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# Tietz syndrome

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Tietz syndrome](#). ORPHA:42665

Tietz syndrome is a genetic hypopigmentation and deafness syndrome characterized by congenital profound bilateral sensorineural hearing loss and generalized albino-like hypopigmentation of skin, eyes and hair.