DISEASE:
Autosomal recessive non-syndromic intellectual disability

NAME:	Autosomal recessive non-syndromic intellectual disability
ORPHACODE:	88616
SYNONYMS:	AR-NSID NS-ARID

XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM OMIM OMIM ICD-10 ICD-10 ICD-10 ICD-10 OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM

ANALYTE(S):	DCPS B3GALNT2 IMPA1 ABCA2 CHKA GRIA1 YIF1B AIMP1 UFSP2 CRBN TUSC3 TECR GRIK2 WASHC4 TRAPPC9 PRSS12 MED23 ST3GAL3 CRADD ZC3H14 NSUN2 LINS1 PGAP1 METTL23 CLIP1 FBXO31 NDST1 FMN2 EDC3 HNMT EZR LMAN2L NEMF GRM7 IQSEC1 TNIK NCDN UBE4A TTC5

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RELATED CONTENT

Related Genetic Tests

- Congenital disorders of glycosylation (79 genes)
- Epilepsy (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ATP binding cassette subfamily A member 2
- aminoacyl tRNA synthetase complex interacting multifunctional protein 1
- alkB homolog 8, tRNA methyltransferase
- beta-1,3-N-acetylgalactosaminyltransferase 2
- chromosome 12 open reading frame 4
- coiled-coil and C2 domain containing 1A
- choline kinase alpha
- CAP-Gly domain containing linker protein 1
- CASP2 and RIPK1 domain containing adaptor with death domain
- cereblon
- decapping enzyme, scavenger
- enhancer of mRNA decapping 3
- eukaryotic translation elongation factor 1 beta 2

- ezrin
- F-box protein 31
- formin 2
- ferric chelate reductase 1 like
- gem nuclear organelle associated protein 5
- glutamate ionotropic receptor AMPA type subunit 1
- glutamate ionotropic receptor kainate type subunit 2
- glutamate metabotropic receptor 7
- histamine N-methyltransferase
- inositol monophosphatase 1
- IQ motif and Sec7 domain ArfGEF 1
- lysine demethylase 5B
- lines homolog 1
- lectin, mannose binding 2 like
- mannosidase alpha class 1B member 1
- membrane bound O-acyltransferase domain containing 7
- mediator complex subunit 23
- mediator complex subunit 25
- methyltransferase 23, arginine
- N-alpha-acetyltransferase 20, NatB catalytic subunit
- neurochondrin
- N-deacetylase and N-sulfotransferase 1
- nuclear export mediator factor
- NOP2/Sun RNA methyltransferase 2
- post-GPI attachment to proteins inositol deacylase 1
- phosphatidylinositol glycan anchor biosynthesis class C
- serine protease 12
- arginine and serine rich coiled-coil 1
- seryl-tRNA synthetase 1
- solute carrier family 45 member 1
- ST3 beta-galactoside alpha-2,3-sialyltransferase 3
- trans-2,3-enoyl-CoA reductase
- TRAF2 and NCK interacting kinase

- translocated promoter region, nuclear basket protein
- trafficking protein particle complex subunit 9
- tetratricopeptide repeat domain 5
- tumor suppressor candidate 3
- ubiquitination factor E4A
- UFM1 specific peptidase 2
- WASH complex subunit 4
- Yip1 interacting factor homolog B, membrane trafficking protein
- zinc finger CCCH-type containing 14

Related Gene Panels

- Congenital disorders of glycosylation (79 genes) - KUL
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Rare epilepsy with developmental delay (> 240 genes) - UZA
