
DISEASE:
Autosomal dominant aplasia and myelodysplasia

NAME:	Autosomal dominant aplasia and myelodysplasia
DESCRIPTION:	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.
ORPHACODE:	314399
SYNONYMS:	Autosomal dominant aplastic anemia and myelodysplasia
XREF(S):	<u>Orphanet</u> <u>ICD-10</u> <u>OMIM</u>
ANALYTE(S):	<u>SRP72</u>
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