

## Opis choroby \*

### Definicja

A rare, genetic, neuromuscular disease characterized by adult-onset muscle weakness and atrophy in a scapulo-peroneal distribution, mild involvement of the facial muscles, dysphagia, and gynecomastia. Elevated serum CK levels and mixed myopathic and neurogenic abnormalities are associated clinical findings.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Kaeser syndrome

Zespół Kaesera

Zespół Starka i Kaesera

Stark-Kaeser syndrome

#### Kod ORPHA

85146

#### Kod OMIM

181400

#### Kod ICD10

G12.1

#### Kod ICD11

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#### \*Źródło

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