GENETIC TEST:
Corneal dystrophy (gene panel)

FULL NAME:	Corneal dystrophy (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Whole Exome Sequencing (WES) Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
TURNAROUND TIME (MAXIMUM):	6 months
CREATED:	03 Aug 2020 - 09:07
CHANGED:	16 Dec 2022 - 10:57

Source URL: http://gentest.healthdata.be/genetic_test/1024

RELATED CONTENT

Related Diseases

- AGel amyloidosis
- Amyotrophic lateral sclerosis
- Axenfeld-Rieger syndrome
- Bietti crystalline dystrophy
- Brittle cornea syndrome
- CINCA syndrome
- Classical Ehlers-Danlos syndrome
- Congenital cornea plana
- Congenital hereditary endothelial dystrophy type II
- Congenital stromal corneal dystrophy
- Corneal dystrophy-perceptive deafness syndrome
- Corneal intraepithelial dyskeratosis-palmoplantar hyperkeratosis-laryngeal dyskeratosis syndrome
- Epithelial recurrent erosion dystrophy
- Familial LCAT deficiency
- Familial cold urticaria
- Fish-eye disease
- Fleck corneal dystrophy
- Fuchs endothelial corneal dystrophy
- Gelatinous drop-like corneal dystrophy
- Granular corneal dystrophy type I
- Granular corneal dystrophy type II
- Intermediate generalized junctional epidermolysis bullosa
- Isolated aniridia
- Isolated congenital megalocornea
- Late-onset junctional epidermolysis bullosa
- Lattice corneal dystrophy type I

- Localized junctional epidermolysis bullosa
- Macular corneal dystrophy
- Meesmann corneal dystrophy
- Muckle-Wells syndrome
- Pitt-Hopkins syndrome
- Posterior polymorphous corneal dystrophy
- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB
- Reis-Bücklers corneal dystrophy
- Ring dermoid of cornea
- Schnyder corneal dystrophy
- Thiel-Behnke corneal dystrophy

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- AGBL carboxypeptidase 1
- chordin like 1
- carbohydrate sulfotransferase 6
- collagen type XVII alpha 1 chain
- collagen type III alpha 1 chain
- collagen type V alpha 1 chain
- collagen type VIII alpha 2 chain
- cytochrome P450 family 4 subfamily V member 2
- decorin
- grainyhead like transcription factor 2
- gelsolin
- keratocan

- keratin 12
- keratin 3
- lecithin-cholesterol acyltransferase
- lipoxygenase homology PLAT domains 1
- NLR family pyrin domain containing 1
- NLR family pyrin domain containing 3
- ovo like zinc finger 2
- paired box 6
- phosphoinositide kinase, FYVE-type zinc finger containing
- paired like homeodomain 2
- PR/SET domain 5
- solute carrier family 4 member 11
- superoxide dismutase 1
- steroid sulfatase
- tumor associated calcium signal transducer 2
- transcription factor 4
- transforming growth factor beta induced
- UbiA prenyltransferase domain containing 1
- visual system homeobox 1
- zinc finger E-box binding homeobox 1
- zinc finger protein 469

Related Gene Panels

- Corneal dystrophy - UGent

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