Goodman syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Goodman syndrome. ORPHA:65798

Goodman syndrome is an extremely rare genetic disorder characterized by marked malformations of the head and face (essentially acrocephaly), abnormalities of the hands and feet (polydactyly, syndactyly, clinodactyly, camptodactyly, ulnar deviation), and congenital heart disease. There have been no further descriptions in the literature since 1979. Goodman syndrome could be a variant of Carpenter syndrome.