

# Nocardial pyomyositis

Boby Varkey Maramattom, Rony K. Varghese, Anila Sudhakaran<sup>1</sup>, Kurian Ninan<sup>2</sup>

Departments of Neurology, <sup>1</sup>Emergency Medicine and <sup>2</sup>Radiology, Aster Medcity, Kochi, Kerala, India

For correspondence:

Dr. Boby Varkey Maramattom, Department of Neurology,  
Aster Medcity, Kothad, Kochi - 682 027, Kerala, India.  
E-mail: bobvarkey@gmail.com

Ann Indian Acad Neurol 2017;20:58-58

A 70-year-old male with Type II diabetes underwent plasmapheresis through the internal jugular vein 2 months ago for chronic inflammatory demyelinating polyneuropathy. Subsequently, he was on high-dose prednisolone and azathioprine. He presented to the emergency department with severe right thigh pain for 2 weeks. Examination revealed a fluctuant tender swelling of 4 cm × 7 cm size in the upper anterior thigh and antalgic weakness of adduction of right thigh. Ultrasound showed a well-defined intramuscular cystic lesion in the medial aspect of the right thigh measuring 8 cm × 4 cm. Magnetic resonance imaging showed a well-circumscribed ovoid-shaped T2-weighted homogeneously hyperintense and T1-weighted hypointense lesion in the right adductor longus muscle, measuring 5.4 cm × 4.2 cm × 8.7 cm [Figure 1]. The lesion showed avid peripheral enhancement and thin enhancing septae and did not encase the underlying femoral artery. Ultrasound-guided aspiration revealed frank pus. Gram smear [Figure 2] and culture showed Gram-positive filamentous bacteria which were weakly acid fast (*Nocardia* sp.). He was initiated on treatment with trimethoprim-sulfamethoxazole for 3 months with good improvement.

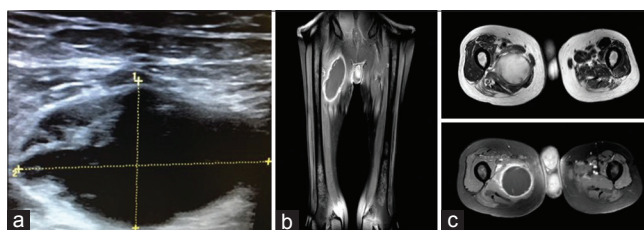
Pyomyositis (tropical pyomyositis) is an uncommon primary infection of skeletal muscles. Seventy-five percent of cases occur in immunocompromised individuals, the thigh being a common site.<sup>[1]</sup> *Staphylococcus aureus* (90%) accounts for the majority of cases. Nocardial pyomyositis is extremely rare but responds well to treatment if initiated early.<sup>[2]</sup> Skeletal muscles are very resistant to infection; however, our patient could have developed nocardial pyomyositis from transient bacteremia during plasmapheresis as there was no evidence of systemic nocardiosis.

**Financial support and sponsorship**  
Nil.

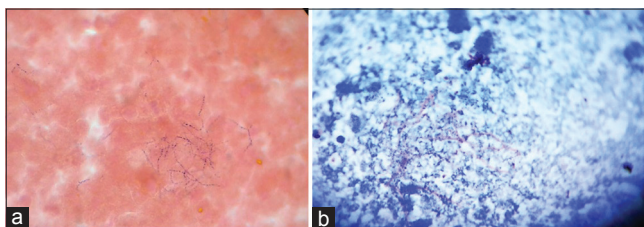
**Conflicts of interest**

There are no conflicts of interest.

Access this article online	
Quick Response Code: 	Website: www.annalsofian.org
	DOI: 10.4103/0972-2327.199915



**Figure 1:** (a) Ultrasound showing 8 cm × 4 cm cystic lesion. (b) Coronal T1-weighted magnetic resonance imaging. (c) Axial T2 magnetic resonance imaging. (d) Axial T1 with contrast. A well-circumscribed ovoid-shaped altered signal intensity lesion in the right adductor longus muscle



**Figure 2:** (a) Gram stain and (b) 1% acid-fast stain showing filamentous bacteria

**References**

1. Agarwal V, Chauhan S, Gupta RK. Pyomyositis. Neuroimaging Clin N Am 2011;21:975-83.
2. Sagar V, Pinto B, Lal A, Kumar M, Rathi M, Sharma K, et al. Nocardia pyomyositis in a patient with granulomatosis with polyangiitis. Int J Rheum Dis 2015. doi: 10.1111/1756-185X.12623. [Epub ahead of print].

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

**For reprints contact:** reprints@medknow.com

**How to cite this article:** Maramattom BV, Varghese RK, Sudhakaran A, Ninan K. Nocardial pyomyositis. Ann Indian Acad Neurol 2017;20:58.

**Received:** 20-10-16, **Revised:** 23-11-16, **Accepted:** 15-12-16