

GENETIC TEST:
Capillary malformation - arteriovenous malformation (2 genes)

FULL NAME:	Capillary malformation - arteriovenous malformation (2 genes)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21

TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	24 Jul 2019 - 11:57
CHANGED:	16 Feb 2024 - 16:22

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Related Analytes

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- [fms related receptor tyrosine kinase 4](#)

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- [gap junction protein gamma 2](#)
- [glomulin, FKBP associated protein](#)
- [hepatocyte growth factor](#)
- [HRas proto-oncogene, GTPase](#)
- [inhibitor of nuclear factor kappa B kinase regulatory subunit gamma](#)
- [integrin subunit alpha 9](#)
- [kinesin family member 11](#)
- [KRAS proto-oncogene, GTPase](#)
- [KRIT1 ankyrin repeat containing](#)
- [NRAS proto-oncogene, GTPase](#)
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- [RAS p21 protein activator 1](#)
- [SMAD family member 4](#)
- [SOS Ras/Rac guanine nucleotide exchange factor 1](#)
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