Schilbach-Rott syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Schilbach-Rott syndrome. ORPHA:2353

Schilbach-Rott syndrome (SRS) is an autosomal dominant dysmorphic disorder that is characterized by dysmorphic facies with hypotelorism, blepharophimosis, and cleft palate, and the frequent occurrence of hypospadias in males.