GENETIC TEST:
Achondrogenesis / Kniest dysplasia / Hypochondrogenesis

FULL NAME:	Achondrogenesis / Kniest dysplasia / Hypochondrogenesis
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	8 weeks
CREATED:	31 Jul 2019 - 10:53
CHANGED:	31 Jan 2023 - 13:21

Source URL: http://gentest.healthdata.be/genetic_test/318

RELATED CONTENT

Related Diseases

- Achondrogenesis type 2
- Hypochondrogenesis
- Kniest dysplasia
- Multiple epiphyseal dysplasia, Beighton type
- Spondyloepiphyseal dysplasia congenita
- Stickler syndrome type 1

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- collagen type II alpha 1 chain