

Opis choroby *

Definicja

A rare retinal vasoproliferative disease affecting preterm infants characterized initially by a delay in physiologic retinal vascular development and compromised physiologic vascularity, and subsequently by aberrant angiogenesis in the form of intravitreal neovascularization.

Dane

Klasyfikacja

Choroba

Synonimy

ROP

Retrolental fibroplasia

ROP

Retrolental fibroplasia

Kod ORPHA

90050

Kod OMIM

133780

Kod ICD10

H35.1

Kod ICD11

9B71.3

*Źródło

orphanet