strong>Vagina</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Texture of vaginal discharge</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>310</td>
<td>188</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>&lt;0.01</td>
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<tr>
<td>Homogeneous</td>
<td>81</td>
<td>162</td>
<td>3.3 (2.4-4.6)</td>
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<tr>
<td>Curdy</td>
<td>53</td>
<td>66</td>
<td>2.1 (1.4-3.1)</td>
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<tr>
<td>Yellowish-frothy</td>
<td>15</td>
<td>38</td>
<td>4.2 (2.2-7.8)</td>
<td>&lt;0.01</td>
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<tr>
<td>Other</td>
<td>37</td>
<td>50</td>
<td>2.2 (1.4-3.5)</td>
<td>&lt;0.01</td>
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<tr>
<td><strong>Color of vaginal discharge</strong></td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Normal</td>
<td>299</td>
<td>174</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>&lt;0.01</td>
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<tr>
<td>White</td>
<td>83</td>
<td>103</td>
<td>2.1 (1.5-3.0)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Yellow</td>
<td>100</td>
<td>209</td>
<td>3.6 (2.7-4.9)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Other</td>
<td>14</td>
<td>18</td>
<td>2.2 (1.1-4.6)</td>
<td>&lt;0.01</td>
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<tr>
<td>Vaginitis</td>
<td>No</td>
<td>320</td>
<td>236</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Vaginitis</td>
<td>Yes</td>
<td>176</td>
<td>268</td>
<td>2.1 (1.6-2.7)</td>
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<td><strong>Cervix</strong></td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Cervicitis</td>
<td>No</td>
<td>280</td>
<td>208</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cervicitis</td>
<td>Yes</td>
<td>216</td>
<td>296</td>
<td>1.8 (1.4-2.4)</td>
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<tr>
<td>Ulceration</td>
<td>No</td>
<td>466</td>
<td>455</td>
<td>1.0&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ulceration</td>
<td>Yes</td>
<td>30</td>
<td>49</td>
<td>1.7 (1.02, 2.8)</td>
</tr>
</tbody>
</table>

<sup>a</sup>Totals vary because of incomplete data; <sup>b</sup>Referent.

**Prevalence of lower genital tract infections**

The overall prevalence of *Candida* spp (more than two yeast cells found) was 11.1%. If we included the border-line cases, the prevalence of *Candida* spp would be 34.2% (95% CI = 31.2-37.1%). The prevalence of this infection was quite similar, both in the asymptomatic and the symptomatic group, as was the case for infection with *T. vaginalis*, *N. gonorrhoeae*, and *C. trachomatis*,
There were nine women (0.9%) who had more than one infection. Among these, one had Candida spp and T. vaginalis, three had Candida spp and C. trachomatis, three had T. vaginalis and C. trachomatis, and two had T. vaginalis and bacterial vaginosis. If the women with co-infections were excluded, the prevalence of any single LGTI was 18.7% (95% CI = 16.2-21.1%).

DISCUSSION

Our study provides prevalence data of five important infections of the lower genital tract in women attending maternal and child care clinics in the capital of Vietnam. These include the prevalence of Candida spp (11.1%), N. gonorrhoeae (0%), T. vaginalis (1.3%), C. trachomatis (4.0%), and bacterial vaginosis (3.5%).

Caution should be used in comparing our results with those of other studies of LGTI because those may have included populations with either low or high risk, eg, women attending ante-natal care clinics, female sex workers, women attending MCH/FP clinics, clients of men visiting STD clinics, etc. Also, there is variation in the settings, eg, in developing countries or in underdeveloped communities in developed countries, STD clinics, community surveys, MCH/FP clinics, etc. We thus choose to compare our data with selected study populations in less-developed communities carried out with a similar design. In addition, detected prevalences may differ depending upon the laboratory methods used, which also varied in these other studies.

Our results showed that the prevalence of Candida spp was similar to those obtained from Sudan (Kafi et al, 2000), Thailand (Thongkrajai et al, 1999), and rural Vietnam (Boon et al, 1999; Hung et al, 2001). The absence of N. gonorrhoeae is consistent with the findings of our pilot study in Hanoi (Anh et al, 1996), Thailand (Thongkrajai et al, 1999), and Turkey (Ortayli et al, 2001); a slightly higher prevalence (<1.0%) was found in indigenous women in Australia (Bowden et al, 1999), Indonesia (Iskandar et al, 2000), and northern Thailand (Kilmarx et al, 1998). The prevalence of T. vaginalis was much lower than that found in India (Vishwanath et al, 2000), Indonesia (Iskandar et al, 2000), Kenya (Feldblum et al, 2000; Fonck et al, 2000), or Thailand (Thongkrajai et al, 1999). Meanwhile, the prevalence of chlamydial infection was similar to that found in our pilot study in Hanoi (Anh et al, 1996), Thailand (Thongkrajai et al, 1999), and Turkey (Ortayli et al, 2001); a slightly higher prevalence (<1.0%) was found in indigenous women in Australia (Bowden et al, 1999), Indonesia (Iskandar et al, 2000), Kenya (Feldblum et al, 2000), and northern Thailand (Kilmarx et al, 1998), but was about ten-fold lower than that found in Jamaica (Dowe et al, 1999). Finally, the prevalence of bacterial vaginosis was much lower than found elsewhere (Dowe et al, 1999; Iskandar et al, 2000).

The low prevalence of condom use as found among women in this study, together with that reported in men who frequent commercial sex
workers in southern Vietnam (Thuy et al., 1999), and the low knowledge of reproductive tract infection among Vietnamese women (Thuy et al., 1999), raise concerns about the control of HIV/AIDS and other sexually-transmitted diseases in Vietnam.

Although the prevalence of LGTI was lower than in some other studies, some pathogen was still found in about one in five women. Many women also used the IUD, and an increased rate of complications of IUD use may be seen in those with concurrent LGTI (Hatcher et al., 1998). The inconsistencies found in reported symptoms, physical examination findings, and laboratory results, underscore the challenges of establishing syndromic diagnostic algorithms (Boon et al., 1999; Vishwanath et al., 2000). Low-cost, simple diagnostic tests for reproductive tract infections remain an urgent need.

This study has some limitations, in that it was carried out in the capital city of Vietnam and does not reflect the general population of Vietnam, especially in rural settings, but it does reflect the situation among women seeking health care in Hanoi. This study provides justification for future risk-factor studies in this area, as well as for prevention and treatment activities.

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