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ABSTRACT

Radiology Section

Cystic lymphangiomas are usually congenital malformations of draining lymphatic channels with most common sites including neck, axilla, mediastinum and retroperitoneum. Occurrence of lymphangiomas in scrotum or inguinal region is a rare entity and epididymal origin of these lesions is even more infrequent. We herein report a case of epididymal lymphangioma detected on USG, which developed after surgical abdominal lymph nodal dissection in an adult patient of carcinoma rectum presenting as painless scrotal swelling.

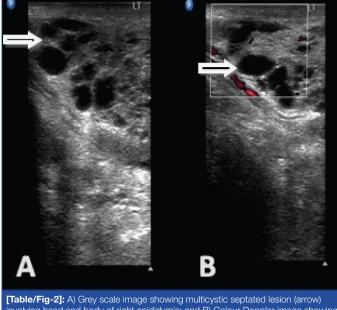
CASE REPORT

A 24-year-old male presented to the institute in GI Oncosurgery Department as a diagnosed case of metastatic adenocarcinoma of rectum. The cross sectional imaging done in the radiology department also showed evidence of polypoidal infiltrative rectal mass with perirectal, bilateral iliac and paraaortic lymphadenopathy and bony lesions [Table/Fig-1]. Patient underwent abdominoperineal resection with abdominal lymph node dissection after receiving neoadjuvant chemotherapy and radiotherapy. Thereafter, patient received multiple cycles of FOLFOX-4 based chemotherapy and was on regular follow up for one and a half years, when he presented with complain of painless right scrotal swelling. USG examination revealed multicystic septated lesion involving head and body of right epididymis measuring 3.0 x 1.6 x 3.6 cm. On colour Doppler imaging there was no internal vascularity [Table/Fig-2a,b]. Mild ipsilateral hydrocele was also present. Bilateral testis were unremarkable. Imaging differentials of cystic lymphangioma, loculated hydrocele, epididymal cysts and spermatocele were considered and patient



[Iable/Fig-1]: High resolution 1200 axial image showing i mass (arrow) with perirectal lymphadenopathy.

Keywords: Epididymis, Lymphatic, Rectum, Scrotum



involving head and body of right epididymis; and B) Colour Doppler image showing no internal vascularity (arrow).

underwent excisional biopsy. Histopathological evaluation was suggestive of epididymal lymphangioma as evidenced by microscopic findings of network of multiple irregular dilated cystic spaces lined by single layer of endothelium with stromal infiltration by lymphocytes. Postoperative period was uneventful and the patient was discharged in stable condition. Follow up with USG imaging after three months showed no evidence of recurrence.

DISCUSSION

Lymphangiomas are benign lymphatic malformations mostly presenting in paediatric population and consist of approximately 26% of all benign vascular neoplasms in children [1]. Occurrence of these lesions is infrequent in adults. The most common sites for children as well as adults are head/neck, axilla, mediastinal and retroperitoneal regions. Scrotum and inguinal canal are uncommon sites for development of lymphangiomas as only about 20 cases have been reported in literature [2]. Epididymal lymphangiomas are even more infrequently encountered, to the best of our knowledge, only six cases have been reported so far with the first case reported

by Thompson in 1936 [3].

Epididymal lymphangioma can occur either as congenital true primary neoplasm or secondary to infection, trauma or postsurgical lymphatic blockage. Out of previously reported 6 cases of epididymal lymphangioma one case was thought to develop secondary neoplasma to herniorrhaphy [4] and rest of the five cases were diagnosed as true primary neoplasms [3,5-8].

In 1976, Whimster first elaborated lymphangiomas, he postulated these lesions to be congenital abnormalities due to anomalous development of the lymphatic channels, originating from sequestered lymphatic tissue which does not develop a normal communication with lymphatic system and thus, present as mass later on [9].

The imaging differentials of cystic epididymal mass lesions mainly consist of epdidymal cysts, spermatoceles and loculated hydrocele. Epididymal cysts and spermatoceles are usually unilocular lesions while lymphangiomas are multicystic septated lesions. However, imaging may be suggestive but definitive diagnosis can only be made by histopathological examination. These lesions frequently show recurrence due to incomplete surgical excision in 25-30% of the cases [10,11].

We herein present a case of right epididymal lymphangioma in a patient of metastatic carcinoma rectum on palliative chemotherapy. Around two years ago, the patient underwent abdominoperineal resection with iliac and paraaortic lymphnode dissection. This patient presented to our institute with painless right scrotal swelling, underwent USG and histopathological evaluation which was finally diagnosed as epididymal lymphangioma possibly resulting as sequelae to previous surgery two years back. Follow up USG examination didn't show any recurrence. To the best of our knowledge, only one such case has been described in literature, where lymphangioma of the epididymis was thought to develop secondarily to previous herniorrhaphy surgery [4]. Our case report is unique in several aspects. First of all lymphangiomas in general are rare in adult patients, inguinoscrotal lymphangiomas are even rarer. A case of adult lymphangiomas, arising from the epidydimis, secondary to surgery is only the second case reported in literature till now, to the best of author's knowledge.

CONCLUSION

The purpose of our case report is to familiarize the radiologists with this rare entity. Although extremely rare, differential of cystic lymphangiomas should also be considered in a patient presenting with scrotal swelling with previous history of surgery.

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