

Opis choroby *

Definicja

Omenn syndrome (OS) is an inflammatory condition characterized by erythroderma, desquamation, alopecia, chronic diarrhea, failure to thrive, lymphadenopathy, and hepatosplenomegaly, associated with severe combined immunodeficiency (SCID; see this term).

Dane

Klasyfikacja

Choroba

Synonimy

Combined immunodeficiency with hypereosinophilia

Złożony Niedobór odporności z hipereozynofilią

Kod ORPHA

39041

Kod OMIM

603554

Kod ICD10

D81.8

Kod ICD11

4A01.10

*Źródło

orphanet