LIPID CONTROL - LEVEL 2 (LPD CONTROL 2)

Cat. No. LE2669 **Lot No.** 2257CH
Size: 5 x 1 ml **Expiry:** 2017-03

INTENDED USE

This product is intended for *in vitro* use in the quality control of Direct HDL, Direct LDL, Lipoprotein (a), Apolipoprotein A-1, Apolipoprotein B, Cholesterol and Triglyceride methods on clinical chemistry systems.

SAFETY PRECAUTIONS AND WARNINGS

Human source material, from which this product has been derived, has been tested at donor level for the Human Immunodeficiency Virus (HIV1 & HIV2) antibody, Hepatitis B surface antigen (HbsAg) and the Hepatitis C virus (HCV) antibody and found to be NON-REACTIVE. FDA approved methods have been used to conduct these tests. However, since no method can offer complete assurance as to the absence of infectious agents, this material and all patient samples should be handled as though capable of transmitting disease. For *in vitro* diagnostic use only.

STORAGE AND STABILITY

Unopened Lipid Control is stable until the expiry date printed on the product label, when stored between +2°C and +8°C. Once reconstituted, the components of the serum are stable for 7 days at +2°C to +8°C, and 4 weeks at -20°C when frozen once. The following exceptions apply: LP(a) is stable for 16 weeks at -20°C when frozen once. Values may drop by up to 10% for Direct LDL Cholesterol when stored for 4 weeks at -20°C.

PREPARATION FOR USE

Open the vial carefully, avoiding any loss of the material and reconstitute with 1 ml of distilled water. Replace the rubber stopper, close the vial and leave to stand for 30 minutes before use. Ensure that all traces of dry material are dissolved by swirling gently.

MATERIALS PROVIDED

Lipid Control - Level 2 5 x 1 ml

MATERIALS REQUIRED BUT NOT PROVIDED

Distilled water
Volumetric pipette

VALUE ASSIGNMENT

Each batch of Lipid Control is submitted to a number of external laboratories. Values are assigned from a consensus of results obtained by these laboratories and internal testing conducted at Randox Laboratories Ltd.

If a method is unavailable, contact Randox Laboratories - Technical Services, Northern Ireland, tel: +44 (0) 28 9445 1070 or email Technical.Services@randox.com

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Analyte	unit	Range target	low	high	methods
Apolipoprotein A-1	g/l	1.50	1.23	1.77	Immunoturbidimetric
	mg/dl	150	123	177	
	g/l	1.45	1.19	1.71	Nephelometric
	mg/dl	145	119	171	
Apolipoprotein B	g/l	1.15	0.94	1.36	Immunoturbidimetric
	mg/dl	115	94.3	136	
	g/l	1.19	0.98	1.40	Nephelometric
	mg/dl	119	97.6	140	
Cholesterol	mmol/l	5.87	5.11	6.63	Cholesterol Oxidase
	mg/dl	227	197	257	
	mmol/l	5.44	4.73	6.15	Siemens Dimension
	mg/dl	210	183	237	
HDL - Cholesterol	mmol/l	1.35	1.15	1.55	Direct Clearance Method
	mg/dl	52.1	44.4	59.8	
	mmol/l	1.07	0.75	1.39	Phosphotungstic acid pptn.
	mg/dl	41.3	28.9	53.7	
	mmol/l	1.25	1.06	1.44	Direct HDL Immunoseparation
	mg/dl	48.3	40.9	55.7	
	mmol/l	1.20	1.02	1.38	Direct HDL PEGME
	mg/dl	46.3	39.4	53.2	
	mmol/l	1.34	1.14	1.54	Direct HDL PPD
	mg/dl	51.7	44.0	59.4	
	mmol/l	1.20	1.02	1.38	Direct HDL Roche 3rd generation
	mg/dl	46.3	39.4	53.2	
	mmol/l	1.45	1.23	1.67	HDL - Ultra
	mg/dl	56.0	47.5	64.5	
LDL - Cholesterol	mmol/l	4.02	3.42	4.62	Direct Clearance Method
	mg/dl	155	132	178	
	mmol/l	3.54	3.01	4.07	Calculated
	mg/dl	137	116	158	
	mmol/l	3.48	2.96	4.00	Selective detergent methods
	mg/dl	134	114	154	
Lipoprotein (a)	mg/dl	19.6	15.7	23.5	Immunoturbidimetric
	nmol/l	45.2	36.2	54.2	
Triglycerides	mmol/l	2.33	1.96	2.70	Lipase/GPO-PAP no correction
	mg/dl	206	173	239	
	mmol/l	2.34	1.97	2.71	Lipase/GK UV no correction
	mg/dl	207	174	240	

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