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Full name:	Dyslipidemia (13 genes) - ULB
Laboratory:	Centre de Génétique Humaine - Erasme ULB
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Related Diseases

- Cholesteryl ester storage disease
- Familial GPIHBP1 deficiency
- Familial apolipoprotein A5 deficiency
- Familial apolipoprotein C-II deficiency
- Familial lipase maturation factor 1 deficiency
- Familial lipoprotein lipase deficiency
- Homozygous familial hypercholesterolemia
- Sitosterolemia
- Wolman disease

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ABCG5	100.00	1	In case of hypercholesterolemia / NM_022436.2
ABCG8	100.00	1	In case of hypercholesterolemia / NM_022437.2
APOA5	100.00	1	In case of hypertriglyceridemia / NM_052968.4
APOB	100.00	1	In case of hypercholesterolemia / NM_000384.2
APOC2	100.00	1	In case of hypertriglyceridemia / NM_000483.4
APOE	100.00	1	In case of hypercholesterolemia / NM_000041.3
GPIHBP1	100.00	1	In case of hypertriglyceridemia / NM_178172.5

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
LDLR	100.00	1	In case of hypercholesterolemia / NM_000527.4
LDLRAP1	100.00	1	In case of hypercholesterolemia / NM_015627.2
LIPA	100.00	1	In case of hypercholesterolemia / NM_000235.3
LMF1	100.00	1	In case of hypertriglyceridemia / NM_022773.3
LPL	100.00	1	In case of hypertriglyceridemia / NM_000237.2
PCSK9	100.00	1	In case of hypercholesterolemia / NM_174936.3