DISEASE:
Zimmermann-Laband syndrome

NAME:	Zimmermann-Laband syndrome
DESCRIPTION:	A rare genetic multiple congenital anomalies syndrome characterized by gingival fibromatosis, coarse facial appearance, and absence or hypoplasia of nails or terminal phalanges of hands and feet.
ORPHACODE:	3473
SYNONYMS:	Gingival fibromatosis-hepatosplenomegaly-other anomalies syndrome Laband syndrome
XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM OMIM
ANALYTE(S):	KCNN3 ATP6V1B2 KCNH1 KCNMA1
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RELATED CONTENT

Related Analytes

- ATPase H+ transporting V1 subunit B2
- potassium voltage-gated channel subfamily H member 1
- potassium calcium-activated channel subfamily M alpha 1
- potassium calcium-activated channel subfamily N member 3