Autosomal dominant Robinow syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant Robinow syndrome. ORPHA:3107

Autosomal dominant Robinow syndrome (DRS) is the more common type of Robinow syndrome (RS, see this term) characterized by mild to moderate limb shortening and abnormalities of the head, face and external genitalia.