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:	PCR based technique
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2022-10-07 / 2027-10-06

EQA:	• Spinal Muscular Atrophy, • Spinal Muscular Atrophy, • Spinal Muscular Atrophy, • Spinal Muscular Atrophy
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	23 Jul 2019 - 12:08
CHANGED:	22 Jan 2024 - 10:07

RELATED CONTENT

Related Diseases

- Proximal spinal muscular atrophy type 1
- Proximal spinal muscular atrophy type 2
- Proximal spinal muscular atrophy type 3
- Proximal spinal muscular atrophy type 4

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- survival of motor neuron 1, telomeric
- survival of motor neuron 2, centromeric