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Monomelic amyotrophy with proximal upper limb ...

[https://jmedicalcasereports.biomedcentral.com/...](https://jmedicalcasereports.biomedcentral.com/) ▾

Mar 17, 2016 · Monomelic amyotrophy is an uncommon, benign, unilateral disorder of the lower motor neurons, affecting predominantly the hand and forearm muscles. Proximal involvement of the arm and shoulder muscles is an unusual presentation that has been rarely reported in the literature. A 28-year-old white man presented with insidious-onset, slowly progressive, unilateral weakness and atrophy of his ...

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Author: Eman Al-Ghawi, Talal Al-Harbi, Adnan Al-...

Publish Year: 2016

Dominant spinal muscular atrophy with lower extremity ...

<https://www.researchgate.net/publication/45583551...>

Spinal muscular atrophy with lower limb predominance (SMA-LED) is an early-onset static or slowly progressive disorder, characterized by proximal muscle weakness and atrophy predominantly ...

Amyotrophic Lateral Sclerosis (ALS) Fact Sheet | National ...

<https://www.ninds.nih.gov/Disorders/Patient-Caregiver-Education/Fact-Sheets/...> ▾

However, the presence of upper and lower motor neuron symptoms strongly suggests the presence of the disease. Physicians will review an individual's full medical history and conduct a neurologic examination at regular intervals to assess whether symptoms such as muscle weakness, atrophy of muscles, and spasticity are getting progressively worse.

(PDF) Monomelic amyotrophy with proximal upper limb ...

<https://www.researchgate.net/publication/297725844...>

Proximal involvement of the arm and shoulder muscles is an unusual presentation that has been rarely reported in the literature. Case presentation A 28-year-old white man presented with insidious ...

Ankylosing spondylitis and central core disease - Case report

<https://www.researchgate.net/publication/9077663...>

We report the case of a 45-year-old man with clinical and radiological diagnosis of AS and proximal muscular weakness in the lower limbs. Needle electromyography showed myopathic features and ...

Congenital Segmental Spinal Muscular Atrophy: A Case Report

Muscular atrophy and weakness in the lower extremities in Behçet's



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Muscular involvement in Behçet's disease: Case report and ...<https://www.sciencedirect.com/science/article/pii/S0960896696000193>

Aug 01, 1996 · Despite its rare occurrence, our **case report** and the review of the literature suggest that the diagnosis of BD should be considered particularly in younger patients presenting with **muscular** symptoms like pain and **swelling** predominantly of the **lower extremities**. CASE REPORT A 22-yr-old white male patient presented for one week with genital ulcers, dolent and fluctuating **swelling** of soft tissue, myalgia in the **lower extremities** ...

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Author: F. Worthmann, J. Bruns, T. Türker, G....

Publish Year: 1996

(PDF) Neuro-Behçet's disease in childhood: A focus on the ...https://www.researchgate.net/publication/235381981_Neuro-Behcet

Jan 29, 2013 · **Behçet disease** reported in the literature from 1971 to ... **muscular atrophy** in **Behçet's disease**. ... Cerebral sinus thrombosis in **Behçet disease**: case report and review of.

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What is the clinical manifestation of spinal muscular atrophy? ▾

When does spinal muscular atrophy occur in children? ▾

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A novel case report of spinal muscular atrophy with ...<https://www.dovepress.com/a-novel-case-report-of...> ▾

May 30, 2019 · Introduction. Spinal **muscular atrophy** (SMA) is a heterogeneous **disease** characterized by degeneration of the **lower** motor neurons and is usually linked to mutations of SMN-1 gene. The condition often presents by decreased motor function and muscle **atrophy** due to degeneration of anterior horn cell in the spinal cord and motor cells of **lower** cranial ...

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Author: Reza Shervin Radv Yalda Nilinour S

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Muscular atrophy and weakness in the lower extremities in Behçet's



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Aug 01, 1996 · Despite its rare occurrence, our case report and the review of the literature suggest that the diagnosis of BD should be considered particularly in younger patients presenting with muscular symptoms like pain and swelling predominantly of the lower extremities. CASE REPORT A 22-yr-old white male patient presented for one week with genital ulcers, dolent and fluctuating swelling of soft tissue, myalgia in the lower extremities ...

Cited by: 22

Author: F. Worthmann, J. Bruns, T. Türker, G. Go...

Publish Year: 1996

[\[PDF\] Hirayama Disease - A Not So Foreign Entity: Case Report ...](#)

https://www.clinicsinsurgery.com/pdfs_folder/cis-v6-id3137.pdf

Hirayama Disease (HD), also known as non progressive juvenile spinal muscular atrophy, is a form of juvenile muscular atrophy. It is a rare self-limiting cervical flexion myelopathy with insidious onset of progressive weakness and muscular wasting of the upper extremity [4,6]. It typically affects young individuals in their second to third

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Feedback

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[Bimelic Hirayama Disease: Clinical Dilemma Solved by Imaging](#)

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3625547>