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 Synonimy

Grupa fenomenów SCID

SCID

Kod ORPHA Kod OMIM Kod ICD10

183660 - D81.1

Kod ICD11 4A01.10

*Źródło

orphanet