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, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-03-25 / 2024-09-09

EQA:	• Spinal Muscular Atrophy, • Spinal Muscular Atrophy, • Spinal Muscular Atrophy , • Spinal Muscular Atrophy
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	23 Jul 2019 - 12:08
CHANGED:	19 Jan 2024 - 09:33
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM05

Source URL: http://gentest.healthdata.be/genetic_test/632

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 Humaine - Erasme ULB

Related Analytes

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