DISEASE:
Sporadic pheochromocytoma

NAME:	Sporadic pheochromocytoma
DESCRIPTION:	A rare tumor of endocrine glands characterized, typically, by a unicentric, unilateral, sporadic, catecholamine-secreting neuroendocrine tumor, arising from chromaffin cells of the adrenal medulla, presenting with highly variable and non-specific manifestations, including hypertension (that can be paroxysmal, persistent, or resistant), cephalea, heart palpitations, anxiety, diaphoresis, unexplained fever, chronic fatigue and weakness, among others.
ORPHACODE:	276624
XREF(S):	Orphanet ICD-10 ICD-10
ANALYTE(S):	EPAS1 VHL RET SDHD SDHB
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RELATED CONTENT

Related Genetic Tests

- Genetic disorders of Calcium and Phosphate metabolism (gene panel)
- Onco-endocrine pathologies (gene panel)
- Paraganglioma and pheochromocytoma (gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- endothelial PAS domain protein 1
- ret proto-oncogene
- succinate dehydrogenase complex iron sulfur subunit B
- succinate dehydrogenase complex subunit D
- von Hippel-Lindau tumor suppressor

Related Gene Panels

- Genetic disorders of Calcium and Phosphate metabolism (31 genes) - KUL
- Onco-endocine pathologies (50 genes) - UCL
- Paraganglioma and pheochromocytoma (29 genes) - UCL
