Simpson-Golabi-Behmel syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Simpson-Golabi-Behmel syndrome. ORPHA:373

Simpson-Golabi-Behmel syndrome (SGBS, also referred to as SGBS type 1) is a rare X-linked multiple congenital anomalies syndrome, characterized by pre- and postnatal overgrowth, distinctive craniofacial features, variable congenital malformations, organomegaly and an increased tumor risk.