DISEASE:
Waardenburg syndrome type 1

NAME:	Waardenburg syndrome type 1
DESCRIPTION:	A subtype of Waardenburg syndrome (WS) characterized by congenital deafness, minor defects in structures arising from neural crest resulting in pigmentation anomalies of eyes, hair, and skin, in combination with dystopia canthorum.
ORPHACODE:	894
SYNONYMS:	WS1 Waardenburg syndrome type I
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PAX3
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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