Clinical and laboratory profiles of genital ulcers (sexually transmitted diseases) in a tertiary care center in northeastern India

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ABSTRACT

Background: Genital ulcers are defined as breaches in the continuity of the genital mucosa and/or skin. Sexually transmitted infections (STIs) that cause genital ulcer disease (GUD) are syphilis, chancroid, donovanosis, lymphogranuloma venereum (LGV), and herpes genitalis. This study aimed to investigate the clinical and laboratory profiles of STI-related genital ulcers. Materials and Methods: A cross-sectional two-year study was conducted on patients attending the Outpatient Department of Dermatology, Venereology and Leprosy in a tertiary care center in northeastern India. Selected were 95 patients who presented themselves with STI-related genital ulcers. Detailed history taking and examination were conducted with basic tests to assist the diagnosis. Results: The male-to-female ratio was 3.32:1, and the most common site was the glans and prepuce in males (28.77%) and the labia majora and minora in females (36.36%). 96.84% of patients had superficial ulcers. The KOH mount was positive in 26 patients. The Tzanck smear was positive in 31 patients. RPR was positive in four. HIV was positive in eleven. Herpes genitalis (96.84%) was the most common GUD. Mixed STIs were attested in 41.05% of patients. Conclusion: GUD can take various forms of presentation. The available laboratory tests should be utilized. The possibility of mixed infections should always be kept in mind.

Key words: Genital ulcer disease; Herpes simplex virus; HIV

INTRODUCTION

Genital ulcers are defined as breaches in the continuity of the genital mucosa and/or skin. Genital ulcer disease (GUD) may be due to sexually transmitted infections (STIs), such as syphilis, chancroid, donovanosis, lymphogranuloma venereum (LGV), herpes genitalis, or non-STIs, such as traumatic ulcers, Behçet’s disease, lichen planus, erythema multiforme, lichen sclerosis et atrophicus, bullous diseases, Fournier gangrene, and squamous cell carcinoma [1].

Documenting the etiological agents of GUD remains especially difficult as the relevant diagnostic tests are often unavailable or are misused. The etiological agent may not be found due to the patient self-medicating or because of the contamination of the ulcers. However, bedside and laboratory tests are, in early lesions, fairly sensitive and specific. A clinical diagnosis may be misleading because the increasing number of HIV coinfections and mixed infections often alter the morphology of the ulcers and the textbook description of GUD may not always be present. The ulcers do not remain confined to the genitalia and may be seen in extragenital sites due to changes in sexual behavioral patterns. Clinical diagnoses have been observed to be incorrect in around 40% of patients with GUD when compared to laboratory tests [1].

In view of the above loopholes in the diagnosis and management of GUD, it is important that a clinical diagnosis is reinforced by the laboratory facilities.
available at the institute of examination so that the newer trends of manifestation, if any, can be recognized and so that any HIV-correlated peculiarity of GUD can be detected. An accurate clinical diagnosis corroborated by laboratory findings will prevent inflicting unnecessary psychological trauma to the patient.

MATERIALS AND METHODS

A cross-sectional study was conducted on patients visiting the Outpatient Department of Dermatology, Venereology and Leprosy in collaboration with the Department of Microbiology, Regional Institute of Medical Sciences (RIMS), Imphal, Manipur, for a two-year period from October 2011 through September 2013.

A total of 95 cases of both sexes and all age groups who presented themselves with STI-related genital ulcers at the Outpatient Department of Dermatology, Venereology and Leprosy, RIMS Hospital, were selected from among all cases displaying genital ulcers after excluding non-STI-related genital ulcers. Clinically diagnosed STI-related genital ulcers for which the minimum laboratory investigations could not be completed due to poor patient compliance were also excluded from the study group. Ethical approval for this study was received from the Institutional Ethical Committee, RIMS, Imphal, Manipur.

After receiving a written informed consent from each of the patients, a detailed history was taken and recorded in a uniform predesigned proforma, followed by a thorough local and systemic examination as well as investigations such as genital swabs for KOH, wet mounts, Gram stain, Giemsa stain, Tzanck smear, and blood samples for VDRL and routine tests, including urine examination, tests for HIV, HBsAg, anti-HCV Ab, serology for herpes simplex virus 1 and 2, and TPHA, wherever needed. The collected data was analyzed with descriptive statistics.

