

GENETIC TEST:

Spinal muscular atrophy (SMA) type 1 (Werdnig-Hoffmann), type 2, type 3 (Kugelberg-Welander) and type 4

FULL NAME:	Spinal muscular atrophy (SMA) type 1 (Werdnig-Hoffmann), type 2, type 3 (Kugelberg-Welander) and type 4
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2023-11-09 / 2024-05-08

EQA:	<ul style="list-style-type: none">• Spinal Muscular Atrophy,• Spinal Muscular Atrophy,• Spinal Muscular Atrophy
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	23 Jul 2019 - 12:08
CHANGED:	22 Jan 2024 - 14:25
URL:	https://labogidsmedgen.uza.be/analyses/spinale-spieratrofie

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