Severe combined immunodeficiency due to IKK2 deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Severe combined immunodeficiency due to IKK2 deficiency. ORPHA:397787

Severe combined immunodeficiency due to IKK2 deficiency is a rare, genetic form of primary immunodeficiency characterized by life-threatening bacterial, fungal and viral infections with the onset in infancy, and failure to thrive. Typically, hypogammaglobulinemia or agammaglobulinemia and normal levels of T and B cells are present.