Autosomal recessive Robinow syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive Robinow syndrome. ORPHA:1507

Autosomal recessive Robinow syndrome (RRS) is the less common type of Robinow syndrome (RS, see this term) characterized by short-limb dwarfism, costovertebral segmentation defects and abnormalities of the head, face and external genitalia.