RESULTS

Among the total 95 cases studied, 73 (76.84%) were males and 22 were females (23.16%), giving a male-to-female ratio of 3.32:1. The most common age group was 21–30 years old (34.73%).

The most common presentation among male patients was painful genital sore (n = 22; 30.14%), the healthcare-seeking behavior in female patients was mainly due to pain (n = 9; 40.90%) (Table 1).

Most of the male patients (n = 52; 43.83%) presented themselves between day 6 and 11, in contrast to the female patients, among whom the majority (n = 14; 63.65%) reported earlier, between day 0 and 5 after the onset of the disease. The mean duration of the disease was 11.88 ± 13.299 days in males and 11.14 ± 16.994 days in females (Table 2).

The most common site in males was the glans and prepuce (n = 21; 28.76%) (Table 3), while, in females, most ulcers appeared in both the labia majora and minora (n = 8; 36.36%) (Fig. 1).

Speculum examination of cervical lesions was skipped so as not to unnecessarily cause more pain to a suffering patient.

Table 1: Distribution of symptoms among males and females

| Symptoms                                      | Male n (%) | Female n (%) |
|-----------------------------------------------|------------|--------------|
| Painful genital sore                           | 22 (30.14) | 9 (40.90)    |
| Itching                                       | 9 (12.33)  | 3 (13.63)    |
| Pain and itching                              | 6 (8.22)   | 0 (0.00)     |
| Pain and dysuria                              | 5 (6.85)   | 1 (4.55)     |
| Pain and discharge from the urethra or vagina | 1 (1.37)   | 5 (22.72)    |
| Asymptomatic rash around the urethral meatus  | 1 (1.37)   | 0 (0.00)     |
| Pain with genital swelling                    | 0 (0.00)   | 1 (4.55)     |
| Genital sore with groin swelling              | 1 (1.37)   | 0 (0.00)     |
| Genital sore with genital swelling            | 1 (1.37)   | 1 (4.55)     |
| Genital sore with growth                      | 1 (1.37)   | 0 (0.00)     |
| Asymptomatic genital sores                   | 14 (19.17) | 1 (4.55)     |
| Painless genital sore with dysuria            | 1 (1.37)   | 0 (0.00)     |
| Itching with dysuria                          | 2 (2.74)   | 0 (0.00)     |
| No complaints (partner examination)          | 1 (1.37)   | 1 (4.55)     |
| Skin eruption over the genitals               | 7 (9.59)   | 0 (0.00)     |
| Painful anal erosion                          | 1 (1.37)   | 0 (0.00)     |
| Total                                         | 73 (100)   | 22 (100)     |

Table 2: Distribution of the time of persistence of lesions upon presentation

| Duration (days) | Male n (%) | Female n (%) |
|-----------------|------------|--------------|
| 0–5             | 22 (30.14) | 14 (63.63)   |
| 6–11            | 32 (43.83) | 3 (13.64)    |
| 12–17           | 7 (9.59)   | 2 (9.09)     |
| 18–23           | 3 (4.11)   | 0 (0.00)     |
| >23             | 9 (12.33)  | 3 (13.64)    |
| Total           | 73 (100)   | 22 (100)     |

| Prepuce, frenulum | 1 (1.37) |
| Prepuce, shaft, corona | 1 (1.37) |
| Shaft             | 2 (2.74) |
| Periurethral area | 2 (2.74) |
| Perianal area     | 1 (1.37) |
| Total             | 73 (100) |
The majority of patients had 1–3 ulcers, 57.53% of males and 40.91% of females (Fig. 2). Among males, more than 16 (21.92%) displayed a single ulcer, which was the most common presentation. The most common shape of ulcer was round (n = 29; 30.53%). Ulcers had well-defined margins in 58 (61.05%) patients, ill-defined in 36 (37.90%), and undermined in one (1.05%). Ninety-two (96.84%) patients had superficial ulcers, while only three (3.16%) had deep ulcers. A majority (n = 67; 70.53%) had clean ulcers. The surface was more erythematous in 29 (30.53%) patients than in the others. Tenderness was present in only 34 (35.79%) patients and induration in 13 (13.68%). Associated vesicles were found in 36 (37.89%) patients, associated discharge from the vagina in 9 (40.90%), and associated urethral discharge in one male (1.05%). Inguinal lymphadenopathy was present in 25 (26.32%) patients, 16 males and 9 females, among which tenderness was present in nine males and four females (Table 4a).

