Infantile myofibromatosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Infantile myofibromatosis. ORPHA:2591

Infantile myofibromatosis (IM) is a rare benign soft tissue tumor characterized by the development of nodules in the skin, striated muscles, bones, and in exceptional cases, visceral organs, leading to a broad spectrum of clinical symptoms. IM contains myofibroblasts.