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Full name:	Erythocyoses, polycythémies, thrombocytoses congénitales (gene panel) - ULG
Laboratory:	Centre de Génétique Humaine - CHU Sart-Tilman
Created:	21 Dec 2022 - 11:31
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Related Diseases

- Autosomal dominant secondary polycythemia
- Congenital amegakaryocytic thrombocytopenia
- Essential thrombocythemia
- Familial thrombocytosis
- Primary familial polycythemia
- Primary myelofibrosis
- Thrombocythemia with distal limb defects

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ANKRD26	0.00	0	
BPGM	0.00	0	
CSF1R	0.00	0	
CSF3R	0.00	0	
DDX41	0.00	0	
EGLN1	0.00	0	
ELANE	0.00	0	
EPAS1	0.00	0	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
EPO	0.00	0	
EPOR	0.00	0	
GATA2	0.00	0	
JAK1	0.00	0	
JAK2	0.00	0	
JAK3	0.00	0	
MPL	0.00	0	
PIEZO1	0.00	0	
SH2B3	0.00	0	
SLC30A10	0.00	0	
STAT3	0.00	0	
STAT5B	0.00	0	
THPO	0.00	0	
VHL	0.00	0	