Case Report

Winging of Scapula due to a Sinister Etiology

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Received 30 May 2020; Revised 17 September 2020; Accepted 26 October 2020; Published 4 November 2020

Academic Editor: Mehmet Turgut

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Background. Scapular winging is a rare but disabling deformity, which is commonly caused by lesions of the long thoracic and spinal accessory nerves that innervate the serratus anterior and trapezius muscles, respectively [1]. Rarely, it can be caused by damage to the dorsal scapular nerve which innervates the rhomboid muscle [1]. The spinal accessory nerve is the XIth cranial nerve, while the long thoracic and dorsal scapular nerves originate from the cervical nerve roots, C5, 6, and 7, and C4, 5, and 6, respectively. Traumatic damage to these nerves causing neuropraxia accounts for the majority of the cases of scapular winging [2].

1. Introduction

Scapular winging is a disabling deformity caused by lesions of the long thoracic and spinal accessory nerves that innervate the serratus anterior and trapezius muscles, respectively [1]. Rarely, it can be caused by damage to the dorsal scapular nerve which innervates the rhomboid muscle [1]. The spinal accessory nerve is the XIth cranial nerve, while the long thoracic and dorsal scapular nerves originate from the cervical nerve roots, C5, 6, and 7, and C4, 5, and 6, respectively. Traumatic damage to these nerves causing neuropraxia accounts for the majority of the cases of scapular winging [2].

Apical lung tumors produce various neurological manifestations owing to its local infiltration or paraneoplastic effects. Reported cases of winging of scapula as the initial mode of presentation of an apical lung malignancy are extremely rare [3]. We present a case where an apical lung malignancy is manifested as winging of the ipsilateral scapula, by the infiltration of C5–C7 roots of the brachial plexus.

2. Case

A 54-year-old man presented with gradual onset slowly progressive weakness of the right arm over a period of 2 months. Weakness was preceded by pain in the right shoulder region which radiated down the right arm. Pain was present even at rest. Weakness was predominant in the proximal muscles of the right upper limb, with preserved fine movements of the right hand. He denied a sensory loss
involving the affected limb. His left upper limb and both the lower limbs were normal with regard to the power and sensation. He did not have dysphagia, dysarthria, diplopia, or unsteadiness. His higher functions were intact. He did not report any fever, loss of appetite or weight, chronic cough, or hemoptysis. There was no recent history of trauma. He was not a smoker and neither did he consume any illicit drugs.

On examination, enlarged, multiple, hard, and fixed lymph nodes were noted in right supraclavicular fossa and anterior triangle of the neck. There was no finger clubbing. Wasting and fasciculation were observed in the muscles of the shoulder girdle and proximal part of the right upper limb, sparing forearm, and the hand muscles. Proximal muscle power was of 3/5 with preserved distal power. Posterior winging of the right scapula was noted, which became more prominent up on pushing against resistance (Figures 1(a) and 1(b)). Rest of the examination including voice, cranial nerves, pupils, contralateral upper limb, and lower limbs was normal. Bronchial breathing was heard in the upper zone of the right lung, with trachea slightly deviated to the opposite side.

The electromyography (EMG) revealed large motor unit potentials with increasing amplitude and duration and poor activation of right serratus anterior and internal scapulae muscles. The nerve conduction studies (NCS) suggested a compressive lesion involving C5–C7 nerve roots of the brachial plexus. The chest radiograph revealed a large homogenous opacity involving the right apical region (Figure 2).

Subsequent contrast-enhanced computerized tomography (CECT) showed an ill-defined enhancing malignant lesion, obliterating the right upper bronchus, right upper lobe, pulmonary artery, and pleura (Figure 3). Probable stage was IV (T4N3M1a). Enlarged necrotic lymph nodes were seen in the mediastinum, with some in the right supraclavicular fossa. Histology and immunohistochemistry (Figure 4) of a cervical lymph node revealed metastatic deposits from a poorly differentiated carcinoma, most probably of pulmonary origin.

