Delta-beta-thalassemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Delta-beta-thalassemia. ORPHA:231237

Delta-beta-thalassemia is a form of beta-thalassemia (see this term) characterized by decreased or absent synthesis of the delta- and beta-globin chains with a compensatory increase in expression of fetal gamma-chain synthesis.