Cronkhite-Canada syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Cronkhite-Canada syndrome. ORPHA:2930

Cronkhite-Canada syndrome (CCS) is a rare gastrointestinal (GI) polyposis syndrome characterized by the association of non-hereditary GI polyposis with the cutaneous triad of alopecia, nail changes and hyperpigmentation.