X-linked agammaglobulinemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked agammaglobulinemia. ORPHA:47

X-linked agammaglobulinemia (XLA) is a clinically variable form of isolated agammaglobulinemia, an inherited immunodeficiency disorder (see this term), and is characterized in affected males by recurrent bacterial infections during infancy.