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Look at the clinical and molecular spectrum of Wiedemann-Steiner syndrome: An emerging member of the chromatinopathies family

Abstract

Wiedemann-Steiner syndrome (OMIM #605130) is a rare cogenital malformation syndrome characterized by hypertrichosis cubiti associated with short stature; consistent facial features, including long eyelashes, thick or arched eyebrows with a lateral flare, wide nasal bridge, and downslanting and vertically narrow palpebral fissures; mild to

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Cited by: 1 **Author:** Laura Avagliano, Ilaria Parenti, Ilaria Par…

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Author: Gabriella Maria Squeo, Bartolomeo A... **Publish Year:** 2020

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A second type of the disorder (Marfan **syndrome** type 2; OMIM 154705) is associated with a second locus, MFS2, at 3p25-p24.2 in a large French **family** (**family** MS1).

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