There were no significant findings suggestive of microorganisms in 40 (42.10%) patients. HIV was positive in 11 (11.58%) patients; ten were positive and already on ART; one was recently diagnosed at the time of the study Table 4b. HCV antibody was positive in one HIV patient (1.05%) on ART, while there was not a single patient positive for HBsAg.

Table 3: Distribution of ulcer site in males

| Ulcer site                  | Number (%)       |
|-----------------------------|------------------|
| Prepuce                     | 16 (21.92)       |
| Corona                      | 3 (4.11)         |
| Prepuce, corona             | 2 (2.74)         |
| Glans                       | 15 (20.55)       |
| Glans, prepuce, corona      | 1 (1.37)         |
| Glans, shaft                | 3 (4.11)         |
| Glans, prepuce, shaft       | 1 (1.37)         |
| Glans, prepuce              | 21 (28.76)       |
| Glans, corona               | 3 (4.11)         |
| Prepuce, scrotum            | 1 (1.37)         |

Table 4a: Other clinical characteristics of genital ulcers

| Cutaneous examination | Male, n (%) | Female, n (%) | Total, n (%) |
|-----------------------|-------------|---------------|--------------|
| Ulcer margin          |             |               |              |
| Well-defined          | 40 (54.79)  | 18 (81.82)    | 58 (61.05)   |
| Ill-defined           | 32 (43.84)  | 4 (18.18)     | 36 (37.90)   |
| Undermined            | 1 (1.37)    | 0 (0)         | 1 (1.05)     |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Surface               |             |               |              |
| Clean                 | 53 (72.60)  | 14 (63.64)    | 67 (70.53)   |
| Slough                | 18 (24.66)  | 8 (36.36)     | 26 (27.37)   |
| Crusts                | 2 (2.74)    | 0 (0)         | 2 (2.10)     |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Depth                 |             |               |              |
| Superficial           | 70 (95.89)  | 22 (100)      | 92 (96.84)   |
| Deep                  | 3 (4.11)    | 0 (0)         | 3 (3.16)     |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Induration            |             |               |              |
| Present               | 8 (10.96)   | 5 (22.73)     | 13 (13.68)   |
| Absent                | 65 (89.04)  | 17 (77.27)    | 82 (86.32)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Tenderness            |             |               |              |
| Present               | 20 (27.04)  | 14 (63.64)    | 34 (35.79)   |
| Absent                | 53 (72.60)  | 8 (36.36)     | 61 (64.21)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Vesicles              |             |               |              |
| Present               | 22 (30.14)  | 14 (63.64)    | 36 (37.89)   |
| Absent                | 51 (69.86)  | 8 (36.36)     | 59 (62.11)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Inguinal lymph node   |             |               |              |
| Present               | 16 (21.92)  | 9 (40.90)     | 25 (26.32)   |
| Absent                | 57 (78.08)  | 13 (59.10)    | 70 (73.68)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
DISCUSSION

There are more than twenty pathogens transmissible through sexual intercourse but the exact magnitude of the burden of STIs is largely unknown due to the lack of in-depth studies, hindered by the stigma associated with STIs and by the lack of reinforcing clinical diagnosis with easy and quick diagnostic laboratory tests. In view of the rising prevalence of HIV/AIDS, proper control of STIs must be given due importance. It has been found that the presence of an untreated STI—ulcerative or nonulcerative—can increase risk of both the acquisition and transmission of HIV by a factor of ten [2]. A study by Desai et al. on the performance of STI syndromes against etiological diagnosis demonstrated that syndromic management based on genital ulcer syndrome (GUS) may miss around 85% of cases with active infection and lead to treatment of ulcers in 43% of cases in the absence of syphilis infection. Such a situation leads to the obvious financial burden of unindicated treatment while depriving the symptomatic cases of necessary care [3]. A study by Bailey et al. found that ulcers with bacterial or mixed etiologies were more likely to have detectable HIV-1 compared to those with HSV or with unknown etiologies, and concluded that

Herpes genitalis (n = 92; 96.84%) was the most common GUD, the second common being syphilis in 3 (3.16%) patients (Table 6).

