Hemoglobin C-beta-thalassemia syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemoglobin C-beta-thalassemia syndrome. ORPHA:231242

Hemoglobin C - beta-thalassemia (HbC - BT) is a form of beta-thalassemia (see this term) resulting in moderate hemolytic anemia.