Stüve-Wiedemann syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Stüve-Wiedemann syndrome. ORPHA:3206

Stüve-Wiedemann syndrome (SWS) is a rare autosomal recessive congenital primary skeletal dysplasia, characterized by small stature, bowing of the long bones, camptodactyly, hyperthermic episodes, respiratory distress/apneic episodes and feeding difficulties that usually lead to early mortality.