Case Report
Management of vulvar Paget’s disease: a case report and review of the literature

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Abstract: Extra-mammary Paget disease is one of the rare neoplastic conditions of the skin. The most common site of involvement is the vulva and presents itself with erythematous plaques. Although surgery is the most common treatment, removing the disease completely is challenging. We reviewed the reports of other patients with vulvar Paget disease and reported one new case. A 64-year-old woman presented to department of gynaecology with complaints of vulvar pruritus, painful vulvar lesion for 10 years. Physical examination revealed an erythematous plaque with a size of 15 cm*18 cm*20 cm extending over the pubic area to crissum. Pathological result was reported as Paget disease. The radical vulvectomy was performed with a safety margin of 2 cm. The patient was followed up for 2 years. No evidences of recurrence were observed throughout this period.

Keywords: Vulva, Paget disease, surgery, vulvar Paget disease

Introduction

Paget’s disease was first described in 1874 by Sir James Paget as a rare intraepithelial neoplasm of the areola skin [1]. In 1901, Dubreuil described the characteristic ‘cake-icing appearance’ of vulvar Paget’s disease [2]. Vulvar Paget’s disease is an extremely rare clinical entity, accounting for only 1% of vulvar malignancies. Our understanding of this disease is limited [3]. Paget’s disease of the vulva (PDV) commonly develops in postmenopausal Caucasian females. Symptoms include pruritus, tenderness or identification of a vulvar lesion [4]. The patients always suffer from these symptoms for several years before seeking medical advice. The typical clinical manifestations presents as a well-demarcated, thickened, pruritic, erythematous, or white scaly plaque with irregular borders.

Wilkinson and Brown has proposed a classification system for extramammary vulvar Paget’s disease based on the origin of neoplastic paget cells: -primary and secondary disease. Primary cutaneous Paget’s disease, an intraepithelial adenocarcinoma, arises within the epidermis or underlying skin appendages. Secondary or noncutaneous Paget’s disease originates from an underlying noncutaneous adenocarcinoma, most commonly anorectal adenocarcinoma, urothelial carcinoma of the bladder or urethra, carcinoma of the cervix, ovary or endometrium [5, 6]. A biopsy can be used to establish the diagnostic, and immunohistochemical studies may be of great use in distinguishing primary and secondary lesions [7].

Standard treatment of vulvar Paget’s disease is surgical excision [8]. Surgical evaluation of these lesions is important, because 20-30% of cases have an underlying malignancy [9]. Histological disease may exceed far beyond the visible lesion and lead to recurrence. Repeated operations may destroy the anatomy. So it is challenging to excise the disease adequately.

We reported a postmenopausal woman with noninvasive PDV. She was treated by only surgery and followed up for two years. No evidences of recurrence were observed throughout this period. We will show and discuss the treatment of this disease.

Case report

Sixty-four years old female was in natural menopause for 14 years. She visited the gynecologi-
cal oncology clinic in November 2012 with an itchy lesion in her vulva for 10 years. The symptom began about 10 years prior to admission with the appearance of a small pimple on the right labia majora. The marked pruritus was associated with the lesion. The patient received cortisone and ACTH, both locally and parenterally, and got temporary improvement. The vulvar pruritus and painful vulvar lesion was worse during three months.

Pelvic examination showed that the mons pubis and labium were markedly reddened and edematous. The involved area extended posteriorly over the perineum and almost to the anal margin and laterally to the right thigh for a distance of about 4.5 cm from the labium. It also extended up to the clitoris but did not approach the external urethral meatus (Figure 1).

Biopsy from the lesion was performed. The result of the biopsy was Paget’s disease (Figure 2). Upon the result of Paget disease from the pathology, other systems were examined. Breast examination, mammography results, skin examination and lymph node palpation results were normal. Colonoscopy, cystoscopy and pelvic and abdominal ultrasonography were performed and no abnormality could be found. Physical examination revealed an elderly female in good general physical condition. Heart and lungs were clear. There was no adenopathy.

The proposed treatment was surgery with radical vulvectomy and reconstruction via skin flap transplantation. The gross lesion was outlined with a 2 cm border, and strips of tissue beyond this border were excised. This was done in a systematic fashion with proper labeling by quadrants. These strips of tissue were sent directly to the pathology laboratory and processed by frozen section analysis to determine whether invasive disease is present. The routine fashion for vulvar malignancies and no special stains, molecular or receptor analyses were routinely performed.

The patient was followed up for 2 years. No evidences of recurrence were observed throughout this period. The pain disappeared and the quality of life was improved greatly.

