Cernunnos-XLF deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Cernunnos-XLF deficiency. ORPHA:169079

Cernunnos-XLF deficiency is a rare form of combined immunodeficiency characterized by microcephaly, growth retardation, and T and B cell lymphopenia.