

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

*Źródło

orphanet