GENETIC TEST:
Phenylketonuria

FULL NAME:	Phenylketonuria
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) MLPA based techniques
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-03-25 / 2024-09-09

EQA:	• Phenylketonuria, • Phenylketonuria
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	23 Aug 2019 - 07:20
CHANGED:	18 Aug 2023 - 09:31
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM65

RELATED CONTENT

Related Diseases

- Classic phenylketonuria

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- phenylalanine hydroxylase