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Beta Globin Gene and Related Diseases: A Review

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ABSTRACT In the present paper an attempt has been made to review the variations Clinical or phenotypic diversity of b-thalassaemia and other b-haemoglobinopathies suggest that it is determined by layer upon layer of complexity. A wide variety of primary mutations at the b-globin gene; two well-defined secondary modifying loci (d and a gene) and several less well characterized tertiary modifiers interact with strong environmental component.

Home Back