

**GENETIC TEST:**  
**Spastic paraplegia-4**

<b>FULL NAME:</b>	Spastic paraplegia-4
<b>TEST TYPE:</b>	Clinical
<b>TEST SPECIALTY:</b>	Molecular Genetics
<b>TEST PURPOSE:</b>	Carrier diagnosis, Post-natal Diagnosis
<b>SPECIMEN:</b>	Peripheral (whole) blood on EDTA
<b>METHOD CATEGORY:</b>	Sequence analysis: entire coding region Deletion/duplication analysis
<b>METHOD TECHNIQUE:</b>	Bi-directional Sanger Sequence analysis MLPA based techniques
<b>RIZIV CODE:</b>	565471-565482
<b>TURNAROUND TIME (MAXIMUM):</b>	4 - 6 months
<b>CREATED:</b>	23 Jul 2019 - 11:58
<b>CHANGED:</b>	21 Oct 2021 - 10:23
<b>URL:</b>	<a href="https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/15335">https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/15335</a>

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## RELATED CONTENT

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### Related Diseases

- [Autosomal dominant spastic paraplegia type 4](#)

### Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [spastin](#)

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