DISEASE:
Keratosis linearis-ichthyosis congenita-sclerosing keratoderma syndrome

NAME:	Keratosis linearis-ichthyosis congenita-sclerosing keratoderma syndrome
DESCRIPTION:	Keratosis linearis-ichthyosis congenita-sclerosing keratoderma syndrome is an inherited epidermal disorder characterized by palmoplantar keratoderma, linear hyperkeratotic papules on the flexural side of large joints (cord-like distribution around wrists, in antecubital and popliteal folds), hyperkeratotic plaques (on neck, axillae, elbows, wrists, and knees), mild ichthyosiform scaling, and sclerotic constrictions around fingers that present flexural deformities.
ORPHACODE:	281201
SYNONYMS:	KLICK syndrome
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	POMP
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