Opis choroby *

Definicja

A rare primary immunodeficiency characterized by recurrent and/or invasive bacterial, viral, and fungal infections, associated with low to absent blood IgM levels, while IgG, IgG subclasses, and IgA levels, as well as IgG antibody response to vaccinations, are normal. Patients may also present allergic diatheses, and the prevalence of autoimmune diseases is increased.

Dane

Klasyfikacja Synonimy

Choroba Selective immunoglobulin M deficiency

Selektywny Niedobór immunoglobuliny M

Kod ORPHA Kod OMIM Kod ICD10

331235 - D80.4

Kod ICD11 4A01.04

*Źródło

orphanet