Ogden syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Ogden syndrome. ORPHA:276432

Ogden syndrome is a rare, genetic progeroid syndrome characterized by a variable phenotype including postnatal growth delay, severe global developmental delay, hypotonia, non-specific dysmorphic facies with aged appearance and cryptorchidism, as well as cardiac arrthymias and skeletal anomalies. Patients typically present with widely opened fontanels, mainly truncal hypotonia, a waddling gait with hypertonia of the extremities, small hands and feet, broad great toes, scoliosis and redundant skin with lack of subcutaneous fat.