Fryns syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fryns syndrome. ORPHA:2059

Fryns syndrome (FS) is a multiple congenital anomaly syndrome characterized by dysmorphic facial features, congenital diaphragmatic hernia, pulmonary hypoplasia, and distal limb hypoplasia, in addition to variable expression of additional malformations.