DISEASE:
3-methylglutaconic aciduria type 3

NAME:	3-methylglutaconic aciduria type 3
DESCRIPTION:	3-methylglutaconic aciduria type III (MGA III) is an organic aciduria characterised by the association of optic atrophy and choreoathetosis with 3-methylglutaconic aciduria.
ORPHACODE:	67047
SYNONYMS:	Autosomal recessive optic atrophy plus syndrome Autosomal recessive optic atrophy type 3 Costeff optic atrophy syndrome Costeff syndrome Infantile optic atrophy with chorea and spastic paraplegia MGA3
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	MICOS13 OPA3
CREATED:	13 May 2019 - 01:02

CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/2792

RELATED CONTENT

Related Genetic Tests

- Cataract (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- mitochondrial contact site and cristae organizing system subunit 13
- outer mitochondrial membrane lipid metabolism regulator OPA3

Related Gene Panels

- Cataract - UGent
- test