Opis choroby *

Definicja

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.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Premature aging appearance-developmental delay-cardiac arrhythmia syndrome
Zespół Ogdena

Kod ORPHA
276432

Kod OMIM
300855

Kod ICD10
E34.8

Kod ICD11
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*Źródło

orphanet