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X-linked dominant chondrodysplasia, Chassaing-Lacombe type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked dominant chondrodysplasia, Chassaing-Lacombe type. ORPHA:163966

X-linked dominant chondrodysplasia Chassaing-Lacombe type is a rare genetic bone disorder characterized by chondrodysplasia, intrauterine growth retardation (IUGR), hydrocephaly and facial dysmorphism in the affected males.