DISEASE:
X-linked non-syndromic intellectual disability

NAME:	X-linked non-syndromic intellectual disability
ORPHACODE:	777

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

ANALYTE(S):	STEEP1 RPS6KA3 ARX DMD FTSJ1 GDI1 MED12 UPF3B AGTR2 ZNF41 DLG3 IL1RAPL1 PAK3 TSPAN7 ARHGEF6 ZNF81 SYP ZNF711 RAB39B HCFC1 ALG13 MID2 PTCHD1 USP9X ACSL4 MECP2 IQSEC2 SLC9A7 CNKSR2 FRMPD4 USP27X CLCN4

CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Epilepsy (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- acyl-CoA synthetase long chain family member 4
- angiotensin II receptor type 2
- ALG13 UDP-N-acetylglucosaminyltransferase subunit
- Rac/Cdc42 guanine nucleotide exchange factor 6
- aristaless related homeobox
- chloride voltage-gated channel 4
- connector enhancer of kinase suppressor of Ras 2
- discs large MAGUK scaffold protein 3
- dystrophin
- FERM and PDZ domain containing 4
- FtsJ RNA 2'-O-methyltransferase 1
- GDP dissociation inhibitor 1
- host cell factor C1
- interleukin 1 receptor accessory protein like 1
- IQ motif and Sec7 domain ArfGEF 2

- methyl-CpG binding protein 2
- mediator complex subunit 12
- midline 2
- p21 (RAC1) activated kinase 3
- patched domain containing 1
- RAB39B, member RAS oncogene family
- ribosomal protein S6 kinase A3
- solute carrier family 9 member A7
- STING1 ER exit protein 1
- synaptophysin
- tetraspanin 7
- UPF3B regulator of nonsense mediated mRNA decay
- ubiquitin specific peptidase 27 X-linked
- ubiquitin specific peptidase 9 X-linked
- zinc finger protein 41
- zinc finger protein 711
- zinc finger protein 81

Related Gene Panels

- Rare epilepsy with developmental delay (> 240 genes) - UZA
