

## Opis choroby \*

### Definicja

A rare mitochondrial disease characterized by adult onset of progressive external ophthalmoplegia, exercise intolerance, muscle weakness, manifestations of spinocerebellar ataxia (e.g. impaired gait, dysarthria) and mild motor peripheral neuropathy. Respiratory insufficiency has been reported in some cases.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Adult-onset CPEO with mitochondrial myopathy  
CPEO o początku w wieku dorosłym z miopatią mitochondrialną

#### Kod ORPHA

329336

#### Kod OMIM

616479

#### Kod ICD10

G71.3

#### Kod ICD11

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#### \*Źródło

orphanet