

Aseptic Arthritis of the Bilateral Temporomandibular Joint Mimicking Rheumatological Diseases

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A 24-year-old male patient was admitted to the Otorhinolaryngology Department with complaints of pain for three days in the bilateral preauricular region. The pain was constant, severe, and increased when he reclined, chewed or talked. There was no medical history of fever or trauma. Physical examination findings included mild edema, allodynia, and hyperalgesia in the bilateral preauricular area and bilateral masseter muscle. Panoramic radiograph showed limited interincisal mouth opening (12 mm) (Figure 1a). A written informed consent was obtained from the patient.

Initial laboratory tests showed serum C-reactive protein of 154 mg/L (0-5), ferritin of 1314 ng/mL (15-200), leukocyte count of $22.1 \times 10^3/\mu\text{L}$ (4-10), hemoglobin of 10.1 g/dL (10-17), and total bilirubin of 2.31 mg/dL (0-1.1). The test for rheumatoid factor was negative. Significant inflammatory processes were seen on magnetic resonance imaging in T₂ (Figure 1b) and necrotic areas in the right ramus of the mandible were also seen as a hypointense signal in T₂ (Figure 1c). Arthritis in the bilateral condylar head secondary to a sickle cell crisis was diagnosed after the analysis of the laboratory tests and images.

The patient was admitted to the hematology ward and received standard analgesia recommended for the treatment of severe pain consisting of nonsteroidal antiinflammatory

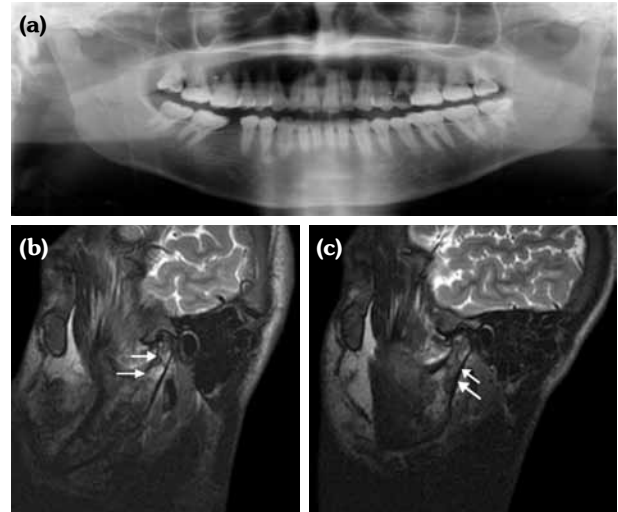


Figure 1. Panoramic radiograph imaging (a), magnetic resonance imaging of left temporomandibular joint showing a signal change at condylar head and neck of bilateral jaw, suggesting bone edema with T₂ hypersensitivity (b) magnetic resonance imaging of right temporomandibular joint showing bilateral ramus infarction as a hypointense signal in T₂ (c), (arrows).

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drugs and parenteral opioid analgesia. The pain gradually subsided after four days of hospital stay, allowing progressive weaning from the analgesic treatment. Mouth opening amplitude improved substantially to 40 mm at six months.

Almost half of sickle cell patients experience orofacial pain.¹ Limb joint pain is common, but temporomandibular joint (TMJ) pain is rare.² Clinicians should be aware that aseptic arthritis may occasionally be caused by sickle cell disease.³ The differential diagnosis should include avascular necrosis of the condylar head, manifestation of rheumatoid arthritis and juvenile rheumatoid arthritis in the TMJ, trauma and lupus arthritis of the TMJ.⁴

Laboratory tests and diagnostic imaging remain important for proper assessment due to the similarity of clinical manifestations of TMJ arthritis. Sickle cell disease may cause TMJ arthritis, and should be considered although rarely seen in the differential diagnosis of TMJ arthritis.

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REFERENCES

1. O'Rourke CA, Hawley GM. Sickle cell disorder and orofacial pain in Jamaican patients. *Br Dent J* 1998;185:90-2.
2. Sansevere JJ, Milles M. Management of the oral and maxillofacial surgery patient with sickle cell disease and related hemoglobinopathies. *J Oral Maxillofac Surg* 1993;51:912-6.
3. Demirbaş AK, Ergün S, Güneri P, Aktener BO, Boyacıoğlu H. Mandibular bone changes in sickle cell anemia: fractal analysis. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2008;106:e41-8.
4. Fernandes EG, Savioli C, Siqueira JT, Silva CA. Oral health and the masticatory system in juvenile systemic lupus erythematosus. *Lupus* 2007;16:713-9.