DISEASE:
Ichthyosis-prematurity syndrome

NAME:	Ichthyosis-prematurity syndrome
DESCRIPTION:	A rare, syndromic congenital ichthyosis characterized by premature birth (at gestational weeks 30-32, in general) in addition to thick, caseous and desquamating epidermis, neonatal respiratory asphyxia, and persistent eosinophilia. After the perinatal period, a spontaneous improvement in the health of affected patients is observed and skin features (vernix caseosa-like scale) evolve into a mild presentation of flat follicular hyperkeratosis with atopy.
ORPHACODE:	88621
SYNONYMS:	Congenital ichthyosis type 4 IPS
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	SLC27A4
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