GENETIC TEST:
Capillary malformation – microcephaly

FULL NAME:	Capillary malformation – microcephaly
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, Amniotic fluid, Chorionic villi
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:	12 Dec 2019 - 13:57
CHANGED:	21 Dec 2023 - 09:30

RELATED CONTENT

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- Microcephaly-capillary malformation syndrome

Related Laboratories

- Centre de Génétique Médicale UCL

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