HINT1 neuropathy in Lithuania: clinical, genetic, and functional profiling

Matilde Malcorps\textsuperscript{1,2†}, Silvia Amor-Barris\textsuperscript{1,2†}, Birute Burnyte\textsuperscript{3†}, Ramune Vilimiene\textsuperscript{4}, Camila Armriola-Ricaute\textsuperscript{1,2}, Kristina Grigalioniene\textsuperscript{3}, Alexandra Ekshteyn\textsuperscript{1,2}, Ausras Morkuniene\textsuperscript{3}, Arunas Vaitkevicius\textsuperscript{4}, Els De Vriendt\textsuperscript{1,2}, Jonathan Baets\textsuperscript{5,6,7}, Steven S. Scherer\textsuperscript{8}, Laima Ambrozaityte\textsuperscript{3}, Algirdas Utkus\textsuperscript{3}, Albena Jordanova\textsuperscript{1,2,9*†} and Kristien Peeters\textsuperscript{1,2†}

Abstract

Background: Recessive loss-of-function variations in HINT1 cause a peculiar subtype of Charcot-Marie-Tooth disease: neuromyotonia and axonal neuropathy (NMAN; OMIM\#137200). With 25 causal variants identified worldwide, HINT1 mutations are among the most common causes of recessive neuropathy. The majority of patients are compound heterozygous or homozygous for a Slavic founder variant (c.110G>C, p.Arg37Pro) that has spread throughout Eurasia and America.

Results: In a cohort of 46 genetically unresolved Lithuanian patients with suspected inherited neuropathy, we identified eight families with HINT1 biallelic variations. Most patients displayed sensorimotor or motor-predominant axonal polyneuropathy and were homozygous for the p.Arg37Pro variant. However, in three families we identified a novel variant (c.299A>G, p.Glu100Gly). The same variant was also found in an American patient with distal hereditary motor neuropathy in compound heterozygous state (p.Arg37Pro/p.Glu100Gly). Haplotype analysis demonstrated a shared chromosomal region of 1.9 Mb between all p.Glu100Gly carriers, suggesting a founder effect. Functional characterization showed that the p.Glu100Gly variant renders a catalytically active enzyme, yet highly unstable in patient cells, thus supporting a loss-of-function mechanism.

Conclusion: Our findings broaden NMAN’s genetic epidemiology and have implications for the molecular diagnostics of inherited neuropathies in the Baltic region and beyond. Moreover, we provide mechanistic insights allowing patient stratification for future treatment strategies.

Keywords: Peripheral neuropathy, Charcot-Marie-Tooth disease, HINT1, Neuromyotonia, Lithuania

\textsuperscript{1}Matilde Malcorps, Silvia Amor-Barris and Birute Burnyte contributed equally to this study as co-first authors

\textsuperscript{2}Albena Jordanova and Kristien Peeters contributed equally to this study as co-last authors

\textsuperscript{9}Correspondence: Albena.jordanova@uantwerpen.vib.be

\textsuperscript{1}Molecular Neurogenomics Group, VIB Center for Molecular Neurology, VIB, Antwerp, Belgium

Full list of author information is available at the end of the article

Background

Biallelic loss-of-function alterations in the histidine triad nucleotide-binding protein 1 (HINT1) cause neuromyotonia and axonal neuropathy (NMAN [OMIM\#137200]) [1]. Patients with HINT1-deficiency show progressive, predominantly motor polyneuropathy typically starting in the first decade of life, leading to lower limb weakness and gait impairment [2]. Neuromyotonia—peripheral nerve hyperexcitability manifesting as spontaneous muscular activity at rest and delayed muscle relaxation after voluntary contraction—is a clinical hallmark of NMAN.
Neuromyotonia is a striking and recognizable feature upon needle electromyography routinely performed in the diagnostic work-up of patients with peripheral neuropathy. So far, the role of HINT1 in the peripheral nervous system is unexplored.

