INTRODUCTION

True cystic lesions of the vagina (excluding cysts arising from the urethra and surrounding tissues) are uncommon and they are usually an incidental finding during routine gynecological examination. Vaginal cysts are mainly classified according to their histology: 1) squamous inclusion cysts lined by stratified squamous epithelium more commonly located on the posterior wall, closer to an episiotomy repair or other surgical procedures; 2) mesonephric or Gartner’s duct cysts lined by non–mucin-secreting low cuboidal cells usually located near the anterolateral wall of the vagina following the route of the mesonephric duct; 3) Mullerian or paramesonephric cysts lined by tall columnar mucin-secreting cells located anywhere in the vagina and grossly indistinguishable from mesonephric cysts; and 4) Bartholin’s gland cysts arising from the Bartholin’s gland duct located near the opening of the Bartholin’s gland into the vestibule. Among them, Mullerian cyst constitutes 30%, Bartholin duct cyst 27.5%, epidermal inclusion cyst 25% and remaining 17.5% is constituted by Gartner duct cyst, endometriotic cyst and unclassified type. These cysts can vary in size and cause a variety of symptoms including a palpable mass, pain, dyspareunia, and voiding dysfunction. We describe a case of a woman with a vaginal cyst diagnosed during a routine gynecological examination.

CASE REPORT

A 20-year-old nulliparous woman presented at her gynecologist for a routine appointment. She had no relevant personal antecedents and denied any abnormal uterine bleeding, urinary or gastro-intestinal symptoms. At gynecological examination a soft, mobile cystic lesion on the posterior vaginal wall was found. There was no cough impulse on the mass and no pain was elicited by the examination. External genitalia was normal.

The patient underwent surgical excision of the cyst under general anesthesia (Figures 1 and 2). A small transverse incision was made on the posterior vaginal wall, which was separated from the cyst. During sharp and blunt dissection the cyst got accidentally ruptured and mucinous material drained out. The cyst wall was stripped away from vaginal wall and the cyst was total-
ly removed, excess of vaginal mucosa was excised and the incision was closed with absorbable sutures. Patient had an uneventful postoperative period and was discharged three hours after surgery.

Histological examination revealed a cyst lined by a single layer of tall columnar cells with mucin at the apex. The epithelium stained positive with mucicarmine and periodic acid-Schiff, confirming the clinical impression of a Mullerian cyst.

On her follow-up reviews, one month and 6 months after surgery, the woman had no complaints, with a normal gynecological examination.

**DISCUSSION**

The vast majority of vaginal cysts are asymptomatic and require no treatment. Symptoms may vary according to cyst size, nature and location – patients may complain of vaginal discomfort, vaginal pressure, cyclic pain, dyspareunia, vaginal bleeding, and urinary or rectal symptoms.

In our patient, differential diagnosis of a rectocele and enterocele were considered and ruled out by rectal examination, cyst location and by absence of cough impulse. The location of the cyst also ruled out a Bartholin cyst and the absence of cyclic pain and nature of the cyst excluded an endometriotic cyst. Inclusion cysts were excluded by absence of previous perineal lacerations or pregnancies. Due to its location, the most probable diagnosis was of a Mullerian cyst although a Gartner duct cyst was still a possibility.

Mullerian cysts are congenital cysts of vagina, usually single but occasionally multifocal. During replacement of Mullerian epithelium with squamous epithelium of the urogenital sinus, Mullerian epithelial tissue can persist anywhere in the vaginal wall. As a consequence, cysts can be found almost anywhere within the vaginal walls, being more commonly found on the anterolateral vaginal wall. Cysts derived from the Mullerian ducts may exhibit histological patterns corresponding to those of any of the tissues normally derived from this duct, principally endocervical (the most common), tubal, or endometrial. Occasionally, a Mullerian cyst may become sufficiently large to be symptomatic and warrant excision.

Gartner’s duct cysts are less common than Mullerian cysts, are almost always located along the anterolateral wall of the vagina and typically are small, with an average diameter of 2 cm. Gartner’s duct cysts can also be associated with abnormalities of the metanephric urinary system such as an ectopic ureter, unilateral renal agenesis, and renal hypoplasia.

Although clinically irrelevant, Gartner’s duct cysts may be distinguished from Mullerian cysts histologically by the presence of a basement membrane and smooth muscle layer. However, clear distinction between the two can be made only on the basis of histo-
chemical staining, with paramesonephric cysts being periodic acid–Schiff and mucin positive, whereas mesonephric cysts are devoid of cytoplasmic mucin or periodic acid–Schiff positive material.

Vaginal cysts are treated via surgical excision. The technique is similar for all cysts and the entire cyst wall must be removed in order to prevent recurrence.

In conclusion, this is an unusual case of a posterior vaginal cyst, an uncommon location for a Müllerian cyst. When evaluating a vaginal cyst, assessment of the lesion via history and pelvic examination is important for the differential diagnosis and decision making regarding treatment. When symptomatic, excision of the cyst is a simple procedure with total relief of symptoms.

REFERENCES

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