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Hyperlipidemia due to hepatic triacylglycerol lipase deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base.

Hyperlipidemia due to hepatic triacylglycerol lipase deficiency. ORPHA:140905

Hyperlipidemia due to hepatic triacylglycerol lipase deficiency is a rare, genetic hyperalphalipoproteinemia disorder characterized by elevated plasma cholesterol and triglyceride (TG) levels with a marked TG enrichment of low- and high-density lipoproteins (HDL), presence of circulating beta-very low density lipoproteins and elevated HDL cholesterol levels, in the presence of a very low, or undetectable, postheparin plasma hepatic lipase activity. Premature atherosclerosis and/or coronary heart disease may be associated.