Omenn syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Omenn syndrome. ORPHA:39041

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).