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## 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

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

Qeios ID: 44NZOH · https://doi.org/10.32388/44NZOH