# Catel-Manzke syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Catel-Manzke syndrome. ORPHA:1388*

Catel-Manzke syndrome is a rare bone disease characterized by bilateral hyperphalangy and clinodactyly of the index finger typically in association with Pierre Robin sequence (see this term) comprising micrognathia, cleft palate and glossoptosis.