

ANALYTE:
HNF4A

NAME:	hepatocyte nuclear factor 4 alpha
SYMBOL:	HNF4A
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SYNONYMS:	HNF4 NR2A1
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
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RELATED CONTENT

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- [Intellectual disability \(virtual gene panel\)](#)
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- [Nephropathies, hereditary \(gene panel\)](#)
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Related Diseases

- [Atypical Fanconi syndrome-neonatal hyperinsulinism syndrome](#)
- [Congenital hyperinsulinism due to HNF4A deficiency](#)
- [HNF1B-related autosomal dominant tubulointerstitial kidney disease](#)
- [MODY](#)

Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)

- Congenital structural heart defects - UGent
- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (genepanel) - UZA
- Hyperinsulinism (5 genes) - UZA
- Intellectual disability (gene panel)
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- MODY (7 genes) - UZA
- MODY - Maturity onset Diabete of the Young (21 genes) - IPG
- Nephrocalcinosis and nephrolithiasis (37 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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