

DISEASE:
Isolated osteopoikilosis

NAME:	Isolated osteopoikilosis
DESCRIPTION:	A rare primary bone dysplasia characterized by multiple, small, round to ovoid osteosclerotic foci with a predilection for the epiphyses and metaphyses of long tubular bones as well as the pelvis, scapula, carpal, and tarsal bones. The condition is usually clinically silent and discovered only incidentally, although some patients may experience mild articular pain with or without joint effusion. Bone strength is normal.
ORPHACODE:	166119
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	LEMD3
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Buschke-Ollendorff / Melorheostosis with Osteopoikilosis

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- LEM domain containing 3

Source URL: <http://gentest.healthdata.be/disease/65>