Mixed STI infections, including those with HIV and HCV, comprised 39 (41.05%) patients. Among males, herpes genitalis was the most common GUD (n = 44). There were two cases of syphilis and one case per each of LGV and chancroid occurring as mixed infections with h. genitalis. In females, h. genitalis was the most common (n = 12) etiology of GUD, followed by h. genitalis with candidal vulvovaginitis (n = 7). However, there was no case of donovanosis (Table 6).

Table 4b: Laboratory tests

| Test                  | Male, n (%) | Female, n (%) | Total, n (%) |
|-----------------------|-------------|---------------|--------------|
| KOH                   |             |               |              |
| Positive              | 19 (26.03)  | 7 (31.82)     | 26 (27.37)   |
| Negative              | 54 (73.97)  | 15 (68.18)    | 69 (72.63)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| Tzanck smear          |             |               |              |
| Positive              | 19 (26.03)  | 12 (54.55)    | 31 (32.63)   |
| Negative              | 54 (73.97)  | 10 (45.45)    | 64 (67.37)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| RPR                   |             |               |              |
| Reactive              | 2 (2.74)    | 2 (9.09)      | 4 (4.21)     |
| Nonreactive           | 71 (97.26)  | 20 (90.91)    | 91 (95.79)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |
| HIV antibody          |             |               |              |
| Reactive              | 8 (8.22)    | 5 (22.73)     | 11 (11.58)   |
| Nonreactive           | 67 (91.78)  | 17 (77.27)    | 84 (88.42)   |
|                       | 73 (100)    | 22 (100)      | 95 (100)     |

Table 5: Gram stain findings (Gm+: Gram-positive; Gm-: Gram-negative)

| Gram stain                                      | Number (%) |
|------------------------------------------------|------------|
| Gm+ cocci singly                                | 10 (10.53) |
| Gm+ cocci in pairs                              | 6 (6.32)   |
| Gm+ cocci in groups                             | 6 (6.32)   |
| Gm+ cocci in groups and pairs                   | 3 (3.16)   |
| Gm+ cocci in groups and chains                  | 3 (3.16)   |
| Gm+ cocci and Gm+ coccobacilli                 | 1 (1.05)   |
| Gm+ cocci in chains                             | 1 (1.05)   |
| Gm+ coccobacilli in groups (rail track appearance) | 1 (1.05)   |
| Gm- bacilli                                     | 8 (8.42)   |
| Gm+ cocci and Gm- bacilli                      | 8 (8.42)   |
| Gm- bacilli and Gm- diplococci                  | 3 (3.16)   |
| Gm- diplococci                                  | 1 (1.05)   |
| Clue cells                                      | 3 (3.16)   |
| Polymorphs                                      | 1 (1.05)   |
| No significant findings                         | 40 (42.10) |
| Total                                          | 95 (100)   |

