Case Report

Urachal remnants presenting as an umbilical sinus in middle age: a rare case report

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ABSTRACT

Urachal sinus is a rare type of Vitello intestinal duct anomalies. Clinical presentation is that of persistent umbilical discharge. It is usually seen in paediatric age group. Moreover, they have a different course in adults than paediatric age group in which they frequently involute and have a benign course. These remnants are prone to infection and development of malignancy. A proper diagnostic workup by clinical and imaging tools is required. Early removal of urachal remnants at first diagnosis is key for preventing future morbidities. In our case, all the three remnants (Vitello intestinal duct, urachus and ligamentum venosum) were found in a middle-aged patient and hence is unusual and a rare presentation.

Keywords: Omphalectomy, Sinogram, Urachus, Umbilical sinus, Vitello-intestinal duct

INTRODUCTION

Vitello intestinal duct or omphalomesenteric duct (OMD) is an embryonic structure, which connects the yolk sac to the midgut and failure of its resorption results in various anomalies including Meckel's diverticulum, patent vitelline duct, fibrous band, sinus tract, umbilical polyp and cyst. These anomalies occur in approximately 2% of the population and may remain silent throughout life, or may present incidentally with an intraabdominal complication. Umbilical disorders are frequently encountered in paediatric surgery. But, the presentation and progression in paediatric and adult population is different. Adults have a higher risk of urachal cancer and incur more morbidity. Thus, proper and early diagnosis of urachal pathologies is must. Due to rarity, urachal anomalies present a diagnostic challenge in adult population.

We report a middle-aged patient with a discharging umbilical sinus in whom all the three remnants of Vitello intestinal duct namely, obliterated Vitello intestinal duct, partially obliterated ligamentum venosum and patent urachus were present.

METHODS

A 55-year-old male presented with creamy foul-smelling discharge from umbilicus for the past 1 year which was at times blood tinged associated with pain over umbilicus. On examination, umbilicus was inverted and centrally placed with creamy non-blood-stained discharge and tenderness around the umbilicus. Contrast enhanced CT abdomen with a sinogram of umbilical sinus revealed multiple blind tracts opening into the umbilicus in the pre-peritoneal space, one extending upwards and the other towards the bladder ending blindly (Figure 1).

Culture and sensitivity of the discharge showed no signs of infection. A working diagnosis of umbilical sinus was made and after pre-operative assessment and pre-anaesthetic work up, patient was taken up for exploration
of the sinus after an informed, written and signed consent.

Figure 1: The umbilical sinus as pointed by the arrow in CT scan.

Intra-operative findings included umbilicus with obliterated urachus, Vitello-intestinal duct and obliterated umbilical vein. Patient underwent omphalectomy with excision of the sinus tract and all the three remnants of umbilicus (Figure 2,3).

Figure 2: Demonstrating the urachal remnants arising from the umbilicus.

Figure 3: Complete specimen after resection.

Post-operative stay was uneventful. Histopathology of the specimen reported keratinized squamous epithelium of the remnants with no features of malignancy.

DISCUSSION

As far as umbilical discharge is concerned, it is quite alarming and in adults only a few studies on the urachal sinus have been conducted. Causes are congenital and acquired, though acquired conditions are more common.

The incidence of urachal pathologies in childhood is approximately 1 in 5000 with a male to female ratio of 3:1.2

In adults it is rare, approximately 2 cases per 100,000 hospital admissions, because urachal anomalies usually involute in early childhood.3

The presentation and progression in pediatric and adult population is different. Adults have a higher risk of urachal cancer and incur more morbidity. Thus, the proper and early diagnosis of urachal pathologies is must. Due to rarity, urachal anomalies present a diagnostic challenge in adult population. However, with proper Clinical and Imaging workup they can be managed effectively.4

Urine discharge from umbilicus suggests patent urachus, hematuria points to vesico urachal diverticulum and pus discharge from umbilicus may be present with urachal sinus. Further complications such as infection and malignancy may present. Imaging has a definitive role in classifying the type of urachal anomaly and further characterizing the disease. In case of external opening, contrast can be given through the umbilicus to delineate the tract.

A relatively anterior location in the preperitoneal space with no obscuration by bowel gas makes ultrasound a good tool for diagnosis. CT is further required for confirmation and to look for malignancy. Findings include a tract/collection extending from the umbilicus with prominent median umbilical vein. In case there is a tract communicating with the bladder then it signifies a patent urachus. Blind ending tract arising from umbilicus suggests umbilical sinus. And double-blind ending cavity is a urachal cyst. Urachal anomalies may get secondarily infected via lymphatic, hematogenous or vesical route by a wide spectrum of microorganisms and may form a urachal abscess. CT may reveal inhomogeneous attenuation or abnormal enhancement (rim or patchy enhancement).5

Treatment includes complete excision of the urachal remnant. In case of infection/abscess, initial control of infection/pus drainage should be followed by surgery. A complete excision of the wall is important as there is a high probability of reinfection and chances of development of malignancy in residual remnants.6,7
CONCLUSION

Urachal anomalies are rare clinical entities and asymptomatic urachal sinuses persisting into late adulthood even more so. Infected urachal remnant should be considered in patients with umbilical discharge and inflammation. Finding all the three remnants is an extremely uncommon anomaly. However, it should be considered as a differential diagnosis of discharging umbilicus. Correct diagnosis with multimodality imaging and complete surgical resection is recommended to prevent subsequent reinfection or malignant transformation.

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REFERENCES
