Opis choroby *

Definicja

A rare congenital nephrotic syndrome characterized by massive protein loss and marked edema manifesting in utero or during the first 3 months of life.

Dane

Klasyfikacja
Choroba

Synonimy
Finnish congenital nephrosis
Fińska wrodzona nefroza

Kod ORPHA
839

Kod OMIM
256300

Kod ICD10
N04.8

Kod ICD11
GB4Z

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*Źródło

orphanet