

Opis choroby *

Definicja

A rare genetic disease that belongs to the group of extreme insulin-resistance syndromes and is due to autoantibodies directed against insulin receptor.

Dane

Klasyfikacja

Choroba

Kod ORPHA

2298

Kod OMIM

-

Kod ICD10

E34.8

Kod ICD11

5A44

*Źródło

orphanet