Voluntary Bilateral Scapulothoracic Dissociation

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ABSTRACT / Study design: Case report.
Objective: To report a rare case of voluntary scapulothoracic dislocation and its treatment.
Background data: Scapulothoracic dissociation is a rare condition. We report a case of a teenager who could voluntarily dislocate her scapulae.
Results: The patient was treated surgically by fixating her scapula to the contralateral transverse process and carrying out associated muscle plication. Now, after a 12-year follow-up, the patient is asymptomatic.
Conclusions: Fixation of the scapula to the contralateral transverse process and plication of associated muscles is a valid treatment for this condition.

Key words: perinatal outcomes, older women, pregnancy.

Introduction
The scapulothoracic joint forms part of the complex shoulder girdle and it contributes to a third of the arm’s elevation. Codman has described this process as the scapulohumeral rhythm.

The scapulothoracic joint is formed by the ventral surface of the scapula covered by the subscapularis muscle, and the posterolateral aspect of the thoracic cage covered by the serratus (posterosuperior and inferior) muscles between the 2nd and 8th rib, and an intermediate fascial sheet that runs between them from the 4th to the 9th rib.

The scapula is attached to the rib cage through the insertion of different muscles (the trapezius, becoming a true suspensor ligament and the levator scapulæ in the superior surface; rhomboideus minor and rhomboideus major and again the trapezius in the posterior surface restricting the anterior translation, antagonized by the serratus anterior inserted in the anterior lip of the scapula; meanwhile the latissimus dorsi as well as the teres major and minor inserted in the inferior angle apply the scapula against the thorax).

Dislocation or scapulothoracic dissociation is a rare condition, occurring mainly in association with a high-energy trauma; only a few reports of atraumatic or voluntary dislocation cases have been published.

We report the case of a teenager who could voluntarily dislocate both her scapulae.

Case Report
The case concerns a 16-year-old girl who came to our hospital with a history of recurrent episodes of bilateral scapular dislocation that had taken place over the previous few months.

The patient had a history of hyperlaxity, with sporadic episodes of subluxation of her hand joints and temporomandibular dislocation. When she was 14 years old, she underwent surgery for snapping hip syndrome. We suspected Ehlers Danlos disease.

No abnormalities were found by general examination. The patient’s cardiovascular and ophthalmologic examination were normal. Osteoarticular physical examination was also normal except for an ability to do a voluntary lateral dislocation of her scapulae, which, from behind, gave the appearance of a swimmer’s back, without altering the function of her upper limbs. (Fig. 1-2)

Figure 1 / Macroscopic aspect preoperative
In January 1997, surgery was performed because the patient could no longer do normal daily activities or sports as a result of the high recurrence of the scapular dislocation.

Fixation of the scapula was by means of a Dacron ligament prosthesis to the contralateral transverse process, associating a plication of the muscles of the inner edge of both scapulae. (Fig.3)

The patient progressed satisfactorily until August 1997, when she presented a left winged scapula that needed a new muscle plication. Later, in November 1997, she suffered an episode of lateral displacement of her right scapula; this was treated by anchoring the scapula to the ribs with Dacron. (Fig.4)

No alterations were found in the histological study of muscle, skin, fat and fibrous tissue.

Biochemical studies revealed a decrease in type III collagen. Genetic studies, however, revealed no mutations, and consequently we could not confirm a diagnosis of Ehlers Danlos disease.

At present, after a 12-year follow-up, results remain satisfactory: there is no shoulder movement limitation and there have been no new episodes of scapula dislocation. (Fig. 5)
Discussion

Scapulothoracic dissociation is usually a result of a high-energy trauma or a violent traction of the upper limb. Neurovascular injuries such as brachial plexus injury, or subclavian artery lesion and shoulder girdle fractures are frequently associated. 4-7, 8

Only a few cases of non-traumatic scapula dissociation, dislocations, or subluxations have been described in the literature. Hollinshead9 reported a case of scapula subluxation in a male patient with an osteochondroma in the ventral surface of his scapula resulting in snapping and blockage of the joint. The patient was treated by resection of the tumor.

Ward10 presented a case of non-traumatic scapulothoracic dislocation secondary to a Pancoast tumor resection. Walker11 described the case of a 19-year-old girl with a unilateral scapulothoracic dislocation after a small fall; treatment was by closed reduction.

In the literature, we have found only one other case of dislocation that had no associated pathology or related trauma: Kushwaha12 reported the case of a woman who suffered a dislocation when changing gears in her car; closed reduction was the treatment.

What distinguishes the case reported here, is that our patient was able to dislocate and relocate both her scapula voluntarily. Studies were conducted to investigate the a possible diagnosis of Ehlers Danlos Syndrome, as this is characterized by joint hyperlaxity that is associated with joint subluxation and dislocation13. The results of these genetic and histological studies were negative for Ehlers Danlos Syndrome.

The surgical treatment was conceived to anchor the scapulae without sacrificing joint movement. To this end, we used Dacroon grafts (used in occasions for ACL reconstruction) and then, to increase stability of the shoulder joint we carried out plication in associated muscles.
Research paper

References


