DISEASE:
Keratosis follicularis spinulosa decalvans

NAME:	Keratosis follicularis spinulosa decalvans
DESCRIPTION:	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.
ORPHACODE:	2340
XREF(S):	Orphanet OMIM OMIM MeSH ICD-10 OMIM
ANALYTE(S):	MBTPS2 SAT1 LRP1
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