Discussion

Paget’s disease was originally described as a breast lesion associated with underlying invasive ductal adenocarcinoma. It was divided into mammary or extramammary disease. According to the location on the body. The most common site of involvement is the vulva, which accounts for up to 60% of primary extramammary. Paget’s disease of the vulva cases account for less than 2% of vulvar malignancies [10]. Erythematous and/or eczematoid area are always accompanied by pruritus in 70% of patients [11]. The lesions in PDV are non-specific. There is a significant delay between the onset of symptoms and diagnosis [12]. The median interval from the onset of symptoms to a histological diagnosis was approximately 20 months [13]. This delay did not correlate with the size or extent of disease, which suggests that this disease is slowly progressive.
Concerning the pathogenesis, it is still a subject of great debate. Paget’s cells are malignant keratinocytes and are transformed in situ. Ellis et al. demonstrated that it was possible that in PDV, cells can migrate and progress to invasive disease by utilizing the existing vasculature, but not formation of new blood vessels [14]. Paget’s cells are larger than keratinocytes, have clear chromatin, and have a nucleolus. The cell cytoplasm is gray-blue, and it appears vacuolated with hematoxylin and eosin staining. On the basis of pathologic diagnosis, the patients were then divided into four groups: Intraepithelial Paget’s disease, invasive Paget’s disease, intraepithelial Paget’s disease with underlying adenocarcinoma and intraepithelial Paget’s disease with a coexisting cancer [15]. The intraepithelial type accounts for the majority of primary cases and has a better clinical course and outcomes.

Underlying adenocarcinoma is found in 10% to 20% of cases [16]. Helwig and Graham reported a higher rate of underlying adenocarcinoma in patients with perianal involvement [17]. Stephen Tebes et al. reported 26% of the women had underlying adenocarcinoma, and twenty-two percent of cases had additional malignancies at other sites, which is consistent with other reports [18]. Commonly associated malignancies mentioned in previous studies are breast, basal cell, rectal, genitourinary, and cervical carcinoma. Women with Paget’s disease should undergo a search for other malignancies, including at least colonoscopy, mammography, and Papanicolaou smear. Any suspicious symptoms of other malignancies should be evaluated additionally.

Paget’s disease of the vulva is a rare locally recurrent chronic disease. The recurrence rate is high, up to 21% to 61% [19]. It may correlate with a patchy tape of distribution with multiple islands of abnormal epithelium, which are not connected and have microscopic extension beyond the gross lesion. The risk factors associated with local recurrence remain controversial, such as a positive surgical margin, dermal invasion, and adnexal involvement.

Standard treatment of vulvar Paget’s disease is surgical excision, depending on the lesion size, lesion location, the presence of invasion, and underlying adenocarcinomas or adjacent skin abnormalities. Surgical procedures include wide local excision, radical hemivulvectomy, and simple or radical vulvectomy. The treatment of noninvasive extramammary Paget’s disease is wide surgical excision with a gross margin of 2-3 cm and resection to the fascia, because the disease usually extends well beyond the gross lesion. Radical vulvectomy may be considered for invasive lesions and adnexal adenocarcinomas. To reduce the width of the margin, mapping biopsies were performed at several selected points where the margin remained clinically ill-defined after infection control. The lesion can then be excised for frozen section analysis to determine whether invasive disease is present and whether radical resection will be necessary. Limiting resection in favor of the preservation of the clitoris, urethra, and anus is recommended.

Recurrences happened in 20% to 60% cases after surgical treatment [20], and microscopic positive margins have been frequently found in 40%-70% cases [21]. So margin status gained interest with intraoperative frozen section analysis. However, the controversy of influence of surgical margins on the recurrence rate also exists. Some studies found that positive margins correlate with increased recurrence rate, and negative margins tend to have longer disease-free periods and lower recurrence rates. However, other studies have found no correlation between margin status and local recurrence [22]. Gunn et al. showed that microscopic disease extended beyond the visible lesion with multifocal islands of microscopic disease [23]. Adamsons et al. also reported that Paget’s cells existed along the basement membrane beyond the apparent border of the lesion [24]. So the high frequency of recurrence and the high rate of positive surgical margins are thought to be a result of the multicentric nature of vulvar Paget’s disease. Because it is unfeasible to freeze all margins and intraoperatively identify sporadic cells. And there was no significant difference in recurrence rate between positive and negative surgical margins. We would recommend selective use of intraoperative frozen section analysis of the vulvectomy specimen, not for margin status, but to determine whether invasive disease is present.

Mohs micrographic surgery (MMS) is performed under microscopic control to remove tumor lesions and conserve tissue loss and function.
Mohs micrographic technique is used by the dermatologists as a standard surgical technique for the cutaneous lesions of extra-mammary Paget disease. A lower recurrence rate in those patients with Mohs surgery than that in the patients who were treated with conventional resection has been documented [25]. Hendi et al. reported a recurrence rate of 16% for the primary disease and 50% in recurrent cases [26]. However, additional clinical trials are warranted to determine whether MMS could be used as a standard surgical technique for EMPDV.

Alternative nonsurgical or combined treatments have been proposed, including radiotherapy, topical chemotherapy, laser, photodynamic treatment with aminolevulinic acid, topical fluorouracil (5-Fu), and imiquimod 5% cream [27]. Systemic chemotherapy may be considered in advanced cases of disease and in cases of metastatic disease although the standard regimens have not been validated. Radiotherapy is an alternative treatment for patients with a medical contradiction for surgery and for patients with recurrent disease. In addition, it could be considered as an adjuvant therapy for patients with a positive margin after surgical. Over the years, many therapeutic modalities have been attempted on patients with PDV in an effort to reduce the significant morbidity associated with the after-radical surgical treatments performed.

The high frequency of recurrence remains the most challenging feature in the management of vulvar intraepithelial Paget’s disease. Long-term monitoring of patients is recommended, and repeat surgical excision is often necessary. The optimum treatment regimen needs to be defined in prospective studies and long-term follow-up data are needed.

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Disclosure of conflict of interest

None.

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