DISEASE:
Waardenburg syndrome type 2

NAME:	Waardenburg syndrome type 2
DESCRIPTION:	An autosomal dominant subtype of Waardenburg syndrome (WS) characterized by varying degrees of deafness and pigmentation anomalies of eyes, hair and skin, but without dystopia canthorum.
ORPHACODE:	895
SYNONYMS:	WS2 Waardenburg syndrome type II
XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM OMIM MeSH ICD-10

ANALYTE(S):	KITLG TYR SNAI2 SOX10 MITF EDNRB
CREATED:	13 May 2019 - 01:02
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- Centrum Medische Genetica - UZ Antwerpen

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- KIT ligand
- melanocyte inducing transcription factor
- snail family transcriptional repressor 2
- SRY-box transcription factor 10
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