GENETIC TEST:
Non Invasive Prenatal Testing (NIPT) of trisomies 13, 18 et 21 and sex chromosomes

FULL NAME:	Non Invasive Prenatal Testing (NIPT) of trisomies 13, 18 et 21 and sex chromosomes
TEST TYPE:	Clinical
TEST SPECIALTY:	Cytogenetics
TEST PURPOSE:	Prenatal diagnosis
SPECIMEN:	Cell free DNA
METHOD CATEGORY:	Detection of chromosome alteration large in size
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565611-565622
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies, • NIPT for common aneuploidies
TURNAROUND TIME (MAXIMUM):	7 calendar days
CREATED:	24 Jul 2019 - 09:55
CHANGED:	01 Mar 2023 - 15:58
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/5464

RELATED CONTENT

Related Diseases

- Down syndrome
- Trisomy 13
- Trisomy 18

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- Chromosome 13
- Chromosome 18
- Chromosome 21