Case Report of a 12-Year-Old Girl with Imperforate Hymen—Review of the Literature on Incision Techniques and Further Treatment of Patients

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Abstract

Background: Imperforate hymen (IH) is formed if no canal is created between the vaginal plate and the urogenital sinus. Treatment of IH is surgical. Different types of incisions and postoperative treatment are proposed. Case: A 12-year-old girl presented with a 3-day history of pain in the lower abdomen. Examination of the external genitalia disclosed a bulging hymen. Ultrasonography examination of her lower abdomen revealed hematocolpos. A diagnosis of hematocolpos caused by IH was established and surgical treatment was decided. Conclusion: Treatment of choice for IH is undeniably surgical. Although different types of procedures have been proposed, the surgeon should choose the one that best suits the patient’s needs without harming anatomical structures closely related to the hymen.

Keywords

Imperforate Hymen, Hematocolpos, Hymenectomy, Incision Techniques

1. Introduction

The hymen is a membrane located in the vaginal opening at the level of the introitus and is composed of urogenital sinus epithelial connective tissue [1] [2] [3] [4] [5]. During the final weeks of the embryologic development, a canal is created between the vaginal plate and the urogenital sinus [1] [2] [4] [6] [7] [8]. An imperforate hymen (IH) is a congenital disorder whereby the hymenal membrane covers the entire vaginal opening, thus causing genital outflow obstruction which leads to hematocolpos. The condition is mostly detected in ado-

2. Case

A 12-year-old girl presented to the emergency department of our hospital with a 3-day history of pain in the lower abdomen. No fever was reported. She had been admitted to the pediatric ward of our hospital 2 days earlier owing to urinary retention. During her hospitalization, a Foley catheter was inserted and urine flow was restored. Routine and culture urine samples were taken, the results of which were normal. The girl was discharged after 24 hours. No menstruation was noted. The rest of the girl’s medical history was unremarkable.

Physical examination of the patient revealed pain in the lower abdomen. Giordano’s sign was negative on both sides. Secondary sexual characteristics were found to be normal. Examination of the external genitalia disclosed a bulging hymen.

An ultrasound examination of her lower abdomen revealed hematocolpos (Figure 1). The rest of the internal genitalia showed normal development in accordance with her age. No further imaging examinations were performed. No pathological parameters resulted from blood and urine test.

A diagnosis of hematocolpos caused by IH was established and surgical treatment was decided. Surgery was performed under general anesthesia with the patient placed in the lithotomy position. The external genitalia area was sterilized and a Foley catheter was placed in the urethra so as to prevent it from being damaged (Figure 2). An X-shaped incision of the hymen was made and 600 ml of dark fluid was slowly drained. No suction was used. Four stitches were applied at the rim of the remaining hymen to ensure adequate haemostasis. The Foley catheter was immediately removed following completion of the procedure. The girl urinated normally after surgery and no postoperative antibiotics or estrogen ointment was prescribed. Sterilization of the external genitalia with Betadine solution three times per day was recommended for the first 24 hours. She was discharged from hospital the following day. The follow-up period was uneventful. Her menstruation was normal and no reformation of the hymen has been reported to date. It was not possible to assess possible sex dysfunction since the patient was not yet sexually active.

3. Discussion

IH is the most common congenital obstructive anomaly of female genital tract
Figure 1. Ultrasound examination of the lower abdomen revealing hematocolpos.

Figure 2. Patient placed in the lithotomy position for the surgery. A Foley catheter is placed in the urethra so as to prevent it from being damaged. A bulging hymen is obvious.

It is usually detected in puberty due to hematocolpos [3] [4] [6] [12] [13]. Although cases of infants have been reported with hydrocolpos secondary to IH [3] [4] [9] [15], misdiagnosis of IH is not uncommon, as occurred in our case. Similar symptoms are common in low transverse vaginal septum [14], obstructed hemivagina and ipsilateral renal agenesis [16]. Delayed diagnosis of IH can cause several complications, the most frequent of which is endometriosis [2] [4] [6] [13], while less frequent complications include retrograde menstruation, pelvic adhesions, fallopian tube damage, infertility [16] [17]. Consequently, physicians should have a high index of suspicion when young girls present to them with symptoms that most commonly relate to IH, i.e. cyclical lower abdominal pain and/or urinary outflow obstruction combined with the absence of menarche [17].

The treatment of IH is surgical and different types of incisions are proposed. In our case, as it was also reported in other articles [2] [16] an X-shaped incision was chosen combined with the insertion of a Foley catheter in the urethra so as to stent it. This method protects the integrity of the urethra [2]. Despite its supe-
priority in maintaining the urethra intact, the X-shaped incision is not suggested in the vast majority of reports published in the literature. A cross-shaped incision of the hymen, with or without the use of a Foley catheter to stent the urethra, seems to be the incision of choice [1] [3] [7] [8] [11]. Other types of incision include: a star-incision so as to protect the Bartholin’s glands [9], vertical midline incision (<1cm) [12], annular incision with [4] or without [6] [10] [13] placement of a Foley catheter in the opening of the hymen and an elliptical incision [16]. The annular incision is claimed to have the least effect on the hymen’s architecture and thus appears to be the preferred choice among cultures and religions where a girl’s virginity is of great importance [4] [6].

Apart from the type of incision, a differentiation has been noticed in the use or not of stitches to prevent the reformation of the hymen. In our case, stitches were used as a means of haemostasis and no reformation of the hymen was reported. Not only in cases where stitches were applied [2] [3] [8] [12] [13] but also in those where they were not used [1] [6], no reformation of the hymen was detected. Only one case of reformation after four repeated surgeries is reported [4].

The surgical instrument used to perform the incision in our case was a No. 11 lancet, as used for most cases in the literature. Only two different techniques have been reported: one demonstrates the use of unipolar electroautery [6] and the other the use of CO₂ laser [10]. Finally, controversy appears to surround postoperative. Some authors report the use of antibiotics [2] [9] and estrogen ointment [9], while others choose to retain the Foley catheter in the incision for two weeks [9]. In our case, none of the above measures were taken. We chose to advise application of Betadine solution to the area, three times a day for the first 24 hours after the procedure. With the exception of one case [4], all patients had an uneventful recovery and follow-up period, as did our patient.

4. Conclusion

In conclusion, IH is a condition whose diagnosis often mandates a high index of suspicion on behalf of the physician. If left untreated, it could result in serious health problems for the patient. The treatment of choice is undeniably surgical. Although different types of procedures have been proposed, the surgeon should choose the procedure that best suits the patient’s needs without harming anatomical structures closely related to the hymen.

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**Ethics**

The procedures followed were in accordance with the ethical standards of the Helsinki Declaration of 1975, as revised in 1983. A written consent form was given by the patient’s parents in which their agreement to publish this report was stated.

**Conflicts of Interest**

The authors declare no conflicts of interest regarding the publication of this paper.

**References**


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