At bronchoscopy, right upper bronchus was seen to be completely obstructed by a tumor with endobronchial invasion. The histology (Figure 5) and immunohistochemistry from the tissue obtained by bronchoscopy confirmed a poorly differentiated adenocarcinoma of the lung. Magnetic resonant imaging (MRI) of cervical spine was planned, but the patient refused to undergo further investigation.

3. Discussion

Scapular winging is a disabling condition that hinders the ability to lift, pull, and push against objects. Winging of scapula results from paralysis of the muscles that keeps the scapula anchored to the chest wall. The commonest cause of scapular winging is isolated serratus anterior muscle (SAM) paralysis [2]. Long thoracic nerve (LTN) innervates the serratus anterior muscle. Due to its lengthy course and superficial location, LTN is susceptible to blunt and penetrating trauma [3]. The causes of nontraumatic injury to the LTN include viral infections such as poliomyelitis, drugs, exposure to toxins (herbicides and tetanus antitoxin), muscular dystrophies (fascio-scapulo-humeral dystrophy), C7 radiculopathy, and aortic coarctation [1]. Iatrogenic causes such as thoracotomies for lung or cardiac surgeries, hematomas secondary to anticoagulation therapy, are also identified as causes of SAM palsy [3].

An apical lung malignancy causing scapular winging has been reported only once in the literature [3]. In the case reported by Toshkezi et al., apart from the winging scapula,
The patient also had Horner’s syndrome, which is a recognized neurological manifestation of Pancoast tumors. This would have been a clue to suspect an infiltrative pathology as the cause for the LTN palsy. But ours is a case where an apical lung malignancy caused ipsilateral winging of scapula, in the absence of any other typical neurological manifestations of a Pancoast tumor, such as hoarse voice and Horner’s syndrome, which are due to the involvement of recurrent laryngeal nerve and sympathetic trunk, respectively. Neither did he have any signs of compression or infiltration of the middle and lower trunks of the brachial plexus, such as pain irradiating to the upper limb or small muscle wasting of the hand [4].

The respiratory symptoms such as cough or hemoptysis are uncommon at the initial stages of apical lung neoplasms due to its peripheral location [4]. Apart from bronchial breathing heard at auscultation and cervical lymphadenopathy, a lung malignancy was not suspected as a cause for

Figure 2: Chest radiograph—posterior-anterior view—right side apical ill-defined mass causing tracheal deviation to opposite side.

Figure 3: CECT chest—ill-defined enhancing malignant lesion—obliterating the right upper bronchus, right upper lobe, pulmonary artery, and pleura.
winging of scapula, at the initial evaluation. Apical tumors can easily be missed on plain radiographs in the early course of the disease, as they might be hidden behind the clavicle and the first rib. CT and MR images would be useful to assess the local spread to adjacent neurovascular structures and the spine. The local invasion was classically seen in squamous cell carcinoma, followed by small cell carcinoma of lung, although the current case was of an adenocarcinoma [5].

4. Conclusion

Winging of the scapula is commonly caused by paralysis of serratus anterior muscle (SAM), due to an injury to the long thoracic nerve (LTN) or its nerve roots.

This case highlights the value of considering a compressive lung pathology with infiltration in the differential diagnosis, when evaluating winging of the scapula.

Consent

Informed written consent for publishing the patient’s details including the photos, in the journal, was obtained and is available for the perusal of the editors upon request.

Disclosure

The abstract of this case was presented as a poster presentation at XXIV World Congress of Neurology held in Dubai, United Arab of Emirates, in 2019.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors’ Contributions

SG collected information, followed up the patient, did literature review, and drafted the manuscript. PY analyzed the lung biopsy. BS and HK were involved in the diagnosis and management of the patient and preparing the manuscript. All the authors read and approved the final manuscript.

Acknowledgments

The authors would like to acknowledge the patient who gave full support in collecting information and pictures to be published.
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