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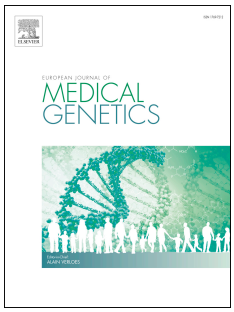

A *de novo* mutation in *RPL10* causes a rare X-linked ribosomopathy characterized by syndromic intellectual disability and epilepsy: A new case and review of the literature

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CLINICAL REPORT

A *de novo* mutation in *RPL10* causes a rare X-linked ribosomopathy characterized by syndromic intellectual disability and epilepsy: a new case and review of the literature

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Running Title:

Rare ribosomopathy caused by mutation in *RPL10*