

DISEASE:

Lattice corneal dystrophy type I

NAME:	Lattice corneal dystrophy type I
DESCRIPTION:	Type I lattice corneal dystrophy (LCDI) is a frequent form of stromal corneal dystrophy (see this term) characterized by a network of delicate interdigitating branching filamentous opacities within the cornea with progressive visual impairment and no systemic manifestations.
ORPHACODE:	98964
SYNONYMS:	Biber-Haab-Dimmer dystrophy Classic lattice corneal dystrophy LCD1 LCDI Lattice corneal dystrophy type 1
XREF(S):	Orphanet ICD-10 OMIM OMIM MeSH
ANALYTE(S):	TGFB1
CREATED:	13 May 2019 - 01:02

CHANGED:	22 Jun 2023 - 16:14
-----------------	---------------------

Source URL: <http://gentest.healthdata.be/disease/3614>

RELATED CONTENT

Related Genetic Tests

- [Corneal dystrophy \(gene panel\)](#)

Related Laboratories

- [Centrum Medische Genetica - UZ Gent](#)

Related Analytes

- [transforming growth factor beta induced](#)

Related Gene Panels

- [Corneal dystrophy - UGent](#)

Source URL: <http://gentest.healthdata.be/disease/3614>