HINT1 encodes a ubiquitous homodimeric purine phosphoramidase belonging to the evolutionary conserved histidine-triad superfamily. In vitro, HINT1 is a promiscuous enzyme, hydrolyzing diverse AMP-linked substrates [3] and acts as a SUMO1-cleaving Cys-protease [4], yet its endogenous substrate(s) remain unknown. HINT1 has been attributed pleiotropic cellular roles, including regulation of transcription factors involved in tumor progression and apoptosis [5, 6], modulating G-protein coupled receptor signaling [7], and controlling calcium signaling via the store-operated calcium entry pathway [8].

Currently, 25 causal variants have been identified in over 100 NMAN-patients from Europe, Asia and America [1, 2, 9–12]. Haplotype analysis demonstrated founder effects for four of the recurrent HINT1 mutations in Europe [1, 2, 13] (p.Arg37Pro, p.Cys84Arg, p.Arg95Gln, p.His112Asn) and one in China [9] (p.Cys38Arg) explaining the elevated prevalence of NMAN in certain geographical areas. The ancient Slavic founder allele p.Arg37Pro, present in the majority of NMAN-patients, has a particularly high carrier frequency (1:67-250) in Central and South-East Europe, Russia and Turkey [1, 10, 14]. As a consequence, NMAN ranks among the most frequent forms of axonal neuropathy in those regions.

Most causal NMAN-variants are recurrent missense changes, targeting conserved but also less conserved amino acid residues all over the protein sequence, making it difficult to assess the pathogenicity of a novel variant based on the commonly used criteria like frequency, conservation, or position. Moreover, there is increasing evidence that the disease-causing alterations have differential effects on HINT1 protein stability and function. This has important implications for future therapeutic strategies, as the mutational category will determine a patient’s treatment options. Therefore, functional characterization of novel HINT1 variations benefits both diagnostics and patient stratification. NMAN-associated HINT1 alterations cause a loss of (enzymatic) function, because they either lead to unstable protein or transcript, or affect key residues in the catalytic cleft [1]. Genetic complementation testing in a HINT1-knockout (KO) growth deficient yeast strain proved that yeast and human HINT1 orthologs are functionally conserved and that the NMAN-variations abolish this function [1]. Notably, this overexpression system can be used to assess the activity of NMAN-proteins that are degraded in endogenous conditions [13]. Here, we performed the first systematic assessment of HINT1 neuropathy in Lithuania and describe a potential new founder event in the Baltic region.

**Results**

**Clinical findings in families with the novel p.Glu100Gly variant**

We studied nine families carrying HINT1 mutations (Table 1). In four of them, a novel variant was found to segregate with the peripheral neuropathy in a homozygous or compound heterozygous state. Their clinical findings are presented below.

The index patient in family Lit1 was a 45-year-old man who developed slowly progressive weakness of his lower limbs, foot drop, difficulty in walking, and frequent falls beginning at age 10 (Table 1). He underwent bilateral surgical correction of foot deformities at age 30, and developed difficulty to straighten fingers and grasp small objects at age 35. The physical examination at age 45 revealed muscle wasting and weakness in distal upper and lower limbs, bilateral pes cavus, foot drop and atrophy of the intrinsic hand muscles and thenar eminence. He did not complain of sensory impairment. His family history was negative for neuromuscular disorders.

In family Lit2, a 40-year-old woman presented with gait impairment, distal weakness of lower limbs and bilateral foot drop at the age of 12. Muscle weakness has slowly progressed in a length-dependent manner. Later, she developed difficulties in releasing grip after a strong voluntary hand contraction. She had undergone bilateral achillotomy. The physical examination (Fig. 1A–C) revealed muscle wasting of the lower limbs, and severe distal weakness that was more prominent in the legs. Diminished biceps brachii, triceps brachii, patellar reflexes and absent ankle reflexes were observed. Sensory examination and coordination were normal. Her 44-year-old brother (Fig. 1D–F) was also affected and presented at the early childhood with slowly progressive weakness and muscle wasting of his feet and calves, foot drop, gait impairment and frequent falls. He noticed hand wasting and weakness at the age of 20.

