Camurati-Engelmann disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Camurati-Engelmann disease. ORPHA:1328

Camurati-Englemann disease (CED) is a rare, clinically variable bone dysplasia syndrome characterized by hyperostosis of the long bones, skull, spine and pelvis, associated with severe pain in the extremities, a wide-based waddling gait, joint contractures, muscle weakness and easy fatigability. Camurati-Englemann disease (CED) is a rare, clinically variable bone dysplasia syndrome characterized by hyperostosis of the long bones, skull, spine and pelvis, associated with severe pain in the extremities, a wide-based waddling gait, joint contractures, muscle weakness and easy fatigability.