

GENETIC TEST:
Spinocerebellar ataxia (types 1, 2, 3, 6, 7) - CAG repeat expansion

FULL NAME:	Spinocerebellar ataxia (types 1, 2, 3, 6, 7) - CAG repeat expansion
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
EQA:	• Spinocerebellar Ataxia's, • Spinocerebellar Ataxia's
TURNAROUND TIME (MAXIMUM):	6 - 8 weeks

CREATED:	23 Jul 2019 - 12:23
CHANGED:	01 Mar 2023 - 16:08
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/15310

Source URL: http://gentest.healthdata.be/genetic_test/156

RELATED CONTENT

Related Diseases

- [Machado-Joseph disease type 1](#)
- [Machado-Joseph disease type 2](#)
- [Machado-Joseph disease type 3](#)
- [Spinocerebellar ataxia type 1](#)
- [Spinocerebellar ataxia type 2](#)
- [Spinocerebellar ataxia type 6](#)
- [Spinocerebellar ataxia type 7](#)

Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

Related Analytes

- [ataxin 1](#)
- [ataxin 2](#)
- [ataxin 3](#)
- [ataxin 7](#)
- [calcium voltage-gated channel subunit alpha1 A](#)

Related Gene Panels

- Spinocerebellar ataxia (types 1, 2, 3, 6, 7) (5 genes) - KUL

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