Opis choroby *

Definicja

Severe combined immunodeficiency (SCID) comprises a group of rare monogenic primary immunodeficiency disorders characterized by a lack of functional peripheral T lymphocytes resulting in early-onset severe respiratory infections and failure to thrive. They are classified according to immunological phenotype into SCID with absence of T cells but presence of B cells (T-B+ SCID) or SCID with absence of both (T-B- SCID) (see these terms). Both of these groups include several forms, with or without natural killer (NK) cells.

Dane

Klasyfikacja
Grupa fenomenów

Synonimy
SCID
SCID

Kod ORPHA
183660

Kod OMIM
-

Kod ICD10
D81.1

Kod ICD11
4A01.10

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*Źródło

orphanet