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Full name:	Brain malformations (34 genes) - ULB
Laboratory:	Centre de Génétique Humaine - Erasme ULB
Created:	09 Dec 2019 - 12:32
Changed:	03 May 2023 - 12:29

Related Diseases

- Autosomal recessive primary microcephaly
- Ear-patella-short stature syndrome
- Microcephalic osteodysplastic primordial dwarfism type II
- Microform holoprosencephaly
- Primary microcephaly-mild intellectual disability-young-onset diabetes syndrome
- Seckel syndrome

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ASPM	98.88	0	
CCDC88C	100.00	0	
CDK5RAP2	99.64	0	
CDON	100.00	0	
CENPJ	100.00	0	
CEP135	100.00	0	
CEP152	99.96	0	
CRIPTO	100.00	0	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
DISP1	100.00	0	
DLL1	100.00	0	
FGF8	99.79	0	
FGFR1	94.19	0	
FOXH1	100.00	0	
GAS1	100.00	0	
GCM2	78.95	0	
GLI2	100.00	0	
L1CAM	96.28	0	
MCPH1	99.98	0	
MPDZ	99.70	0	
NODAL	100.00	0	
ORC4	100.00	0	
ORC6	100.00	0	
PCNT	100.00	0	
PTCH1	99.88	0	
SHH	98.82	0	
SIX3	99.47	0	
SLC25A15	99.87	0	
SMAD2	99.61	0	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
STIL	100.00	0	
SUFU	100.00	0	
TGIF1	100.00	0	
TRMT10A	100.00	0	
WDR62	100.00	0	
ZIC2	100.00	0	