Reflex Sympathetic Dystrophy Syndrome of the Ankle in Pregnancy

Gebelikte Ayak Bileğinin Refleks Sempatik Distrofi Sendromu

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Abstract
Reflex sympathetic dystrophy (RSDS) is a type of arthropathy combining a painful syndrome with regional trophic disturbances. The condition primarily affects women during pregnancy, but it also affects middle-aged men. Although hip involvement is the most common form of the disease in pregnancy, other joints such as knee and ankle may be involved. Diagnosis is based on typical clinical and radiological observations. Pain may persist for six months or longer. Fracture is the major risk for the disease. Treatment consists of analgesics, protection against stress fractures, and the prevention of contractures. Reflex sympathetic dystrophy does not appear to affect the course of pregnancy but conclusion of the pregnancy is necessary for cure. We described a patient with pain and local osteopenia of the ankle occurring in the third trimester of pregnancy. We prefer the name reflex sympathetic dystrophy for this condition and we discussed the different terms which determine the same condition. (Rheumatism 2007; 22: 76-9)

Key words: Reflex sympathetic dystrophy, transient osteoporosis, pregnancy

Introduction
Reflex sympathetic dystrophy (RSD) is a pain syndrome that is characterized by autonomic, motor and sensory disturbances with unknown etiology that can encompass a wide range of manifestations in affected limbs (1). The disease usually characterized a self-limiting syndrome, but recurrence may be seen in other joints. Treatment consists of analgesics, protection against stress fractures, and physical therapy for prevention of complications. Transient osteoporosis is an uncommon condition which affects previously healthy women in the third trimester of pregnancy (2). There are several published similar cases which were named different terms such as transient osteoporosis, Sudeck's atrophy, algodistrophy which determine this condition (3). Lequesne (1968) proposed a non-traumatic form of RSD. However, whether all terms were determine the same condition or not, remains to be explored.

This poorly understood disorder, which is frequently deceptive and sometimes clinically misleading especially in pregnant women, characterized by pain in the involved joint (5, 6). We described a woman who developed local osteoporosis in the ankle and foot during pregnancy.

Case Report
A 25 year old woman (gravida 3) at 30 weeks gestation was referred with pain on right foot and ankle. Her...
pregnancy was otherwise uneventful. She had no history of trauma or preceding illness.

Physical examination was showed blue discoloration of the right leg distal to the knees. The skin felt cool and moist. There was marked tenderness below the knees to both light and deep palpation and to percussion. The pain was aggravated by weight-bearing and relieved with rest. Pulses were normal and there were no neurological deficits. Clinical examination revealed decreased skin temperature in the affected areas, change in skin color, and normal articular mobility. Marked hypersensitivity to touch and periarticular soft tissue swelling was evident around the ankles. Her gait was antalgic. No similar symptoms were occurred at prior pregnancies. She was no smoking and had no alcohol consumption. There was no family history for osteoporosis or bone disorders. She had not taken any drugs during pregnancy.

Laboratory evaluations were as follows: Hemoglobin: 11.2 g/dL (12-18 g/dl), ESR: 54 mm/1 h (0-15 mm/h), CRP: 15.3 mg/dL (0.0-0.8), white blood cell: 10 000/mm³ Ca: 8.5 L (8.6-10.2 mg/dl), P: 3.2 (2.7-4.5 mg/dl), ALP: 115 (50-270 U/l), PTH: 41.30 (15-65pg/ml).

The patient refused any form of imaging during pregnancy. A healthy child was delivered at 40 weeks. After delivery, plain radiographs showed that diffuse osteopenia of her right foot and ankle (Figure 1). A magnetic resonance imaging (MRI) was performed. There was low signal intensity on T1-weighted images and high signal intensity on T2-weighted images in ankle and foot (Figure 2). In addition, MRI demonstrated changes consistent with bone marrow oedema affecting bone including the talus, cuboid, calcaneus and lateral cuneiform, and generalized surrounding soft tissue oedema. Repeat radiographs showed results similar to the first one. Bone mineral density revealed decreased bone mass, and dual energy X-ray absorbiometry (DXA) measurement of bone density demonstrated osteopenia of the spine with a T score (femur): -2.2, BMD (femur): 0.607 g/cm², T score (spine): -2.1, BMD (spine): 0.815 g/cm². Plain radiographs of spine and MRI of hip were normal (Fig 3-4). Bone scintigram usually provides early diagnosis in transient regional osteoporosis. Scintigraphy was not performed in this patient, because we did not have a nuclear medical imaging system in our hospital.

The patient was advised to avoid weight bearing, and treated with a combination of salmon calcitonin at a dose of 200 U/l 3 times a week, and 1000 mg Calcium-800 mg vitamin D, and analgesics for four weeks. Symptoms were resolved over the next three weeks. MRI showed resolution of bone marrow oedema in foot and ankle bones (Figure 5).
Discussion

Pregnancy can alter the course of some inflammatory disorders. It can also lead to the emergence of mechanical joint diseases, such as reflex sympathetic dystrophy of the lower extremities (7). Reflex sympathetic dystrophy is a disease with a wide range of clinical manifestations. Thus, before it was recognized as a single disease, it received a large number of names, such as algodystrophy, Sudeck atrophy, shoulder-hand syndrome, and causalgia. Transient osteoporosis has also been thought by many authors to be a form of reflex sympathetic dystrophy. It has been proposed that an objective differentiation was not possible between them (8). Although pathogenetic mechanism of the disease is not totally clear, sympathetic regulation of micro vessels is certainly (9).

Transient osteoporosis in pregnancy was first described by Curtiss and Kincaid (10) in three cases of transient demineralization of the hip during pregnancy. Lequesne (4) in 1968 described the same pathology and called it transient osteoporosis or RSD. De Marchi et al. (11) reported five men and a woman with the same clinical picture. Later, additional reports of cases have been appeared and further defined, but the different terms was used to determine the same condition in each case. In 1994, a working group of the International Association for the Study of Pain (IASP) developed a consensus definition and proposed a new terminology. Thus, the term complex regional pain syndrome (CRPS) type I replaces the name RSD, and the term CRSP type II, which requires demonstrable peripheral injury, replaces the term causalgia (12).

The pathophysiology of transient osteoporosis which occurs in pregnancy is contributed to microtrauma in weight-bearing bones or chemical alterations. It has been proposed regarding the pathogenetic mechanism which an unknown stimulus activates a large number of bone turnover foci in the involved joint, initiating intensive osteoclast resorption. (13,14). There are relatively rare reports on this issue, but the association of RSD and pregnancy does not randomly appear. Factors which contribute to RSD related to pregnancy have not been elucidated. During pregnancy, significant weight gain, hyperlordosis, hypertriglyceridemia, vascular stasis induced by the compression on the inferior vena cava may be risk factors for RSD (7).

The relation between RSD and transient osteoporosis remains controversial, but pathophysiology and clinical findings appears to be same. However, RSD often associated with sensory, motor, and autonomic disturbances as well as trophic changes, such as increased hair growth, redness, and warmth (15). But, patients with transient osteoporosis presented with only pain and osteoporosis. We preferred to name it as RSD of foot in pregnancy because the clinical findings of our patient met the diagnostic criteria for CRPS type I.

We described the pregnant patient with RSD of the foot and ankle. We thought that RSD should be included in the differential diagnosis of patients with acute foot and ankle pain. Failure to recognize this unusual cause of pain can result in delayed or mistaken diagnosis especially during pregnancy. The recognition of this condition provides us to avoid invasive investigations in pregnant women.

References


