Gigantic retroperitoneal metastatic seminoma encasing the aorta in a young man

Abdul Wahab,1 Muhammad Hashim Hayat,2 Amman Yousaf,3 Raseen Tariq4

DESCRIPTION
A 30-year-old man with no significant medical history presented with worsening of lower abdominal discomfort and a weight loss of 70 pounds during the past 6 months. Physical examination was significant for a large, firm, non-tender, irregular mass in the lower abdomen, more prominent on the left of the midline, and a small-sized non-tender swelling of the right testicle. Lactate dehydrogenase was 1562 U/L (135–225 U/L), uric acid 8.2 mg/dL (3.4–7.0 mg/dL), serum ß-human chorionic gonadotropin (HCG) 85 mIU/mL and alpha-fetoprotein (AFP) 1.3 ng/mL (0.0–9.0 ng/mL). CT scan of the abdomen and pelvis showed a large retroperitoneal mass encasing the abdominal aorta and its branching (figure 1A) and compressing the inferior vena cava (figures 1B, C and 2). Testicular ultrasound (US) showed a small right testicular mass with internal vascularity (figure 3A, B).

The patient underwent right-sided orchiectomy and US-guided biopsy of the retroperitoneal mass. Histopathology and immunopathology confirmed metastatic seminoma (figure 4A, B). After semen cryopreservation, the patient was started on chemotherapy. The patient has received four cycles of bleomycin, etoposide and cisplatin-based chemotherapy and no complications have been report till date (on 3-month follow-up).

Testicular germ cell tumours are the most common cancers in men aged 15–39 years with an increasing incidence rate during the last four decades.1 Seminoma is the most common subtype of testicular malignancy and it originates from seminiferous tubules, due to the malignant transformation of primordial germ cells.2 Testicular tumours usually present as a painless swelling of the testis found by the patient, his partner or during physical examination. Patients with metastatic disease can present...
Images in...

Figure 4  Histopathological features of the biopsied specimens. (A) H&E-stained slide from the testicular specimen (200× magnification). (B) H&E-stained slide from the retroperitoneal mass. Both images show epithelioid cells present in sheets with abundant clear to pink granular cytoplasm, polygonal nuclei and distinct nucleoli consistent with seminoma.

with local signs and symptoms (eg, abdominal discomfort, neck mass, the difficulty of breathing, bone pain, extremity swelling, etc) based on the location of the metastasis. Moreover, a giant retroperitoneal mass can have a mass effect on the surrounding vessels such as the aorta, its major branches and a mass effect on veins can lead to thrombosis such as inferior vena cava (IVC) thrombosis in our patient.2 3 Scrotal ultrasonography confirms the existence of mass and usually shows hypoechoic lesion without calcifications and cystic component in cases of seminoma.

In cases of seminoma, LDH (lactate dehydrogenase) can be elevated and occasionally slightly elevated ß-HCG but seminomas never secrete AFP.2 Radical orchiectomy is the standard of care for primary tumour and histopathology of the specimen confirms the diagnosis and differentiates the type of germ cell tumour.1 3 CT scan of the chest, abdomen and pelvis and MRI of the brain are used for staging. Patients must be provided with details about adolescent and young adults with cancer programmes and fertility preservation options.2

For patients with stage 1 seminoma, post orchiectomy options include active surveillance, retroperitoneal lymph node dissection or chemotherapy (one or two cycles).1 For most advanced tumours, chemotherapy remains the standard of care.1 2

Learning points

► Seminoma is the most common subtype of testicular malignancy that usually presents as a painless scrotal swelling and metastatic disease can demonstrate focal manifestations.
► A huge abdominal mass can be misleading and physicians might ignore examining the scrotum.
► Scrotal examination and ultrasonography can confirm the existence of a mass lesion that is usually solitary, hypoechoic without internal vascularity. Management can include orchiectomy, retroperitoneal lymph node dissection or chemotherapy, depending on the tumour stage.

Acknowledgements  The authors are grateful to Dr Anthony N. Snow, MD (clinical associate professor of pathology at University of IOWA Health Care) for his valuable contribution in histopathological slides for this case.

Contributors  AW, MHH and AY: Drafted and revised the manuscript. RT: Drafted and revised the manuscript and gave final approval for submission.

Funding  The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests  None declared.

Patient consent for publication  Obtained.

Provenance and peer review  Not commissioned; externally peer reviewed.

ORCID iD  Amman Yousaf http://orcid.org/0000-0003-0646-508X

REFERENCES