ANALYTE:
CHD7

NAME:	chromodomain helicase DNA binding protein 7
SYMBOL:	CHD7
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ20357 FLJ20361 KIAA1416
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Autism (gene panel)
- CHARGE syndrome
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Hearing loss (deafness), (gene panel)
- Hypogonadotropic hypogonadism (33 genes)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Malformations of cortical development (235 genes)
- Microphthalmia / Anophthalmia / Coloboma-Anterior Segment Dysgenesis (MAC-ASD) (gene panel)
- Myopathy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- CHARGE syndrome
- Kallmann syndrome

- Normosmic congenital hypogonadotropic hypogonadism
- Omenn syndrome

Related Gene Panels

- Autism (57 genes) - IPG
- CHARGE (2 genes) - UZA
- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Heterotaxie PCD - UGent
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Hypogonadotropic hypogonadism (33 genes) - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Microphthalmia/Anophthalmia/Coloboma - Anterior Segment Dysgenesis - UGent
- Myopathy (genepanel) - UZA
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia - UGent
