Shprintzen-Goldberg syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Shprintzen-Goldberg syndrome. ORPHA:2462

Shprintzen-Goldberg syndrome (SGS) is a very rare genetic disorder characterized by craniosynostosis, craniofacial and skeletal abnormalities, marfanoid habitus, cardiac anomalies, neurological abnormalities, and intellectual disability.