Table 6: Distribution of the diagnosis of the various GUDs

| Diagnosis                                      | Male n (%) | Female n (%) | Total n (%) |
|------------------------------------------------|------------|--------------|-------------|
| H. genitalis                                   | 44 (60.27) | 12 (54.55)   | 56 (58.95)  |
| Primary syphilis                               | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| Secondary syphilis                             | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| Secondary syphilis with candidal vulvovaginitis| 0 (0)      | 1 (4.55)     | 1 (1.05)    |
| Chancroid, h. genitalis                        | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| LGV with h. genitalis, HIV                     | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| H. genitalis with candidal balanoposthitis      | 16 (21.92) | 0 (0)        | 16 (16.85)  |
| H. genitalis, genital warts, BV, HIV            | 0 (0)      | 1 (4.55)     | 1 (1.05)    |
| H. genitalis, BV                               | 0 (0)      | 1 (4.55)     | 1 (1.05)    |
| H. genitalis, genital warts                     | 2 (2.74)   | 0 (0)        | 2 (2.11)    |
| H. genitalis, NGU, candidal balanoposthitis     | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| H. genitalis, candidal vulvovaginitis           | 0 (0)      | 3 (13.62)    | 3 (3.16)    |
| H. genitalis, candidal vulvovaginitis, HIV      | 0 (0)      | 4 (18.18)    | 4 (4.21)    |
| H. genitalis, NGU                              | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| H. genitalis, HCV, HIV                         | 1 (1.37)   | 0 (0)        | 1 (1.05)    |
| H. genitalis, HIV                              | 2 (2.74)   | 0 (0)        | 2 (2.11)    |
| H. genitalis, candidal balanoposthitis, genital warts, HIV | 2 (2.74) | 0 (0) | 2(2.11) |
| Total                                         | 73 (100)   | 22 (100)     | 95 (100)    |
the presence of lesional HIV-1 in almost one-half of HIV-positive males underscores the importance of GUD in increasing HIV infectivity [4]. Choudhry et al. suggested that syndromic management of GUD is not very effective in identifying herpes genitalis and syphilis. They also found a high incidence of HIV seropositivity (10.3%) in their study population, suggesting a close association of STI with HIV [5]. A study by Coovadia, Kharsany, et al. conducted in Durban, South Africa, demonstrated that, because of the lack of laboratory facilities in most STD clinics, microbial diagnosis is based on clinical criteria alone; their findings indicate that clinical and epidemiological features alone are not reliable in diagnosing the etiology of genital ulcers. Multiple ulcers, which are said to be characteristic of chancroid, were also noted in 36% (16/44) of patients with syphilis and in only 60% (24/40) of those with chancroid. The presence of buboes and the length of the incubation period proved to be of some help in diagnosing chancroid [6]. In another study, the accuracy of a clinical diagnosis in men was 66% in lymphogranuloma venereum (LGV), 65% in donovanosis, 42% in chancroid, 39% in genital herpes, 32% in primary syphilis, and 8% in mixed infections. In women, it was 94% in secondary syphilis, 83% in donovanosis, 60% in genital herpes, 58% in primary syphilis, 57% in chancroid, 40% in LGV, and 14% in mixed infections. Differences between clinical and laboratory diagnostic accuracies may reflect the similarities between the clinical appearance of the various causes of GUD, the presence of mixed infections, atypical ulceration due to longstanding disease, and insensitive laboratory tests [7].

Although there are more than 500 STD clinics in India, only 42 are functioning properly and collecting adequate records of cases. With the advocacy of the syndromic approach to the treatment of GUD and discharge, cases are diagnosed on the basis of flow charts, without laboratory aid, and are treated accordingly. This results in the provision of less information on HIV-positive cases among unregistered patients, which becomes a considerable social issue because HIV-positive cases remain unrecorded and become sources further propagating infection [8]. A study by DiCarlo and Martin demonstrated that the results of reaching an accurate clinical diagnosis of GUD were dependent on the clinician’s skills and are thus influenced by factors such as previous experience, knowledge of epidemiological risk factors, and disease prevalence, due to which the results may not be generalized. They found that comparing individual signs underlined statistically significant differences between ulcers in chancroid, genital herpes, and primary syphilis. Despite the statistical significance of these differences, however, there was a considerable overlap between the three groups [9].

Behets et al., in a study to determine the etiology of genital ulcers in Antananarivo, Madagascar, using a multiplex polymerase chain reaction (M-PCR) assay to detect DNA targets from Haemophilus ducreyi, Treponema pallidum, and herpes simplex virus in a single ulcer specimen and to assess the accuracy of clinical diagnosis and locally performed laboratory tests for comparison with the M-PCR results of 196 patients, found that syphilis and chancroid were clinically diagnosed in 42.9% of patients, syphilis in 36.7%, chancroid in 7.1%, LGV in 4.1%, syphilis and LGV in 4.1%, scabies in 4.1%, genital herpes in 0.5%, and multiple agents in 3.1% [10]. In HIV-infected patients, genital herpes can result in severe and atypical clinical presentations. Syphilis in an HIV patient appears to progress more rapidly through the clinical stages of syphilis and may often have an atypical clinical presentation or a refractory course after appropriate intramuscular penicillin, and lead to unusual serologic test results [11]. A study by Gopalkrishna et al. found that a majority of chlamydia-infected patients had syphilis (48%) besides other concomitant sexually transmitted infections, such as gonorrhea, genital herpes simplex type 2, condyloma acuminata, candidiasis, and bacterial vaginosis [12].

