DISEASE:
Syndromic recessive X-linked ichthyosis

NAME:	Syndromic recessive X-linked ichthyosis
DESCRIPTION:	A rare genetic skin disease belonging to the Mendelian Disorders of Cornification (MeDOC) characterized by a generally mild cutaneous desquamation in association with extracutaneous manifestations as part of a syndrome.
ORPHACODE:	281090
SYNONYMS:	Recessive X-linked ichthyosis with extracutaneous manifestations Syndromic RXLI
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	STS
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