Paramedian chest wall dermoid cyst

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DESCRIPTION

Congenital dermoid cysts occur at lines of embryological fusion due to ectoderm sequestration. Despite being congenital, dermoid cysts may not present until late childhood or even later.1

Implantation dermoids (epidermoid cysts) have different pathogenesis. They happen secondary to mechanical implantation of epidermis into the subcutaneous tissue secondary to injury or surgery.

The incidence of dermoid tumours is roughly 3 per 10,000 paediatric patients, with greater than 80% of paediatric dermoids occurring in the head and neck.2 Other sites include midline of the chest. Literature review shows case reports of exception to this rule, but no truncal non-midline lesions have been reported.3 4

A 14-year-old male patient presented to outpatient clinic with a painless swelling in left upper part of chest wall of 2 years duration. He had no history of previous injury or surgery to the site. He was otherwise fit and well.

Clinical examination revealed a 1 cm spherical swelling superficial to the left 1st intercostal space 4 cm from midline (figure 1). The swelling was soft, mobile, not tender and not tethered to skin or deep structures. The overlying skin was unremarkable. There were no other swellings or palpable lymphadenopathy. A provisional diagnosis of a lipomatous tumour was made.

Complementary ultrasound scan was requested but was unfortunately indeterminate, and a specific diagnosis could not be made.

The decision was made to excise the lump under local anaesthesia.

Intraoperatively, a transverse incision was made over the lump. The lump was accidentally punctured by the blade. Expressing the lump contents, that is, creamy material and hair. The lump was completely excised and sent for histopathology.

The analysis of lump revealed ruptured cyst containing some keratin. Adjacent to the cyst, there were skin adnexal structures such as sebaceous glands and sweat glands. There was no significant adipose tissue (figure 2). The features were consistent with the congenital dermoid cyst.4 Implantation dermoids have no adnexal structures and,

Figure 1 An arrow is pointing to location of the cyst on chest wall.

Figure 2 Histological features of the excised cyst.

Patient’s perspective

I am mother of the patient and my son gives permission and consent for this publication. I am eager to get it published as a rare incident.

Learning points

► Dermoid cysts scan present in unusual places.
► Unusual places can be due to migration of the cyst or atypical embryological fusion lines.
► Dermoid cysts may present in late childhood.
therefore, they are easily distinguished from congenital dermoid cysts on histology.

Dermoids occurring outside classical embryological fusion lines are not described. This has occurred at the junction between cervical and thoracic innervated tissue, but this is not understood to be an embryological fusion line. We are not sure if this cyst had migrated from midline. As such it is interesting, and a potential diagnosis for paediatric surgeons to be aware of.

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REFERENCES