

# Zespół progerii Hutchinsona i Gilforda

## Kod Orpha: 740 Kod OMIM: 176670

### Opis choroby \*

#### Definicja

Hutchinson-Gilford progeria syndrome is a rare, fatal, autosomal dominant and premature aging disease, beginning in childhood and characterized by growth reduction, failure to thrive, a typical facial appearance (prominent forehead, protuberant eyes, thin nose with a beaked tip, thin lips, micrognathia and protruding ears) and distinct dermatologic features (generalized alopecia, aged-looking skin, sclerotic and dimpled skin over the abdomen and extremities, prominent cutaneous vasculature, dyspigmentation, nail hypoplasia and loss of subcutaneous fat).

#### Dane

Klasyfikacja	Synonimy
--------------	----------

Choroba	HGPS
	Progeria
	Progeria

Kod ORPHA	Kod OMIM	Kod ICD10
-----------	----------	-----------

740	176670	E34.8
-----	--------	-------

Kod ICD11
-----------

LD2B
------

---

\*[Źródło](#)

[orphanet](#)

### Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie [www.orphanet.pl](http://www.orphanet.pl)