Athyreosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Athyreosis. ORPHA:95713

Athyreosis is a form of thyroid dysgenesis (see this term) characterized by complete absence of thyroid tissue that results in primary congenital hypothyroidism (see this term), a permanent thyroid deficiency that is present from birth.