LCAT deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. LCAT deficiency. ORPHA:650

LCAT (lecithin-cholesterol acyltransferase) deficiency is a rare lipoprotein metabolism disorder characterized clinically by corneal opacities, and sometimes renal failure and hemolytic anemia, and biochemically by severely reduced HDL cholesterol.