DISEASE:
Oculocutaneous albinism type 7

NAME:	Oculocutaneous albinism type 7
DESCRIPTION:	A form of oculocutaneous albinism (OCA) characterized by skin and hair hypopigmentation (light blond to dark brown), nystagmus, iris transillumination, visual acuity ranging from 6/9 to 3/60 and hypopigmentation of the peripheral ocular fundus. Photophobia is not a major feature.
ORPHACODE:	352745
SYNONYMS:	OCA7
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	LRMDA
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

- Centrum Medische Genetica - UZ Gent

Related Analytes

- leucine rich melanocyte differentiation associated

Related Gene Panels

- Ocular and oculocutaneous albinism - UGent