In family Lit3, two affected sisters, now 18 and 19 years old, presented with bilateral foot drop, gait impairment and exercise intolerance at age of 4, followed by gradual progression of muscle weakness and wasting. Patients complained about muscle cramps. Their physical examination revealed muscle wasting of the lower limbs, bilateral foot drop, severe distal weakness that was more prominent in the legs. Diminished reflexes in the upper limbs and absent reflexes in the lower limbs were observed. Sensory evaluation and coordination were normal in both siblings.
Patient USA1 is a 58-year-old man who had a distal hereditary motor neuropathy with a clinical onset at age 12. Regular neurological follow-up since age 36 showed slowly progressive weakness and atrophy predominantly in distal lower limbs, which in the recent years rendered him mostly wheelchair-dependent. The patient now has
severe muscle wasting of the lower limbs, bilateral foot drop, absent tendon reflexes. Although initially distal upper limb strength was mildly affected, over the last couple of years, atrophy and weakness in the intrinsic hand muscles increased substantially. Vibration and pinprick sensation had been normal until age 46, but have subsequently declined to the point that vibration is now reduced at the big toes (1 with a Rydell-Seiffer tuning fork) and pinprick is reduced to just below the knees. The patient had prominent muscle twitches and paresthesia in both distal lower and upper limbs; these largely disappeared in his 30s. The patient suffered from chronic anxiety, depression, and insomnia, which was treated with multiple medications including bupropion,
buspirone, gabapentin, and melatonin. His two siblings were unaffected.

**Electrophysiological studies**

In patient Lit1.1, nerve conduction studies (NCS) performed at age 45 revealed a marked reduction of compound muscle action potentials (CMAPs) in upper limbs and absent in the lower limbs (Table 2). The moderate NCS slowing of the median and the ulnar responses is likely caused by the reduction of CMAPs. Results of the sensory nerve conduction studies were not available. Needle electromyography (EMG) revealed evidence of reinnervation with sparse spontaneous activity in the muscles of the hands.

NCS of patient Lit2.4 at age 40 revealed pure motor axonal polyneuropathy in the lower limbs. Sural nerve action potential and conduction velocity were normal. The motor and sensory responses in the upper limbs were normal. EMG showed chronic reinnervation without hallmarks of active denervation. There were several neuromyotonic discharges in the first dorsal interosseous and deltoid muscles. The NCS of her affected brother was not available but was reported as a severe motor axonal neuropathy with absent peroneal CMAPs.

In family Lit3, assessment of peripheral nerves of the hands revealed similar findings in both siblings at the ages of 18 and 19, respectively. CMAPs, sensory nerve action potentials, and conduction velocities were normal. Prolonged distal latencies of the motor median and the motor ulnar nerve on the right were detected. The findings of the NCS of the lower limbs were compatible with axonal motor polyneuropathy, with preserved sensory responses. EMG recording provides evidence of a neurogenic pattern with sparse spontaneous activity. In the younger sister Lit3.4, EMG displayed neuromyotonic discharges in the first dorsal interosseous muscle.

The NCSs of patient USA1 at ages 40 and 47 showed a progressive chronic length-related motor axonal neuropathy. Peroneal motor nerve responses were absent in both studies. The right tibial CMAP amplitude was barely detectable in the first study and undetectable in the follow-up. The right median nerve motor responses were normal, and right ulnar CMAP amplitude was mildly decreased in both studies. Needle EMG showed signs of length-related chronic denervation in the muscles of the right arm that was severe in distal muscles. The sensory nerve responses in the arm and leg were normal at both time points.

**Genetic results**

We investigated the occurrence of HINT1 neuropathy in Lithuania by testing a cohort of 46 unrelated index patients with suspected peripheral neuropathy that were excluded for the most common genetic causes: *PMP22* duplication/deletion, *GJB1, MFN2* and *MPZ*. Using next-generation sequencing targeting a custom-designed panel of 150 genes associated with Charcot-Marie-Tooth disease and related hereditary neuropathies, we identified biallelic variations in *HINT1* in eight index patients: five affected individuals carried the most common known pathogenic variant NM_005340.7:c.110G>C (p.Arg37Pro) in homozygous state, two patients carried it in compound heterozygous state together with a variant of unknown significance (VUS) NM_005340.7:c.299A>G (p.Glu100Gly), and finally, one patient carried this VUS in homozygous state (Table 1). In addition, in our in-house database of neuropathy patients, we identified an American individual with distal motor neuropathy who carried the same compound heterozygous *HINT1* variation (c.110G>C/c.299A>G). Segregation analysis was performed in two families harboring the unknown c.299A>G variant (Lit2 and Lit3), demonstrating compliance with a recessive inheritance model and uncovering two additional affected individuals (siblings of the index patients) with biallelic *HINT1* variants (Fig. 1G).

