

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
METHOD CATEGORY:	Methylation analysis Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21

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, • Prader-Willi and Angelman Syndromes
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	24 Jul 2019 - 11:08
CHANGED:	24 Jan 2023 - 14:36

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- [Angelman syndrome due to maternal 15q11q13 deletion](#)
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- [Prader-Willi syndrome due to imprinting mutation](#)
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- [Prader-Willi syndrome due to paternal deletion of 15q11q13 type 1](#)

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