A study by Shaw et al. conducted in a rural Gambian community found that HSV2 positivity rose markedly with age in women while in men remained low in the younger age groups. In men, they found no association between HSV2 and the level of education or ethnicity [13]. In a study on rural Ugandans with genital ulcers, participants infected with HIV reported 26 episodes of genital ulcers per 100 person-years, compared with 7 per 100 person-years among those not infected [14]. A study by Sanchez et al. conducted in the Dominican Republic and Peru found that the clinical etiologic diagnosis of GUD was inaccurate and unsuitable as a basis for the management of genital ulcer lesions. Laboratory tests other than RPR are unavailable in many developing countries; even in industrialized countries, tests such as dark-field microscopy and H. ducreyi culture are troublesome to receive, and the latter is insensitive [15].
A study by Ahmed and Mbwana conducted in two Tanzanian cities found that HSV-2 was the most commonly identified agent in genital ulcer specimens with a high prevalence of HIV among STD patients. HSV-2 was detected at significantly higher rates among HIV-seropositive than HIV-seronegative patients with GUD [16]. A study by Renzi et al. on MSM (men who have sex with men) found a 1.7-fold increase in the risk of HIV acquisition among MSM who did not report herpes lesions or symptoms in the last twelve months and a low proportion of HSV-2-seropositive MSM who reported a history of genital herpes. Given the potential difficulty in recognizing reactivation of perianal herpes, they suggested that serological testing might be an important prevention strategy aimed at the identification of HSV-2-infected MSM [17].

One of the most critical drawbacks of diagnosing STIs through laboratory diagnostics is the need to send the specimens to a central reference laboratory for analysis, thus forcing the patient to return to receive the results. A significant number of patients do not return and are lost to follow up [18]. An STI study on a female Brazilian population by de Lima Soares et al. found that multiple infections were common and that 51% of women had at least one STI [19]. A study by Shepherd et al. conducted in Pune, India, found the cofactors in the susceptibility to HIV to include circumcision, urethritis, and genital ulceration [20]. Earlier, Zamzachin et al. conducted a study in the same setting and found the incidence of STI to be 3% of the total OPD visits for skin issues, while VDRL reactivity gave a median percentage of 8.49% and HIV prevalence among STI patients gave a median percentage of 8.12% [21]. A study by Shivswamy, Thappa, et al. conducted in Mumbai found that, among patients with clinical evidence of genital herpes, 94.2% were positive for both HSV-1 and HSV-2. In cases of the first episode of herpes genitalis, 66% were positive for HSV-1 and HSV-2, whereas in recurrent genital herpes, 96.4% were positive for HSV-1 and HSV-2. 72.6% of patients with no history suggestive of genital herpes were seropositive for HSV-2 [22].

In India, the main strategy aimed at achieving effective management for people with established infections has been to integrate STD services into the existing health care system, with a special emphasis on integration at the primary health care (PHC) level. The National AIDS Control Organization (NACO) for case management at this level recommends syndromic management [23]. A study by Reynolds, Risbud, Shepherd, et al. found that the first six months following the estimated time of exposure to syphilis is the period characterized by the greatest risk of HIV-1 infection compared to individuals who either have never had syphilis or who have had syphilis but more than six months have passed since the estimated time of syphilis exposure [24].

An evaluation by Creggan, Bauer, et al. of the relative sensitivities of the VDRL and TP-PA tests found that the sensitivity of the diagnostic strategy of VDRL confirmed by TP-PA was 71%. Substituting RPR for VDRL in a subset of 51 patients produced the same sensitivity. The sensitivity of TP-PA as the first-line diagnostic was 86% [25]. In a study on genital ulcers in women by Gomes et al., the most frequent etiology was herpetic lesion followed by autoimmune ulcers. Histological diagnosis was conclusive in only 26.4% of patients. They found, therefore, that the etiologic diagnosis of herpetic ulcers is difficult to reach, even if a variety of diagnostic methods are applied [26].

The treatment of GUD is often based only on the patient’s medical history, while physical examination is inaccurate. Tests specifically for the evaluation of genital ulcers include syphilis serology and either the DGI or DIF test for Treponema pallidum; the culture or antigen test for HSV; and the culture for Haemophilus ducreyi. Mandatory HIV testing should be performed on all patients with genital ulcers caused by T. pallidum or H. ducreyi and HSV. In most cases, health care providers based in developing countries must treat patients on the basis of clinical presentation and epidemiological circumstances since testing facilities are often unavailable [27]. A study by Richards, Krantz, et al. aimed to describe the healthcare-seeking behavior among patients with newly-acquired symptomatic genital herpes and found that the presence of exquisite tenderness as the chief complaint may impede the diagnosis in the absence of a careful genital examination, which underscores the need to use various laboratory methods, such as follow-up HSV serology, to confirm the diagnosis of HSV infection [28].

