Primary ciliary dyskinesia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Primary ciliary dyskinesia. ORPHA:244

Primary ciliary dyskinesia (PCD) is a rare, genetically heterogeneous, primarily respiratory disorder characterized by chronic upper and lower respiratory tract disease. Approximately half of PCD patients have an organ laterality defect (situs inversus totalis or situs ambiguus/heterotaxy; see these terms).