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Maffucci syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Maffucci syndrome. ORPHA:163634

Maffucci syndrome is a very rare genetic bone and skin disorder characterized by multiple enchondromas, leading to bone deformities, combined with multiple dark, irregularly shaped hemangiomas or less commonly lymphangiomas.