

GENETIC TEST: Carney syndrome

FULL NAME:	Carney syndrome
DESCRIPTION:	Part of the Onco-endocrine pathologies panel (Panel OE)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal 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:	565530-565541
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21

TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	25 Jul 2019 - 07:57
CHANGED:	24 Jan 2023 - 14:57

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