DISEASE:
Multiple osteochondromas

NAME:	Multiple osteochondromas
DESCRIPTION:	A primary bone disorder characterized by development of two or more cartilage capped bony outgrowths (osteochondromas) at the surface of the bones.
ORPHACODE:	321
SYNONYMS:	Bessel-Hagen disease Multiple cartilaginous exostoses
XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM
ANALYTE(S):	EXT1 EXT2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Multiple osteochondromas (2 genes)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- exostosin glycosyltransferase 1
- exostosin glycosyltransferase 2

Related Gene Panels

- Multiple osteochondromas (2 genes) - UZA