DISEASE:
Oculocutaneous albinism type 4

NAME:	Oculocutaneous albinism type 4
DESCRIPTION:	A form of oculocutaneous albinism characterized by varying degrees of skin and hair hypopigmentation, numerous ocular changes and misrouting of the optic nerves at the chiasm.
ORPHACODE:	79435
SYNONYMS:	OCA4
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SLC45A2
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Ocular albinism and oculocutaneous albinism type 1, 2, 3, 4, 6, 7, 8 (gene panel)

Related Laboratories

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- solute carrier family 45 member 2

Related Gene Panels

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