Primary cutaneous follicle center lymphoma

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Primary cutaneous follicle center lymphoma. ORPHA:178540

A rare, indolent primary cutaneous B-cell lymphoma characterized by a solitary or grouped erythematous plaques or tumors, preferentially located on the head, neck or trunk region, and composed of centroblasts and centrocytes arranged in a follicular, diffuse, or mixed growth pattern. The lesions are smooth and typically do not ulcerate. The neoplastic cells express pan B cell markers and Bcl-6, and typically lack Bcl-2.