GENETIC TEST:
Beckwith-Wiedemann syndrome (11p15 methylation)

FULL NAME:	Beckwith-Wiedemann syndrome (11p15 methylation)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Methylation analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• Beckwith - Wiedermann & Silver-Russell syndromes, • Beckwith - Wiedermann & Silver-Russell syndromes, • Beckwith - Wiedermann & Silver-Russell syndromes
TURNAROUND TIME (MAXIMUM):	8- 13 weeks
CREATED:	18 Jul 2019 - 09:17
CHANGED:	04 Dec 2023 - 12:53
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB/12268

RELATED CONTENT

Related Diseases

- Beckwith-Wiedemann syndrome
- Beckwith-Wiedemann syndrome due to imprinting defect of 11p15

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- H19 imprinted maternally expressed transcript
- insulin like growth factor 2
- KCNQ1 opposite strand/antisense transcript 1