Okamoto syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Okamoto syndrome. ORPHA:2729

Okamoto syndrome is characterised by congenital hydronephrosis, intellectual deficit, growth retardation, cleft palate, generalised hypotonia and a characteristic face. Cardiac anomalies have also been reported. To date, 6 cases have been reported.