

## Opis choroby \*

### Definicja

A rare, genetic neuromuscular disease characterized by progressive external ocular, facial and pharyngeal muscle weakness, leading to variable degrees of ptosis, ophthalmoparesis, facial muscle atrophy, dysarthria and dysphagia, as well as distal muscle weakness and atrophy of lower and upper extremities. Respiratory muscle involvement is common, but sensorineural hearing loss, asymmetrical extremity weakness and severe proximal weakness are rare.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Choroba	OPDM Miopatia oczno-gardłowa dystalna OPDM Oculopharyngeal distal myopathy

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
98897	618940	G71.0

**Kod ICD11**  
9C82.1

---

### \*Źródło

orphanet