Mature Cystic Teratoma of Adrenal Gland Masquerading as Myelolipoma

Kumar Anmol a, Reecha Singh b*, Umakant Prasad c and M. Vivek Nayak a*

a Kasturba Medical College, Manipal, India.
b Department of Pathology, Indira Gandhi Institute of Medical Sciences, Patna, India.
c Department of Radiology, Indira Gandhi Institute of Medical Sciences, Patna, India.

Authors’ contributions
This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information
DOI: 10.9734/JAMMR/2022/v34i234857

Open Peer Review History:
This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here:
https://www.sdlarticle5.com/review-history/93367

Received 05 September 2022
Accepted 11 November 2022
Published 12 November 2022

ABSTRACT

Introduction: Extragonadal teratomas are uncommon tumors and are less common in adults than children. They are mostly retroperitoneal. Primary adrenal teratomas are even rarer.

Objective: We report an unusual case of primary adrenal teratoma in a young adult.

Case Summary: A 29 year old male presented with complains of intermittent right sided flank pain for 6 months, dull aching, radiating to groin. On CT Abdomen a large heterogenous mass was seen having predominantly fat component and punctuate calcification in right suprarenal region. Histopathology confirmed a diagnosis of adrenal teratoma.

Conclusions: Adrenal teratomas are difficult to diagnose clinically or on imaging as these mimic other lipomatous lesions. Although rare this should be considered as a differential diagnosis for adrenal tumors.

Keywords: Adrenal; teratoma; lipomatous.
1. INTRODUCTION

Teratomas are totipotent germ cell tumors which usually arise from one or all three germ cell layers i.e., ectoderm, mesoderm, and endoderm. Teratoma is a rare neoplasm with an incidence of 0.9/100,000 population [1]. Most common sites are gonads, sacrococcygeal, mediastinal, and pineal region [2]. Retroperitoneal teratomas account for 4% of all primary teratomas whereas adrenal teratomas are even rarer [3]. Adrenal teratomas can pose a diagnostic challenge because radiologically they mimic myelolipoma, angiomylolipoma, liposarcoma and pheochromocytoma. We report an unusual case of mature teratoma in right adrenal gland in a young adult male.

2. CASE SUMMARY

A 29 year old male presented with complains of intermittent right sided flank pain since 6 months, dull aching, radiating to groin. There was no associated vomiting, hematuria, dysuria or no lower urinary tract symptoms. Physical examination and complete blood count were within normal limits. Serum cortisol, plasma metanephrine and urinary VMA level were also within normal limits. On Ultrasonography a well defined, heterogenous mass of size (10 x 8)cm, oval shaped was seen between right lobe of liver and right kidney having hyperechoic central component with posterior acoustic shadow was seen.

Fig. 1. CT image of upper abdomen: A spherical mass in right adrenal region with predominantly fat and punctate calcification

Fig. 2. Gross findings: Cut specimen show cyst with tufts of hair and cheesy material
Fig. 3. Microscopy: H&E stain (200x) Components of cyst – Skin with hair shaft, mature adipose tissue and lymphoid aggregates

Fig. 4. Microscopy: H&E stain (200x) Shows adrenal gland tissue (Arrow) compressed at periphery and mature adipose tissue
On CT Abdomen a large heterogenous mass, measuring about (7x 8) cm was seen having predominantly fat component, soft tissue density lesion and punctate calcification in right suprarenal region suggestive of Adrenal Myelolipoma. (Fig. 1) Right sided laparoscopic adrenalectomy was done without any complications. The specimen was sent to histopathologic laboratory. On gross examination, a cystic mass measuring (10x10x5) cm with a wall thickness of 1 cm was seen. External surface was congested, cut section showed thick cheesy material with hairtuft. (Fig. 2) On microscopic examination, derivatives of all three germ layers were seen. Compressed adrenal cortical tissue was seen in the cyst wall (Fig. 3) Thus a confirmatory diagnosis of mature teratoma of Adrenal gland was established.

3. DISCUSSION

Gonadal teratomas are known to occur however extra gonadal teratomas are uncommon neoplasms. Adrenal teratomas form about 0.13% of all adrenal tumours [4]. “Clinically, adrenal teratomas usually occur in young women, in the right adrenal gland, and 90% are benign” [5]. Patients are mostly asymptomatic, however in our case the patient presented with right sided flank pain. Abdominal CT is the choice of imaging which usually shows well-circumscribed heterogenous masses mainly comprising of fatty component with multiple tissue types such as bone, teeth, soft tissue and fluid component. In a study by Davidson et al. [6] comprising 21 cases, it was suggested that “the presence of a heterogeneous mass containing a well-circumscribed fluid volume, fat-fluid and calcifications were the most characteristic radiologic findings on CT”. These findings were consistent with our case report.

“Indicators for malignant transformation include significant enhancement of the cyst wall, septations as well as mural nodules and abnormal levels of hormones including cortisol, ACTH, aldosterone, and VMA” [7]. No such findings were evident in our case.

“Elevated levels of CA 19-9 and CA 125 may be an indicator of bilaterality. Since levels of CA 19-9 and CA 125 may be elevated in both benign and malignant conditions, interpretation of these findings must be made in light of the clinical condition of the patient” [8]. In our case these findings were within normal range.

“Outcomes regarding the comparison of elevated CA 19-9 in patients with MCT and ovarian cancer were available” by Cho et al. who showed that “despite the fact that simultaneous elevation of CA 19-9 and CA 125 was more prevalent in patients with malignancy, single elevation of CA 19-9 was more frequently detected in MCTs compared to malignancies” [9].

“Analysis of the multi-region exome sequencing data was used to study the clonal relationship of immature and mature teratoma elements, as well as admixed foci of yolk sac tumor and also disseminated teratoma in the peritoneum” by Heskett MB et al. [10]. They found that “all these different tumor components were indistinguishable based on chromosomal copy number alterations and loss of heterozygosity patterns, indicating a shared clonal origin. This finding suggested that epigenetic differences were likely responsible for the striking variation in differentiation patterns in teratomas” [10].

Histopathology is essential for a confirmatory diagnosis of mature teratoma. Surgery is the method of choice for treatment of mature teratoma. Laparoscopic surgery is the gold standard for adrenal tumors [11] with excellent prognosis and 5 year survival rate of 100%

4. CONCLUSION

This case report emphasizes that adrenal teratoma should be considered as an important differential diagnosis for lipomatous lesions involving the Adrenal gland

CONSENT

As per international standard or university standard, patient(s) written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

As per international standard or university standard written ethical approval has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES


© 2022 Anmol et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here: https://www.sdiarticle5.com/review-history/93367