Autosomal agammaglobulinemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal agammaglobulinemia. ORPHA:33110

Agammaglobulinemia, non-Bruton type (autosomal agammaglobulinemia) is a rare form of agammaglobulinemia, a primary immunodeficiency disease, and is characterized by variable immune dysfunction with frequent and recurrent bacterial infections and/or chronic diarrhea.