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

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*Źródło

orphanet