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Late-onset multiple acyl-CoA dehydrogenase deficiency (MADD) is an autosomal recessive inherited disease of metabolic dysfunction clinically characterized by fluctuating proximal muscle weakness, exercise intolerance, and dramatic riboflavin responsiveness. Dropped head syndrome can occasionally be ...

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A Case of Late Onset Riboflavin-responsive Multiple Acyl-CoA Dehydrogenase Deficiency Manifesting as Recurrent Rhabdomyolysis and Acute Renal Failure Rumikolzumi

1,NaokiSuzuki,MariNagata1,TakafumiHasegawa1,YuAbe2,YukaSaito2,

HiroshiMochizuki3,MakiTateyama1 andMasashiAoki1 Abstract We report an adult case of late-onset ...

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<https://www.sciencedirect.com/science/article/pii/S003537871600432X>

Late-onset multiple acyl-CoA dehydrogenase deficiency (MADD) is a rare, treatable, beta-oxidation disorder responsible for neuromuscular symptoms in adults. This case series describes the clinical and biochemical features of 13 French patients with late-onset MADD.

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Jul 24, 2015 · **Multiple acyl-CoA dehydrogenase deficiency** ... it is the first **case report** of late-onset MADD presenting with bent spine syndrome as the initial symptom. ... Wang N, Wu ZY. Molecular analysis of 51 unrelated pedigrees with late-onset **multiple acyl-CoA dehydrogenation deficiency** (MADD) in southern China confirmed the most common ETFDH mutation ...

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We report an adult case of **late-onset riboflavin-responsive multiple acyl-CoA dehydrogenase deficiency**

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