Belgian Genetic Tests database

ANALYTE:
NBN

NAME:	nibrin
SYMBOL:	NBN
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SYNONYMS:	AT-V1 AT-V2 ATV
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Breast and Ovarian cancer, HBOC, familial (gene panel - 26 genes)
- Breast cancer, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation (gene panel)
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - Primary Adrenal Insuffiency (gene panel)
- Familial cancer predisposition (gene panel)
- Hereditary cancer (Breast, ovary, colon) (26 genes)
- Hereditary cancer (gene panel)
- Hereditary cancer panel (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Malformations of cortical development (235 genes)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders (gene panel)
- Nijmegen Breakage Syndrome
- Nijmegen breakage syndrome
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Familial prostate cancer
- Hereditary breast and/or ovarian cancer syndrome
- Nijmegen breakage syndrome

Related Gene Panels

- Extended Breast Cancer Panel (26 gene) - VUB
- Breast/Ovarian cancer (26 genes) - ULB
- Cancer (Breast, ovary, colon,…) (26 genes) - ULG
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - Primary Adrenal Insufficiency - UGent
- Hematologic Familiar Forms - ULG
- Hereditary Cancer Solution (35 genes) - UCL
- Hereditary breast and ovarian cancer (26 genes) - CHULg
- Hereditary cancer predisposition - UGent
- Hereditary predisposition to cancer (47 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability (gene panel) - UZA
- Intellectual disability/Epilepsy (859 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies - UGent