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Full name:	Glaucoma - UGent
Type of panel:	in-house
Version number:	v2
Laboratory:	Centrum Medische Genetica - UZ Gent
Created:	17 Dec 2019 - 17:01
Changed:	21 Dec 2022 - 08:03

Related Diseases

- Autosomal dominant vitreoretinochoroidopathy
- Cataract-glaucoma syndrome
- Congenital glaucoma
- Glaucoma secondary to spherophakia/ectopia lentis and megalocornea
- Glaucoma-ectopia lentis-microspherophakia-stiff joints-short stature syndrome
- Juvenile glaucoma
- MRCS syndrome
- Nanophthalmos
- Peters anomaly
- Walker-Warburg syndrome
- Weill-Marchesani syndrome

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ADAMTS10	99.99	1	
ADAMTS17	99.99	1	
ASB1	99.98	1	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
B3GLCT	99.90	1	
BEST1	99.86	1	
COL18A1	99.99	1	
COL4A1	99.99	1	
CPAMD8	99.97	1	
CREBBP	99.97	1	
CYP1B1	100.00	1	
FBN1	99.85	1	
FOXC1	100.00	1	
FOXD3	99.85	1	
FOXE3	99.29	1	
GJA1	100.00	1	
IFIH1	99.84	1	
LMX1B	100.00	1	
LTBP2	99.97	1	
MYOC	99.98	1	
NTF4	100.00	1	
OCRL	99.89	1	
OPTN	99.98	1	
PAX6	99.95	1	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
PITX2	99.98	1	
PITX3	100.00	1	
RIGI	99.84	1	
SBF2	99.77	1	
SH3PXD2B	100.00	1	
TBK1	99.07	1	
TEK	99.98	1	
WDR36	99.46	1	