

GENETIC TEST: Fanconi anemia (gene panel)

FULL NAME:	Fanconi anemia (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS) MLPA based techniques digitalMLPA based techniques
RIZIV CODE:	565552-565563
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	30 Jul 2019 - 09:58

CHANGED:	24 Jan 2024 - 15:22
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Related Laboratories

- [Centrum Medische Genetica - UZ Gent](#)

Related Analytes

- [BRCA1 DNA repair associated](#)
- [BRCA2 DNA repair associated](#)
- [BRCA1 interacting helicase 1](#)
- [ERCC excision repair 4, endonuclease catalytic subunit](#)
- [FA complementation group A](#)
- [FA complementation group B](#)
- [FA complementation group C](#)
- [FA complementation group D2](#)
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- [mitotic arrest deficient 2 like 2](#)

- partner and localizer of BRCA2
- RAD51 recombinase
- RAD51 paralog C
- ring finger and WD repeat domain 3
- SLX4 structure-specific endonuclease subunit
- ubiquitin conjugating enzyme E2 T
- X-ray repair cross complementing 2

Related Gene Panels

- Fanconi anemia - UGent

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