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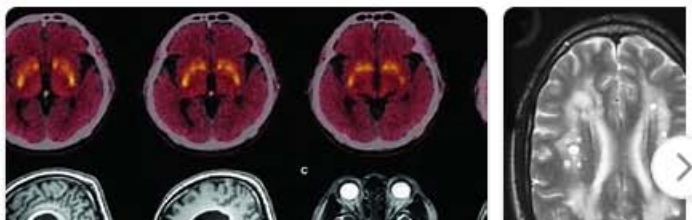
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A genetically confirmed case of **spinocerebellar ataxia type 3 (SCA 3)**, presenting with disabling foot dystonia, peripheral neuropathy, and minimal cerebellar signs is reported. The **dystonia** improved...

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Spinocerebellar ataxia type 3 with dopamine-responsive dystonia: A case report

Zhang X *et al.* Rare SCA3 with dopamine-responsive dystonia

Xiao-Le Zhang, Xiao-Bo Li, Fa-Feng Cheng, Shu-Ling Liu, Wen-Chao Ni, Fei-Fei Tang,
Qing-Guo Wang, Xue-Qian Wang

Abstract

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Spinocerebellar ataxia type 3 with dopamine-responsive dystonia:



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A genetically confirmed **case of spinocerebellar ataxia type 3 (SCA 3)**, presenting with disabling foot dystonia, peripheral neuropathy, and minimal cerebellar signs is reported. The **dystonia** improved dramatically with **levodopa** treatment in the absence of additional parkinsonian feature. A trial of **levodopa** for **dystonia** in **SCA 3** may be of therapeutic

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