Fibroepithelial polyp of vagina in a two-year-old girl and review of the literature

Alev Suzen¹, Nazile Erturk¹, Ozgur Ilhan Celik²

¹ Department of Pediatric Surgery, Faculty of Medicine, Mugla Sıtkı Kocman University, Mugla, Turkey
² Department of Pathology, Faculty of Medicine, Mugla Sıtkı Kocman University, Mugla, Turkey

Abstract
Fibroepithelial polyp of vagina (FEPV) is a mucosal polypoid lesion with a fibro vascular stalk and oedematus or hypocellular stroma covered by squamous epithelial surface. FEPV is extremely rare before menarche and after menopause. This article describes a two year old girl with a vaginal polyp and the histological analysis of which revealed to be a FEPV. Surgical excision of the mass was performed, and in 1 year of follow-up no recurrence was detected. A review of the literature showed that there were four cases of FEPV in neonatal girls. While these reported cases were neonatal, our case is the first case of FEPV in a two year old girl.

Keywords
Child; fibroepithelial polyp; vagina.

Introduction
A fibroepithelial polyp of vagina (FEPV) in an infant girl is extremely rare [1]. Common interlabial masses in infants and little girls include prolapsed ectopic ureterocele, prolapsus of urethra, paraurethral Skene cyst, hydro (metro) colpos, botryoid sarcoma of the vagina and FEPV [2,3]. We have to distinguish the origin (genital or urological) of the interlabial masses with good clinical examinations [3].

Case report
A two-year-old girl came to our hospital because of a vaginal polypoid mass which was discovered by her parents. She had no other systemic problems. The vaginal mass was discovered 6 months ago by her parents
and the parents said that vaginal mass was disappearing intermittently. On our first physical examination at the outpatient clinic, there was no vaginal mass as the vaginal mass entered inside the hymenal ring. When the mass came out of the hymenal ring, physical examination revealed a 1.2x0.5 cm polypoid mass, with a tall neck, originating from superior vaginal wall [Fig. 1].

The other urogenital examination was normal. Palpation showed it was painless and there were no signs of bleeding and inflammation.

Surgical excision of the vaginal polyp was performed and great attention was paid to preservation of the hymeneal ring. Histopathological findings showed a fibroepithelial polyp rich in vascular strictures [Fig. 2]. After 1 year of follow-up, the patient has shown no recurrence.

**Fig. 1.** A 1.2x0.5 cm polypoid mass, with a tall neck, originating from vaginal wall.

**Fig. 2.** Fibroepithelial polyp of vagina with a fibrovascular stroma containing congestive vascular structures covered by benign epithelial surface, (H&E x40).

**Discussion**

A FEPV is a polypoid lesion with a central fibrovascular core covered by squamous epithelium. Stroma may be oedematous, hypo cellular, or mixoid [4, 5]. FEPV is rare and is mostly seen in adult women or neonates. It is suggested that FEPV occur due to hormonally induced hyperplasia of subepithelial connective tissue or end stage of granulation tissue as a result of local injury. It is a benign procedure. During pregnancy, these lesions may be enlarged [4, 5].

Hymenal polyps are seen in the neonatal period and early childhood. They are probably due to oestrogen stimulation in the fetal period. Most of them disappear
spontaneously within a few weeks. Hymenal polyps originate from vaginal hymenal ring [6]. As polypoid formation in our case originates from inside of the hymenal ring, we eliminated hymenal polyp. Tubulo-squamous polyp of vagina has been described recently as a rare benign lesion. It is histologically characterized by both epithelial and mesenchymal component, embedded in a fibrous stroma [7]. It has not been detected in a child in the literature. After reviewing the literature, we found four cases of a FEPV in neonates: In 1990, Pul et al [8], in 2008 Jallouli et al [9], in 2010 Calleja et al [3], in 2015 Abdulhakim et al [1]. Our case is the first case of FEPV in a two year old girl. A FEPV should be considered in interlabial masses in prepubertal girls [8]. Therapy involves excision of the polyp [9]. As, the presentation of sarcoma and polyp could be similar, histological analysis is important [3]. Although it is benign and rarely recurs, it must be eliminated from botryoid sarcoma by histopathological examination [1, 3].

Conclusion
A FEPV is most seen in adult women. A few neonates were reported as a FEPV, but like in our case FEPV should be regarded in differential diagnosis of the interlabial masses at any age.

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References
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