

## Opis choroby \*

### Definicja

A muscular channelopathy presenting with a pure myotonia dramatically aggravated by potassium ingestion, with variable cold sensitivity and no episodic weakness. This group includes three forms: myotonia fluctuans, myotonia permanens, and acetazolamide-responsive myotonia.

### Dane

#### Klasyfikacja

Grupa fenomenów

#### Synonimy

K+-aggravated myotonia  
K+-aggravated myotonia  
K-aggravated myotonia  
PAM  
K-aggravated myotonia  
PAM

#### Kod ORPHA

612

#### Kod OMIM

608390

#### Kod ICD10

G71.1

#### Kod ICD11

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\*Źródło

orphanet