Genitopalatocardiac syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Genitopalatocardiac syndrome. ORPHA:2075

Genitopalatocardiac syndrome is a rare, multiple congenital anomalies/dysmorphic syndrome characterized by male, 46,XY gonadal dysgenesis, cleft palate, micrognathia, conotruncal heart defects and unspecific skeletal, brain and kidney anomalies.