GENETIC TEST:
Angelman / Prader Willi Syndrome

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

EQA:	• Prader-Willi and Angelman Syndromes, • Prader-Willi and Angelman Syndromes , • Prader-Willi and Angelman Syndromes , • Prader-Willi and Angelman Syndromes, • Prader-Willi and Angelman Syndromes
TURNAROUND TIME (MAXIMUM):	8- 13 weeks
CREATED:	17 Jul 2019 - 13:32
CHANGED:	04 Dec 2023 - 12:33
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB/13673

RELATED CONTENT

Related Diseases

- Angelman syndrome due to imprinting defect in 15q11-q13
- Angelman syndrome due to maternal 15q11q13 deletion
- Angelman syndrome due to paternal uniparental disomy of chromosome 15
- Prader-Willi syndrome due to imprinting mutation
- Prader-Willi syndrome due to maternal uniparental disomy of chromosome 15
- Prader-Willi syndrome due to paternal deletion of 15q11q13 type 1

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- chromosome 15 - 15q11-q13
- ubiquitin protein ligase E3A