GENETIC TEST:
Kallmann syndrome / Hypogonadotropic Hypogonadism (FGFR1 gene)

FULL NAME:	Kallmann syndrome / Hypogonadotropic Hypogonadism (FGFR1 gene)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	02 Sep 2019 - 07:26
CHANGED:	01 Dec 2023 - 17:07

URL:	https://www.chu.ulg.ac.be/jcms/c2_23451207/fr/hypogondisme-hypogonadotrope-etud…

RELATED CONTENT

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