

DOMINANT OR DE NOVO MONOGENIC DISEASES

GENE	DISEASES	
BRAF	Cardiofaciocutaneous syndrome 1, LEOPARD syndrome 3, Noonan syndrome	●
CBL	Noonan syndrome-like disorder with or without juvenile myelomonocytic leukemia	●
CHD7	CHARGE syndrome	●
COL1A1	Osteogenesis imperfecta	●
COL1A2	Osteogenesis imperfecta	●
COL3A1	Ehlers-Danlos syndrome	●
CREBBP	Rubinstein-Taybi syndrome	●
EFNB1	Craniofrontonasal dysplasia	●
EP300	Rubinstein-Taybi syndrome	●
FBN1	Marfan syndrome	●
FGFR1	Pfeiffer syndrome	●
FGFR2	Crouzon syndrome	●
FGFR3	Hypochondroplasia	●
FOXP1	Congenital Rett-like syndrome	●
G6PD	Glucose-6-phosphate dehydrogenase deficiency	●
HDAC8	Cornelia de Lange syndrome 5	●
HRAS	Costello syndrome, Noonan syndrome	●
JAG1	Alagille syndrome	●
KAT6B	Genitopatellar syndrome	●
KMT2D	Kabuki syndrome	●
KRAS	Noonan syndrome	●
MAP2K1	Cardiofaciocutaneous syndrome 3, Noonan syndrome	●
MAP2K2	Cardiofaciocutaneous syndrome 4, Noonan syndrome	●
MECP2	Rett syndrome	●
NIPBL	Cornelia de Lange syndrome 1	●

GENE	PATOLOGIE	
NOTCH2	Alagille syndrome 2	●
NRAS	Noonan syndrome	●
NSD1	Sotos syndrome 1	●
OTC	Ornithine transcarbamylase deficiency	●
PAFAH1B1	Lissencephaly 1	●
POLR1C	Treacher Collins syndrome	●
PTPN11	LEOPARD syndrome	●
RAD21	Cornelia de Lange syndrome 4	●
RAF1	Noonan syndrome	●
SCN1A	Dravet syndrome	●
SHH	Holoprosencephaly type 3	●
SHOC2	Noonan-like syndrome with loose anagen hair, Noonan syndrome	●
SMC1A	Cornelia de Lange syndrome 2	●
SMC3	Cornelia de Lange syndrome 3	●
SOS1	Noonan syndrome	●
STXBP1	Early infantile epileptic	●
TBX5	encephalopathy Holt-Oram	●
TCF4	syndrome Pitt-Hopkins syndrome	●
TCOF1	Treacher Collins syndrome	●
TSC1	Tuberous sclerosis 1 Tuberous	●
TSC2	sclerosis 2 Lissencephaly	●
TUBA1A		●
TUBB2B	Tubulinopathy with polymicrogyria/lissencephaly	●
TWIST1	Saethre-Chotzen syndrome	●
ZIC2	Nonsyndromic holoprosencephaly	●

● screening test **NEGATIVE = LOW RISK**

● screening test **POSITIVE = HIGH RISK**