FAMILIAL HYPERCHOLESTEROLEMIA/DYSLIPIDEMIA REQUISITION

LAB USE ONLY	PATIENT INFORMATION (INCOMPLETE REQUESTS WILL BE BANKED)
Received date:	Name:
Notes:	Address:
SAMPLE COLLECTION	Date of Birth:
Date drawn:	Health Card No.:
EDTA blood (lavender top) (5ml at room temp)	Sex: M F Unknown Unspecified
DNA (100ng minimum) Other:	Birthsex: M F Unknown Unspecified
TEST REQUEST	ELIGIBILITY CRITERIA FOR TESTING
Familial Hypercholesterolemia- Focused Panel (8 genes) ABCG5, ABCG8, APOB, APOE, LDLR, LDLRAP1, LIPA, PCSK9	Individual must meet one or more of the following: 1. Confirmed FH disease-causing pathogenic/likely pathogenic variant in a close blood relative
Dyslipidemia - Comprehensive Panel (25 genes) ABCA1, ABCG5, ABCG8, ANGPTL3, APOA1, APOA5, APOB, APOC2, APOC3, APOE, CETP, GPD1, GPIHBP1, LCAT, LDLR, LDLRAP1, LIPA, LIPC, LMF1, LPL, MTTP, PCSK9, SAR1B, SCARB1, STAP1	2. High LDL-cholesterol level of ≥8.5 mmol/L at any age 3. Untreated elevated LDL-cholesterol level (not due to secondary causes) Specify: mmmol/L
Carrier testing/Known Family Mutation	Untreated LDL-cholesterol level ≥5.0 mmol/L for age 40 years and over Untreated LDL-cholesterol level ≥4.5 mmol/L for age 18 to 39 years
LHSC MD#/Name of index case in the family (include copy of report)	Untreated LDL-cholesterol level ≥3.5 mmol/L for age under 18 years
	AND at least one of the following: Tendon xanthomas and/or corneal arcus in proband
Date of Birth:	First-degree relative (FDR) with high LDL- cholesterol level (not due to secondary causes)
Relationship to this patient:	Proband or FDR with early onset ASCVD (men under 55 years; women under 65 years)
Gene: RefSeq:NM:	Limited family history information (e.g., adopted)
Mutation:	4. Clinical judgement:Criteria above not met, but suspicion remains
	Describe:
REFERRING PHYSICIAN Authorized Signature is	s Required CC REPORT TO
Physician Name (print):	Name:
Signature: Email:	Address:
Clinic/Hospital:	
Address:	Telephone:
Telephone: Fax:	Fax:

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