Letterer-Siwe disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Letterer-Siwe disease. ORPHA:99870

Letterer-Siwe syndrome (LSS) is the acute disseminated multisystem form of Langerhans cell histiocytosis (see this term) characterised by proliferation of nonlipid histiocytes in the viscera and bones.