# DISEASE:
# Hereditary neuropathy with liability to pressure palsies

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| **NAME:** | Hereditary neuropathy with liability to pressure palsies |
| **DESCRIPTION:** | A rare neurologic disease characterized by recurrent mononeuropathies usually triggered by minor physical activities innocuous to healthy people. |
| **ORPHACODE:** | 640 |
| **SYNONYMS:** | Current pressure-sensitive neuropathy<br>HNPP<br>Heterozygous microdeletion 17p11.2p12<br>Potato-grubbing palsy<br>Tomaculous neuropathy<br>Tulip-bulb digger's palsy |
| **XREF(S):** | Orphanet<br>MedDRA<br>ICD-10<br>OMIM |
| **ANALYTE(S):** | PMP22<br>PMP22 |
| **CREATED:** | 13 May 2019 - 01:02 |

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| **CHANGED:** | 22 Jun 2023 - 16:14 |

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Source URL: http://gentest.healthdata.be/index.php/index.php/disease/160

# RELATED CONTENT

## Related Genetic Tests

- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Charcot-Marie-Tooth type 1A / Hereditary Neuropathy with Liability to Pressure Palsies
- Charcot-Marie-Tooth type 1A / Hereditary Neuropathy with Liability to Pressure Palsies
- Charcot-Marie-Tooth type 1A / Hereditary Neuropathy with Liability to Pressure Palsies
- Charcot-Marie-Tooth type 1A (CMT1A) / Hereditary Neuropathy with Liability to Pressure Palsies
- Charcot-Marie-Tooth type 1A (CMT1A) / Hereditary Neuropathy with Liability to Pressure Palsies (HNPP)
- Hereditary Neuropathy with Liability to Pressure Palsies (HNPP)
- Neuropathy (gene panel)

## Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

## Related Analytes

- peripheral myelin protein 22

## Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Neuropathy (148 genes) - IPG

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