

## GENETIC TEST: Peripheral neuropathy (gene panel)

<b>FULL NAME:</b>	Peripheral neuropathy (gene panel)
<b>TEST TYPE:</b>	Clinical
<b>TEST SPECIALTY:</b>	Molecular Genetics
<b>TEST PURPOSE:</b>	Post-natal Diagnosis
<b>SPECIMEN:</b>	Peripheral (whole) blood on EDTA, DNA
<b>METHOD CATEGORY:</b>	Sequence analysis: entire coding region Whole Exome Sequencing (WES)
<b>METHOD TECHNIQUE:</b>	Next Generation Sequencing (NGS)
<b>RIZIV CODE:</b>	565493-565504
<b>ACCREDITATION (ISO 15189):</b>	2023-11-09 / 2024-05-08
<b>TURNAROUND TIME (MAXIMUM):</b>	6 months
<b>CREATED:</b>	11 Dec 2020 - 09:59
<b>CHANGED:</b>	22 Jan 2024 - 14:20

URL:	<a href="https://labogidsmedgen.uza.be/analyses/whole-exome-sequencing-wes-ikv-perifere-...">https://labogidsmedgen.uza.be/analyses/whole-exome-sequencing-wes-ikv-perifere-...</a>
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## RELATED CONTENT

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### Related Diseases

- [Autosomal dominant Charcot-Marie-Tooth disease type 2D](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2DD](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2F](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2K](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2L](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2N](#)
- [Autosomal dominant Charcot-Marie-Tooth disease type 2W](#)
- [Autosomal dominant intermediate Charcot-Marie-Tooth disease type F](#)
- [Autosomal recessive Charcot-Marie-Tooth disease with hoarseness](#)
- [Autosomal recessive axonal neuropathy with neuromyotonia](#)
- [Autosomal recessive intermediate Charcot-Marie-Tooth disease type A](#)
- [Autosomal recessive intermediate Charcot-Marie-Tooth disease type D](#)
- [Charcot-Marie-Tooth disease type 1D](#)
- [Charcot-Marie-Tooth disease type 2H](#)
- [Charcot-Marie-Tooth disease type 4A](#)
- [Charcot-Marie-Tooth disease type 4E](#)
- [Charcot-Marie-Tooth disease type 4G](#)
- [Charcot-Marie-Tooth disease type 4H](#)
- [Charcot-Marie-Tooth disease type 4J](#)
- [Dejerine-Sottas syndrome](#)
- [Distal hereditary motor neuropathy type 2](#)
- [Distal hereditary motor neuropathy type 5](#)
- [Muscular dystrophy, Selcen type](#)
- [Neuropathy with hearing impairment](#)
- [Severe neurodegenerative syndrome with lipodystrophy](#)
- [X-linked Charcot-Marie-Tooth disease type 1](#)

- [X-linked Charcot-Marie-Tooth disease type 4](#)

## Related Laboratories

- [Centrum Medische Genetica - UZ Antwerpen](#)

## Related Analytes

- [alanyl-tRNA synthetase 1](#)
- [ATP binding cassette subfamily A member 1](#)
- [abhydrolase domain containing 12, lysophospholipase](#)
- [ADP-ribosylserine hydrolase](#)
- [ATP/GTP binding carboxypeptidase 1](#)
- [apoptosis inducing factor mitochondria associated 1](#)
- [Rho guanine nucleotide exchange factor 10](#)
- [arylsulfatase A](#)
- [atlastin GTPase 1](#)
- [atlastin GTPase 3](#)
- [ATM serine/threonine kinase](#)
- [ATPase Na+/K+ transporting subunit alpha 1](#)
- [ATPase copper transporting alpha](#)
- [beta-1,4-N-acetyl-galactosaminyltransferase 1](#)
- [BAG cochaperone 3](#)
- [BICD cargo adaptor 2](#)
- [BSCL2 lipid droplet biogenesis associated, seipin](#)
- [cell adhesion molecule 3](#)
- [CD59 molecule \(CD59 blood group\)](#)
- [cilia and flagella associated protein 276](#)
- [coiled-coil-helix-coiled-coil-helix domain containing 10](#)
- [clathrin heavy chain like 1](#)

- contactin associated protein 1
- cytochrome c oxidase assembly factor 7
- cytochrome c oxidase assembly factor COX20
- cytochrome c oxidase subunit 6A1
- CTD phosphatase subunit 1
- catenin beta 1
- cytochrome P450 family 27 subfamily A member 1
- aspartyl-tRNA synthetase 2, mitochondrial
- DDB1 and CUL4 associated factor 8
- dynactin subunit 1
- diacylglycerol O-acyltransferase 2
- deoxyguanosine kinase
- dehydrogenase E1 and transketolase domain containing 1
- DnaJ heat shock protein family (Hsp40) member B2
- DnaJ heat shock protein family (Hsp40) member C3
- dynamin 2
- DNA methyltransferase 1
- dystrophin related protein 2
- dystonin
- dynein cytoplasmic 1 heavy chain 1
- early growth response 2
- elongator complex protein 1
- electron transfer flavoprotein dehydrogenase
- fibulin 5
- F-box protein 38
- FYVE, RhoGEF and PH domain containing 4
- FIG4 phosphoinositide 5-phosphatase
- FLVCR heme transporter 1
- frataxin
- galactosylceramidase
- gigaxonin
- glycyl-tRNA synthetase 1
- glucosylceramidase beta 2

