Lecithin Acyltransferase Deficiency

National Cancer Institute

Source

National Cancer Institute. Lecithin Acyltransferase Deficiency. NCI Thesaurus. Code C84813.

A disorder of lipoprotein metabolism caused by mutations in the LCAT gene. It is characterized by deficiency of the enzyme lecithin cholesterol acyltransferase. It is manifested with corneal opacity, hemolytic anemia, and proteinuria.