The Unusual Case of A Sternoclavicular Joint Cyst In A 21 Months Old Child

Stefan Bittmann, Elisabeth Luchter, Gloria Villalon, Elena Moschüring-Alieva

Abstract
Sternoclavicular joint cysts are extremely rare in children. We report the 6th case of a ganglion of the sternoclavicular joint in a child, which was diagnosed by ultrasonography.

Keywords: Sternoclavicular Joint; Cyst; Synovial Cyst; Ganglion; Child

Introduction
Sternoclavicular joint cysts are extremely rare in childhood. Only 5 cases have been reported thus far in the medical literature [1]. We describe a case of a paraarticular cyst of the right sternoclavicular (SC) joint in a child.

Case Report
A child referred to our pediatric department with a 6 month history of a firm, painless swelling at the ventral aspect of the right sternoclavicular joint which enlarged slowly over time. At clinical examination, the size of the lesion was estimated to be approximately in 2 cm in diameter. A conventional radiograph (Figure 1) showed no obvious osseous or soft tissue abnormality. Ultrasound examination (Figure 2) revealed a paraarticular cyst with a small stalk communicating with the right sternoclavicular joint. Due to the asymptomatic nature of the lesion, no surgical intervention was undertaken.

Discussion
Many different conditions may be associated with the sternoclavicular (SC) joint. Hyperostosis of the sternoclavicular joint, osteitis condensans, Friedrich’s syndrome, Tietze’s disease, subluxation of the SC joint, and ganglion cystic lesions. All of these conditions are rare in childhood. Ganglia are structures of cystic form surrounded by fibrous layer without a synovial structure and are full of mucinous fluid. They most commonly present as masses around hand that can induce compressive symptoms. They may be found near most major joints. Their etiology is unclear. On ultrasound they typically described as anchoic masses, and
may have a communication with a joint or tendon tissue (see Figure 1). In adults, synovial cysts originating from the sternoclavicular joints were described with rheumatoid arthritis, spondyarthropathies, enthesis and psoriatic arthritis [2,3,4-7]. Ganglion cysts that have their origin from the sternoclavicular joint have only been reported once in childhood. Haber et al described the first 5 cases of sternoclavicular joint cysts in children. In this study, only 1 patient was symptomatic. Biopsy was performed in 3 patients and the diagnosis of a ganglion cyst was confirmed by histological analysis. No recurrence was found and present in this study. Haber recommended observation of asymptomatic cystic lesions that arise in the sternoclavicular joint. No extensive study concerning the treatment of ganglion cysts of the SC-joint has been reported in the medical literature so far. The most often described and published cases are single cases. Asymptomatic patients should be observed and symptomatic patients may be best treated with steroid injection or cyst excision.

Figure 1: No abnormality is discernable in a radiograph of the right SC joint with a lead marker taped to the superior aspect of the subcutaneous swelling.

Figure 2: An axial ultrasound image of both sternoclavicular joints, using a broadband (frequency 8-14 MHz) linear array transducer and a standoff gel pad, shows an anechogetic cystic structure communicating via a stalk (white arrow) with the right SC joint.

References

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