Levocardia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Levocardia. ORPHA:95854

A rare, congenital, non-syndromic, developmental defect during embryogenesis characterized by the heart located in the normal (levo) position associated with abdominal viscera located in the dextro position. Cardiac (e.g. interrupted inferior vena cava with azygous continuation) and/or splenic (asplenia, polysplenia) anomalies, as well as intestinal malrotation, are frequently associated.