Opitz G/BBB syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Opitz G/BBB syndrome. ORPHA:2745

Opitz G/BBB syndrome (OS) is a multiple congenital anomalies disorder characterized by malformations of the midline including hypertelorism, laryngo-tracheo-esophalgeal defects and hypospadias. There are two clinically indistinguishable genetic subtypes of Opitz G/BBB: X-linked Opitz G/BBB syndrome (XLOS), and autosomal dominant Opitz G/BBB syndrome (ADOS) (see these terms).