DISEASE:
KID syndrome

NAME:	KID syndrome
DESCRIPTION:	A rare congenital ectodermal disorder characterized by vascularizing keratitis, hyperkeratotic skin lesions and hearing loss.
ORPHACODE:	477
SYNONYMS:	Ichthyosis hystrix Rheydt type KID/HID syndrome Keratitis-ichthyosis-deafness/Hystrix-like ichthyosis-deafness syndrome Keratitis-ichthyosis-hearing loss/Hystrix-like ichthyosis-hearing loss syndrome Senter syndrome
XREF(S):	Orphanet ICD-10 MedDRA OMIM OMIM OMIM
ANALYTE(S):	GJB2 GJB6
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