Open Peer Review on Qeios

Keratosis follicularis spinulosa decalvans

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Keratosis follicularis spinulosa decalvans. ORPHA:2340

Keratosis follicularis spinulosa decalvans is a rare genodermatosis occurring during infancy or childhood, predominantly affecting males, and characterized by diffuse follicular hyperkeratosis associated with progressive cicatricial alopecia of the scalp, eyebrows and eyelashes. Additional findings can include photophobia, corneal dystrophy, facial erythema, and/or palmoplantar keratoderma.