Congenital rubella syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital rubella syndrome. ORPHA:290

Congenital rubella syndrome (CRS) is an infectious embryofetopathy that may present in an infant as a result of maternal infection and subsequent fetal infection with rubella virus. CRS can lead to deafness, cataract, and variety of other permanent manifestations including cardiac and neurological defects.