

GENETIC TEST:
Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)

FULL NAME:	Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)
DESCRIPTION:	Analysed genes: ANKRD26;BPGM; CSF1R; CSF3R; DDX41; EGLN1; ELANE; EPAS1; EPO; EPOR; GATA2; JAK1; JAK2; JAK3; SH2B3=LNK; MPL; PIEZO1 ; SLC30A10 ;STAT3; STAT5B; THPO; VHL
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Prognostic diagnosis, Therapeutic Management
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565530-565541

ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23
TURNAROUND TIME (MAXIMUM):	30
CREATED:	20 Dec 2022 - 17:05
CHANGED:	24 Jan 2024 - 12:18

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Related Analytes

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- [erythropoietin](#)

- [erythropoietin receptor](#)
- [GATA binding protein 2](#)
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- [Janus kinase 2](#)
- [Janus kinase 3](#)
- [MPL proto-oncogene, thrombopoietin receptor](#)
- [piezo type mechanosensitive ion channel component 1](#)
- [SH2B adaptor protein 3](#)
- [solute carrier family 30 member 10](#)
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