

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):	2021-09-11 / 2026-09-10

EQA:	• Spinal Muscular Atrophy, • Spinal Muscular Atrophy, • Spinal Muscular Atrophy
TURNAROUND TIME (MAXIMUM):	6 weeks
CREATED:	23 Jul 2019 - 12:08
CHANGED:	07 Dec 2022 - 10:27

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