- 1,4-alpha-glucan branching enzyme 1
- golgi brefeldin A resistant guanine nucleotide exchange factor 1
- ganglioside induced differentiation associated protein 1
- gap junction protein beta 1
- gap junction protein beta 3
- gap junction protein gamma 2
- galactosidase alpha
- G protein subunit beta 4
- gelsolin
- hydroxyacyl-CoA dehydrogenase trifunctional multienzyme complex subunit alpha
- hydroxyacyl-CoA dehydrogenase trifunctional multienzyme complex subunit beta
- histidyl-tRNA synthetase 1
- histidine triad nucleotide binding protein 1
- hexokinase 1
- heterogeneous nuclear ribonucleoprotein A1
- homeobox D10
- heat shock protein family B (small) member 1
- heat shock protein family B (small) member 3
- heat shock protein family B (small) member 8
- isoleucyl-tRNA synthetase 2, mitochondrial
- interferon related developmental regulator 1
- immunoglobulin mu DNA binding protein 2
- inverted formin, FH2 and WH2 domain containing
- inositol 1,4,5-trisphosphate receptor type 3
- jagged canonical Notch ligand 1
- kinesin family member 1A
- kinesin family member 5A
- lipopolysaccharide induced TNF factor
- lamin A/C
- leucine rich repeat and sterile alpha motif containing 1
- lysosomal trafficking regulator
- minichromosome maintenance complex component 3 associated protein
- mitofusin 2

- metabolism of cobalamin associated C
- membrane metalloendopeptidase
- MORC family CW-type zinc finger 2
- mitochondrial inner membrane protein MPV17
- myelin protein zero
- myotubularin related protein 2
- mitochondrial translation release factor in rescue
- myosin heavy chain 14
- alpha-N-acetylgalactosaminidase
- N-acetyl-alpha-glucosaminidase
- asparaginyl-tRNA synthetase 1
- N-myc downstream regulated 1
- neurofilament heavy chain
- neurofilament light chain
- nerve growth factor
- neurotrophic receptor tyrosine kinase 1
- phosphate cytidylyltransferase 2, ethanolamine
- pyruvate dehydrogenase kinase 3
- peroxisomal biogenesis factor 7
- phytanoyl-CoA 2-hydroxylase
- piezo type mechanosensitive ion channel component 2
- phospholipase D family member 3
- pleckstrin homology and RhoGEF domain containing G5
- proteolipid protein 1
- phosphomannomutase 2
- peripheral myelin protein 2
- peripheral myelin protein 22
- polynucleotide kinase 3'-phosphatase
- patatin like phospholipase domain containing 6
- DNA polymerase gamma, catalytic subunit
- RNA polymerase III subunit B
- protoporphyrinogen oxidase
- PR/SET domain 12

- protein kinase C gamma
- prion protein
- phosphoribosyl pyrophosphate synthetase 1
- periaxin
- phosphatase and tensin homolog
- protein tyrosine phosphatase non-receptor type 11
- peptidyl-tRNA hydrolase 2
- RAB7A, member RAS oncogene family
- receptor accessory protein 1
- reticulophagy regulator 1
- replication factor C subunit 1
- sacsin molecular chaperone
- SET binding factor 1
- SET binding factor 2
- scavenger receptor class B member 2
- sodium voltage-gated channel alpha subunit 10
- sodium voltage-gated channel alpha subunit 11
- sodium voltage-gated channel alpha subunit 9
- synthesis of cytochrome C oxidase 2
- SCY1 like pseudokinase 1
- septin 9
- senataxin
- SH3 domain and tetratricopeptide repeats 2
- sigma non-opioid intracellular receptor 1
- solute carrier family 12 member 6
- solute carrier family 25 member 19
- solute carrier family 25 member 46
- solute carrier family 52 member 2
- solute carrier family 52 member 3
- solute carrier family 5 member 7
- sorbitol dehydrogenase
- SRY-box transcription factor 10
- spastin

- SPG11 vesicle trafficking associated, spatacsin
- SPG7 matrix AAA peptidase subunit, paraplegin
- spectrin alpha, non-erythrocytic 1
- spectrin beta, non-erythrocytic 4
- serine palmitoyltransferase long chain base subunit 1
- serine palmitoyltransferase long chain base subunit 2
- serine palmitoyltransferase long chain base subunit 3
- SURF1 cytochrome c oxidase assembly factor
- synaptotagmin 2
- tubulin folding cofactor E
- TBC1 domain containing kinase
- tectonin beta-propeller repeat containing 2
- trafficking from ER to golgi regulator
- tripartite motif containing 2
- transient receptor potential cation channel subfamily V member 4
- alpha tocopherol transfer protein
- transthyretin
- tubulin beta 3 class III
- twinkle mtDNA helicase
- ubiquitin C-terminal hydrolase L1
- VAMP associated protein B and C
- valosin containing protein
- vacuolar protein sorting 13 homolog A
- VRK serine/threonine kinase 1
- von Willebrand factor A domain containing 1
- tryptophanyl-tRNA synthetase 1
- WNK lysine deficient protein kinase 1
- X-ray repair cross complementing 1
- tyrosyl-tRNA synthetase 1
- zinc finger homeobox 2
- zinc finger FYVE-type containing 26

## Related Gene Panels

- [Neuropathy \(genepanel\) - UZA](#)

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