In studying the effects of genital ulcer disease and herpes simplex virus type 2 on the efficacy of male circumcision as HIV prevention, Gray, Serwadda, et al. suggested that ulcers prevented by circumcision are likely to be non-herpetic in origin. They also suggested that most of the reduction in the risk of HIV afforded by circumcision is attributable to the removal of vulnerable foreskin tissue that contains HIV target
cells [29]. Nath and Thappa showed the importance of Tzanck smear in the rapid diagnosis of genital herpes lesions by identifying multinucleate giant cells, although with less sensitivity than by viral cultures, and the feasibility of undertaking histopathology occasionally in cases such as a chronic herpes infection in HIV-infected individuals with atypical disease morphology and clinical course [30]. Since HIV shares numerous aspects with other STIs—including modes of transmission, behavioral and other cofactors, and potential control measures—HIV prevention can logically be situated within the larger encompassing domain of STI control [31]. In a study conducted in Brazil, Barbosa et al. verified the association between *C. trachomatis* infection and gonorrhea with urethral discharge, highlighting that STIs are often transmitted by the same route and treating them may reduce the incidence of HIV and other STIs, which would be an important step in prevention, planning, and care [32].

A prospective study of genital ulcer disease by Hope-Rapp et al. in Paris found primary syphilis to be the most common (35%) cause, followed by herpes genitalis (27%) and chancroid (3%). A high frequency of HIV infection (27%) was found particularly in patients with primary syphilis (33%). They stressed that no clinical characteristic is predictive of etiology, undermining the importance of performing a thorough microbiologic evaluation [33]. Not much data is available regarding the current status of STIs in India and how their epidemiological and clinical picture is being modified in the context of the ongoing HIV/AIDS epidemic. More care needs to be exercised in order to inculcate safe sex practices and, thereby, reduce the incidence of preventable STIs, since intercurrent STIs are known to hasten the progression of an HIV infection toward the stage of fully-developed AIDS [34].

In a study to determine the etiology of STIs in Maputo, Mozambique, Zimba et al. stressed that syndromic management (SM) guidelines need to be area-specific and based on the knowledge of the prevalence of etiological syndromes and their drug susceptibility. The overall prevalence of HIV-1 and HIV-2 infection was 55%. GUD was associated with a significantly higher HIV prevalence (p < 0.001) [35]. A prospective study by Bala et al. found that the frequency of biological false positive (BFP) reactions with the VDRL test was quite low (0.2%) and concluded that the TPHA test should be employed for the routine confirmation of a positive VDRL test irrespective of its titer, especially to accurately diagnose and confirm syphilis in cases with a titer less than 1:8. They further noted that some of the TPHA-positive cases with a VDRL result of less than 1:8 might have been treated cases of syphilis and, therefore, attempts should be made to differentiate between past and active infections [36].

The most common presentation among male patients was painful genital sores (n = 22; 30.14%), followed by asymptomatic genital sores (n = 14; 19.17%) and itching (n = 9; 12.33%). The healthcare-seeking behavior in female patients was mainly due to pain (n = 9; 40.90%), which were all herpetic cases, similarly to a study by Richards et al. [28]. Only one (1.37%) male patient with herpes genitalis produced clear urethral discharge. Internal dysuria was present in nine (9.47%) patients with herpes genitalis, out of which eight were males. Fourteen (19.17%) male patients, one patient with secondary syphilis and the others with herpes genitalis, and one (4.55%) female patient with herpes genitalis displayed asymptomatic GUD. Most (n = 32; 43.83%) of male patients presented themselves between day 6 and 11, most frequently on day 7 (n = 24; 32.88%), among which the majority (n = 23) were herpetic. This was in contrast to females, among which the majority (n = 14; 63.63%) reported earlier, between day 0 and 5 of the onset of the disease. The mean duration of disease was 11.88 ± 13.299 days (mean ± SD) in males and 11.14 ± 16.994 days in females with an overall mean of 11.71 ± 14.145 days, which is almost identical to a study by Hope-Rapp et al. [33]. Pain in herpes genitalis reaches its maximum intensity during day 7–11 [37].

