DISEASE:
Autosomal dominant palmoplantar keratoderma and congenital alopecia

NAME:	Autosomal dominant palmoplantar keratoderma and congenital alopecia
DESCRIPTION:	A rare genetic skin disorder characterized by absence of scalp and body hair and palmoplantar keratoderma, without other hand complications.
ORPHACODE:	1010
SYNONYMS:	Autosomal dominant palmoplantar hyperkeratosis and congenital alopecia PPK-CA, Stevanovic type Palmoplantar keratoderma and congenital alopecia, Stevanovic type
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	GJA1
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