GENETIC TEST:
Osteogenesis Imperfecta (gene panel)

FULL NAME:	Osteogenesis Imperfecta (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Whole Exome Sequencing (WES)
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	14 Dec 2022 - 16:30
CHANGED:	23 Aug 2023 - 09:33
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB/16397

Source URL: http://gentest.healthdata.be/genetic_test/1095

RELATED CONTENT

Related Diseases

- Autosomal dominant osteopetrosis type 1
- Bruck syndrome
- Classical Ehlers-Danlos syndrome
- Ehlers-Danlos/osteogenesis imperfecta syndrome
- Gnathodiaphyseal dysplasia
- High bone mass osteogenesis imperfecta
- Osteogenesis imperfecta
- Osteogenesis imperfecta type 1
- Osteogenesis imperfecta type 2
- Osteogenesis imperfecta type 3
- Osteogenesis imperfecta type 4

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- anoctamin 5
- bone morphogenetic protein 1
- collagen type I alpha 1 chain
- collagen type I alpha 2 chain
- cAMP responsive element binding protein 3 like 1

- cartilage associated protein
- FKBP prolyl isomerase 10
- interferon induced transmembrane protein 5
- LDL receptor related protein 5
- membrane bound transcription factor peptidase, site 2
- prolyl 3-hydroxylase 1
- prolyl 4-hydroxylase subunit beta
- procollagen-lysine,2-oxoglutarate 5-dioxygenase 2
- plastin 3
- peptidylprolyl isomerase B
- SEC24 homolog D, COPII coat complex component
- serpin family F member 1
- serpin family H member 1
- sphingomyelin synthase 2
- Sp7 transcription factor
- secreted protein acidic and cysteine rich
- terminal nucleotidyltransferase 5A
- transmembrane protein 38B
- Wnt family member 1
- xylosyltransferase 2

Related Gene Panels

- Osteogenesis Imperfecta (25 genes) - KUL
