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):	2021-03-25 / 2024-09-09

EQA:	• Prader-Willi and Angelman Syndromes, • Prader-Willi and Angelman Syndromes
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	24 Jul 2019 - 11:08
CHANGED:	17 Aug 2023 - 10:39
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM78

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
- Prader-Willi syndrome due to paternal deletion of 15q11q13 type 2

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

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