DISEASE:
Autosomal recessive epidermolytic ichthyosis

NAME:	Autosomal recessive epidermolytic ichthyosis
DESCRIPTION:	A rare, inherited, non-syndromic ichthyosis characterized by congenital, generalized erythroderma with cutaneous blistering and erosions, resembling collodion presentation at birth, replaced by progressive hyperkeratosis later in life without palmoplantar involvement. The ultrastructural pathology consists of sparse keratin filaments and keratin clumps that show a nearly homogeneous, amorphous structure.
ORPHACODE:	512103
SYNONYMS:	AREI
XREF(S):	Orphanet ICD-10
ANALYTE(S):	KRT10
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