Olecranon Bursitis Causing Reflex Sympathetic Dystrophy Syndrome in a Hemiplegic Extremity

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Received July 11th, 2012; revised August 14th, 2012; accepted August 20th, 2012

ABSTRACT

Olecranon bursitis is rarely seen in patients with stroke, but reflex sympathetic dystrophy syndrome is not. Here, we report a stroke case, which represented the clinical symptom of reflex sympathetic dystrophy syndrome after development of olecranon bursitis in few days, and describe his clinical association between both conditions. Sonographic picture of the right elbow in this case is shown.

Keywords: Olecranon Bursitis; Reflex Sympathetic Dystrophy Syndrome; Stroke

1. Introduction

Olecranon bursitis (OB) is rarely seen in patients with stroke, but in contrast, reflex sympathetic dystrophy syndrome (RSDS) is not true. RSDS is an actual condition defined by localized or diffused pain [1]. The affirmed etiology between RSDS and stroke, the degree of severity and recover of motor deficit with sensory problem was not clear [2]. Meanwhile, there is no universally accepted tools for measuring outcomes inpatients with RSDS are confirmed [3]. RSDS of the upper extremity after stroke is famous as shoulder-hand syndrome. The incidence of shoulder-hand syndrome is varying by diverse clinical data or report [2]. Combination of both conditions has never been reported in that kind of patients yet. We herein report a case of stroke, whose hemiplegic extremity developed olecranon bursitis and ensuing RSDS in a short time.

2. Case

A 53-year-old man was admitted to our hospital on 24 December 2010 for inpatient rehabilitation after transfer from another hospital following cerebral infarction in the left hemisphere on 23 October 2010. Three weeks after admission, the patient suffered from painful swelling of the right elbow region, which continued to worsen. The local swelling raised with increased temperature in the lesion side (Figure 1). After confirming the diagnosis of OB clinically, he underwent aspiration of the swelling bursa under sonoguide on 2 January 2011, where showed moderate effusion (Figure 2). Cultures of the aspirated bursal fluid failed to make diagnosis of bacterial and fungal infections.

Shortly after the procedure, the patient grumbled strong sharp pain accompanied with diffuse swelling of his right upper limb, particularly in the wrist region. The pain worsened while stretching the elbow and wrist joints, which was particularly obvious at night. The diagnosis of RSDS of the right hemiplegic limb was made by physical examination as well as triple phase bone scan, which was performed on 24 January 2011, and revealed mildly to moderately increased tracer uptake during the arterial and soft-tissue phases, and moderately to markedly at the right wrist and MP joints of the right hand (Figure 3).

Figure 1. Lateral view of olecranon bursitis of the patient’s right elbow in slight flexion. The local swelling of the elbow was shown.
Figure 2. The ultrasonic image of right olecranon bursitis—before arthrocentesis. The bursa was observed to be filled with fluid and was then confirmed laboratorially as aseptic.

Rehabilitation modality treated the dystrophic condition with some benefit. Another OB took place in the left elbow one month later, but remitted spontaneously in one week.

3. Discussion

This is the first report describing the development of OB and RSDS in the same limb. OB is commonly seen in players in sports, such as wrestling, weight lifting, gymnastics, skateboarding, and rollerblading, particularly when they do not wear elbow protection. OB is also common seen in clinical practice, which etiology can be distinguished as septic and aseptic. There are various kinds of organism resulting in OB, including bacteria, Tubercle Bacillus, fungus, protothecosis etc. The aseptic OB can be related to systemic inflammatory disease, for instance, rheumatoid arthritis or gouty arthritis. According to the literature, the only possible factor contributing to both conditions is infection, but the aspirated bursal fluid failed to confirm the existence of an infection in our patient. The elbow region is the dependent part of the upper limb in supine or seated position, and it may had some degrees of vascular or nerve compromise when it is compressed. We hypothesized that motor deficit might contribute to localized inflammation after repetitive microtrauma, which inflammation result in the subsequent OB.

The peculiar phenomenon of RSDS characterized by autonomic dysfunction and severe pain that may lead to contracture of the affected limbs [4,5]. The incidence for RSDS in the affected paretic upper extremity after stroke varies between 1.5% and 61% [6]. Diagnostic criteria include the occurrence of uncommon regional pain and other sensory changes following an injury. RSDS may develop after brain lesions, and also is mostly often commenced by trauma to peripheral nerve or soft tissue [7]. We suggested that an OB lesion would probably

Figure 3. Images of three-phase bone scan of the right hemiplegic limb. Uptake of Tc-99m was shown increased in blood-pool phase (a), tissue phase (b) and bone phase (c). Stronger isotope uptake can be found in wrist bones and 2nd - 4th metaphalangeal joints in bone phase (d).
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The exact cause of RSDS had not been understood well. The development of unilateral sweating abnormalities after stroke led to the hypothesis that autonomic disturbances occurred in stroke and RSDS might share a common central pathophysiology [8]. Important risk factors for development of a RSDS are subluxation, paresis of the shoulder girdle, moderate spasticity and visual deficits for the paretic extremity. Pain is also one of the frequent reasons. A study demonstrated that peripheral traumatic injury of the shoulder joint capsules led to painful extremities after stroke [9]. Some authors claimed that peripheral lesions might initiate a self-perpetuating vicious cycle of pain followed by the full picture of RSDS [10]. We assumed that in stroke patients with extremity OB, like our case, appears to be at a significantly higher risk of developing RSDS. Based on clinical sequence with OB first and then RSDS in the affected limb, it was difficulty excluding the association between OB and RSDS and further assumption that RSDS might be induced by OB. We hypothesize that, if OB develops, the condition should be treated early to prevent the potential development of RSDS.

This report describes the development of RSDS following OB in a hemiplegic limb, and we emphasized that the important role of motor deficit in hemiplegic limb, because it did not happen in the healthy limb even OB occurrence.

REFERENCES


