

DISEASE:
Achondrogenesis type 1A

NAME:	Achondrogenesis type 1A
DESCRIPTION:	A rare, lethal type of achondrogenesis characterized by dwarfism with extremely short limbs, narrow chest, short ribs that are easily fractured, soft skull bones and distinctive histological features of the cartilage.
ORPHACODE:	93299
SYNONYMS:	Achondrogenesis, Houston-Harris type
XREF(S):	Orphanet ICD-10 OMIM MeSH
ANALYTE(S):	TRIP11
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/3164>

RELATED CONTENT

Related Analytes

- [thyroid hormone receptor interactor 11](#)

Source URL: <http://gentest.healthdata.be/disease/3164>