SCALP syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. SCALP syndrome. ORPHA:370052

SCALP syndrome is a rare skin disease characterized by the association of sebaceous nevus and aplasia cutis congenita (usually on the scalp and face) in conjunction with limbal dermoid of the eye, a giant congenital melanocytic nevus and variable central nervous system abnormalities, including seizures, hydrocephalus, neurocutaneous melanosis, arachnoid cysts, and diffuse unilateral hemishpere enlargement.