DISEASE:
Waardenburg syndrome type 3

NAME:	Waardenburg syndrome type 3
DESCRIPTION:	A very rare subtype of Waardenburg syndrome (WS) that is characterized by limb anomalies in association with congenital hearing loss, minor defects in structures arising from neural crest, resulting in pigmentation anomalies of eyes, hair, and skin.
ORPHACODE:	896
SYNONYMS:	Klein-Waardenburg syndrome WS3 Waardenburg syndrome type III Waardenburg syndrome with limb anomalies
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PAX3
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RELATED CONTENT

Related Genetic Tests

- Waardenburg Syndrome types I and III
- Waardenburg syndrome (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- paired box 3

Related Gene Panels

- Waardenburg syndrome (6 genes) - UZA