

LIPOPROTEIN, VERY LOW DENSITY (VLDL; pre-ß-lipoprotein) from Human Plasma

Product No. L2264

ProductInformation

Product Description

HDL, LDL and VLDL are isolated sequentially from plasma by using the modified methods of Rudel, L.L.³ and Burstein, M⁴. Each lipoprotein is then concentrated and dialyzed extensively against 0.15 M NaCl, 0.01% EDTA, pH 7.4-7.5.

HDL and LDL are then filtered through a 0.2 μ membrane and VLDL is filtered through a 0.45 μ membrane.

Each lipoprotein class has a characteristic electrophoretic mobility and chemical composition. Each class is essentially free from contamination by other lipoprotein as determined by agarose electrophoresis using sudan black B staining for lipid. However, it is common for some serum proteins, foreign to the lipoprotein itself, to be present.

Storage

All lipoproteins should be stored at 2-8°C. Freezing may cause structural or composition changes.

Product Profile

Protein concentration: 0.3-1.2 mg/ml assayed by modified Lowry method using BSA as standard (see vial label for lot specific number).

Buffer: 0.15 M NaCl, 0.01% EDTA, pH 7.4-7.5

Source: Fresh, non-frozen plasma

Size: M.W. 6-27 x 10⁶; diameter 25-90 nm¹

Chemical composition: 88-95% lipid and 5-12% protein²

References

- 1. Mills, G.L., *et al.*, <u>A Guidebook to Lipoprotein</u>
 <u>Technique</u>, Elsevier:Amsterdam, New York and Oxford, 1984: p. 3.
- 2. Olivecrona, T., *et al.*, Biochim Biophys. Acta, **98**, 81 (1965)
- 3. Rudel, L.L., et al., Biochem. J., 139, 89 (1974)
- 4. Burstein, M., et al., Can. J. Biochem., 55, 766 (1977)
- 5. Frederickson, D.S., Circulation, 31, 321 (1965)

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