Spina bifida-hypospadias syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spina bifida-hypospadias syndrome. ORPHA:3176

Spina bifida-hypospadias syndrome is a rare developmental defect during embryogenesis disorder characterized by the specific association of glandular hypospadias and lumbo-sacral spina bifida. Affected individuals may or may not present additional congenital anomalies, such as hydrocephaly, microstomia, patent ductus arteriosus, cryptorchidism, intestinal malrotation, rocker-bottom feet, and hypertrichosis.