

## Opis choroby \*

### Definicja

A rare, genetic, neuromuscular disease characterized by adult-onset muscle weakness and atrophy in a scapuloperoneal 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	Synonimy	
Choroba	Kaeser syndrome	
	Zespół KAESERA	
	Zespół Starka i KAESERA	
	Stark-Kaeser syndrome	
Kod ORPHA	Kod OMIM	Kod ICD10
85146	181400	G12.1

### Kod ICD11

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\*Źródło

orphanet