Belgian Genetic Tests database

ANALYTE:
EHHADH

NAME:	enoyl-CoA hydratase and 3-hydroxyacyl CoA dehydrogenase
SYMBOL:	EHHADH
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/2268

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- End-stage renal disease, ESRD (gene panel)
- Hepatology (gene panel)
- Hepatorenal disorders (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Metabolic diseases with hepatic disorders (20 genes)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Bifunctional enzyme deficiency
- Primary Fanconi renotubular syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Hepatology panel - UGent
- Hepatorenal disorders (13 genes) - UCL
- Metabolic diseases with hepatic disorders (20 genes) - UCL
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent

- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
