PARC syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. PARC syndrome. ORPHA:2825

PARC syndrome is a rare genetic developmental defect during embryogenesis syndrome characterized by the association of congenital poikiloderma (P), generalized alopecia (A), retrognathism (R) and cleft palate (C). There have been no further descriptions in the literature since 1990.