GENETIC TEST:
Pheochromocytoma - paraganglioma syndrome (gene panel)

FULL NAME:	Pheochromocytoma - paraganglioma syndrome (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:	MLPA based techniques Next Generation Sequencing (NGS)
RIZIV CODE:	565530-565541
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
EQA:	• Phaeochromocytoma and Paraganglioma Disorders
TURNAROUND TIME (MAXIMUM):	3 - 4 months

CREATED:	22 Jul 2019 - 14:27
CHANGED:	13 Dec 2022 - 11:54

Source URL: http://gentest.healthdata.be/genetic_test/344

RELATED CONTENT

Related Diseases

- Hereditary pheochromocytoma-paraganglioma

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- MYC associated factor X
- ret proto-oncogene
- succinate dehydrogenase complex flavoprotein subunit A
- succinate dehydrogenase complex assembly factor 2
- succinate dehydrogenase complex iron sulfur subunit B
- succinate dehydrogenase complex subunit C
- succinate dehydrogenase complex subunit D
- succinate-CoA ligase GDP-forming subunit beta
- transmembrane protein 127
- von Hippel-Lindau tumor suppressor

Related Gene Panels

- Pheochromocytoma - paraganglioma syndrome - UGent