The most common site in males was in the glans and prepuce (n = 21; 28.76%). In females, most ulcers appeared in both the labia majora and minora (n = 8; 36.36%), which may be due to the greater contact of these sites during sexual contact. A study by Gomes et al. found that a majority of female genital ulcers were in the midline near the clitoris or vaginal introitus [26].

Among male patients, more than 16 (21.92%) displayed a single ulcer, which is the most common presentation, similarly to what was observed by Hope-Rapp et al. [33]. Tenderness was present in only 34 (35.97%) patients, which may be due to the superficial nature of most of the ulcers. Associated vesicles, mostly minute, were found in 36 (37.89%) patients, almost identically to a study by Sanchez et al. [15], and associated discharge from the vagina in nine (40.90%) patients. Inguinal lymphadenopathy was present in 25 (26.32%) patients, which again mirrors other studies [10,15].
The KOH mount was positive in 26 (27.37%) patients, 19 males and 7 females, highlighting the importance of attending not only to the complaint at hand but also to the surrounding areas in order to rule out the possibility of other associated STIs. Multiple infections were also found in a study by de Lima Soares [19]. Tzanck smear was positive in only 31 (32.65%) cases, comparably to a study in Manaus, Brazilian Amazon, in which it was positive in 30.3% of GUD patients [38]; this, in turn, may be due to the fact that most of the patients presented themselves late, with a mean duration of 11.71 days. Besides, as described above, associated vesicles, mostly minute, were present in only 36 patients, which may likewise contribute to the lesser positivity of Tzanck smear. Indeed, Tzanck smear has a sensitivity of only 50% [39], may be negative in older lesions, and is inapplicable to dry lesions and crusts [40].

The RPR test was positive in four (4.21%) patients, among which TPHA was positive in three patients. One female patient with herpes genitalis, who was TPHA-negative, had a titer of as high as 1:64, which may be explained by the fact that acute biological false positive reactions are caused by herpes genitalis [41] and other viral infections [42]. The seroprevalence of syphilis in our study was 0.35%, which is quite low compared to the 3.5% of a study in Delhi [36]. This may be due to the widespread use of penicillin injection for all infections in this part of the country, which may have contributed to the relatively low prevalence of syphilis and other bacterial STIs as well. Histopathological examination was not deemed the gold standard for diagnosis in one study of female genital ulcers, where a low accuracy rate was found [26]. However, biopsies of genital ulcers might be helpful in identifying the cause of unusual ulcers or those that do not respond to initial therapy [43].

Herpes genitalis (n = 92; 96.84%) was the most common GUD in our study, similarly to other studies in different parts of the world [5,26,37,39,44,45]. The second most common cause was syphilis in three (3.15%) patients. In most countries around the world, syphilis ranks as the second most frequent cause of GUD, with a worldwide prevalence of 2–25% [16,39]. Hontelez et al. [46] opined that, since *T. pallidum* is susceptible to the commonly available antibiotics, increased antibiotic exposure would be expected to have a secondary effect on the prevalence of syphilis. In one study, primary syphilis was found to be the most frequent etiology of genital ulcerations in MSM [33]. There was only one (1.05%) case per each of chancroid and LGV, both as mixed infections with herpes genitalis. Mixed etiologies were also observed in other studies [19,32]. There was no case of donovanosis in our study, which may be due to its low prevalence in this region, whereas it is endemic along the east coast of Orissa, Andhra Pradesh, and Tamil Nadu [47]. The decrease in the prevalence of bacterial GUD, particularly chancroid, may be due to the improved bacterial STD management as well as the improved HSV-2 diagnostics, which may suggest a relative increase in the role of HSV-2 as a cause of GUD [16].

**CONCLUSION**

GUDs can have various modes of presentation with varied symptomatology, while atypical cases are a diagnostic dilemma. GUDs should never be presumed to be due to a single etiological agent, and all available laboratory facilities within reach should be utilized as far as practicable.

**Statement of Human and Animal Rights**

All the procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the 2008 revision of the Declaration of Helsinki of 1975.

**Statement of Informed Consent**

Informed consent for participation in this study was obtained from all patients.

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