

## Opis choroby \*

### Definicja

A rare premature aging syndrome characterized by atrophy of the skin and subcutaneous tissue involving predominantly the distal parts of the extremities, resulting in prematurely aged appearance of the hand and feet. Another prominent feature is the characteristic facies with hollow cheeks, beaked nose, and owl-like eyes. Additional, non-dermatological manifestations, like bone anomalies have been described in some patients. Mode of inheritance has not been definitively established.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Acrogeria, Gottron type
	Akrogeria, typu Gottrona
	Akrometageria
	Zespół Gottrona
	Acrometageria
	Gottron syndrome

**Kod ORPHA**  
2500

**Kod OMIM**  
201200

**Kod ICD10**  
L90.8

**Kod ICD11**  
LD2B

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### \*Źródło

orphanet