Autosomal dominant aplasia and myelodysplasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant aplasia and myelodysplasia. ORPHA:314399

Autosomal dominant aplasia and myelodysplasia is a rare, genetic, hematologic disorder characterized by bone marrow failure which manifests with aplastic anemia and/or myelodysplasia, associated with hearing/ear abnormalities (such as deafness, labyrinthitis), inherited in an autosomal dominant manner.