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

EQA:	• Spinal Muscular Atrophy, • Spinal Muscular Atrophy, • Spinal Muscular Atrophy
TURNAROUND TIME (MAXIMUM):	6 - 8 weeks
CREATED:	23 Jul 2019 - 12:08
CHANGED:	07 Dec 2022 - 10:26
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/12968

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

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

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