The three Lithuanian families carrying the novel c.299A>G *HINT1* variation (Lit1-3) originated from different ethnolinguistic regions in the country and were seemingly unrelated. Yet, haplotype analysis of all c.299A>G variant carriers, including the American patient who had a Polish ancestry, demonstrated a shared chromosomal background of 1.9 Mb (D5S809-D5S2110) surrounding the VUS, suggestive of a single mutational origin. We also confirmed that the c.110G>C carriers share the previously established disease haplotype surrounding this known founder allele (Fig. 1G) [1].

In the global population the c.299A>G variation is extremely rare; for instance, in the Genome Aggregation Database (GnomAD v2.1.1) [15] it was observed only once in heterozygous state in a non-Finnish European female (allele frequency 1:251,448). In a cohort of 98 unrelated Lithuanian control individuals, the c.110G>C variation was observed once in heterozygous state (allele frequency 1:196) and the c.229A>G variant was not detected.

The p.Glu100Gly substitution targets a conserved residue (GERP: 3.52) but its predicted effect on HINT1 function is variable (Polyphen-2 v2.2.2r398: probably benign, score = 0.050; SIFT v6.2.1: tolerated, score = 0.38; Mutation Taster: disease causing, probability = 1.00) [16–18].

**Functional characterization**

To investigate the impact of the p.Glu100Gly VUS on HINT1 protein function, we performed genetic complementation testing in yeast [1, 13]. Using a vector with a
| ID  | Age (y) | Side | Motor | Sensory | EMG |
|-----|---------|------|-------|---------|-----|
|     |         |      | Median | Ulnar | Peroneal | Tibial | Median | Ulnar | Peroneal | Tibial | Median | Ulnar | Sural |
|     |         |      | CMAP (mV) | CMAP (mV) | CV (m/s) | CMAP (mV) | CV (m/s) | SNAP (µV) | CV (m/s) | SNAP (µV) | CV (m/s) | SNAP (µV) | CV (m/s) |
| Lit1.1 | 45 | L | 0.09 | 18.2 | 0.39 | 39.8 | ND | ND | ND | ND | NA | NA | NA | ND | ND | + | ND |
| Lit2.4 | 40 | R | 7.6 | 52.5 | 6.62 | 56.4 | ND | ND | 0.55 | 35.1 | 750 | 52.2 | 57.4 | 50.5 | 27.8 | 45.5 | + | + |
| Lit3.3 | 18 | R | 5.15 | 51.3 | 7.7 | 46.8 | 1.12 | 34.4 | 2.9 | 36.7 | 92.1 | 57.5 | 57.0 | 61.1 | 21.5 | 42.1 | + | ND |
| Lit3.4 | 19 | R | 6.4 | 48.8 | 3.87 | 58.8 | ND | ND | 0.8 | 34.8 | 95.0 | 60.2 | 78.3 | 53.9 | 20.1 | 36.5 | + | + |
| Lit4 | 18 | R | 0.32 | 47.9 | 0.78 | 50.0 | ND | ND | 0.3 | 39.7 | 32.0 | 53.3 | 26.0 | 52.6 | – | – | + | + |
| Lit5 | 21 | R | 5.13 | 49.0 | 3.2 | 49.0 | 0.77 | 35.9 | 2.3 | 36.7 | 37.3 | 53.3 | 140 | 52.9 | ND | ND | + | ND |
| Lit6 | 15 | R | 4.3 | 50.0 | 1.2 | 50.0 | – | – | – | – | – | – | – | – | – | – | + | + |
| Lit7 | 42 | R | ND | ND | 0.8 | 50.0 | ND | ND | ND | ND | 33.3 | 54.2 | 313 | 50.0 | ND | ND | ND | ND |
| Lit8 | 33 | R | 8.7 | 47.0 | 4.1 | 46.2 | ND | ND | ND | ND | 23.5 | 47.1 | 12.8 | 45.1 | ND | ND | ND | ND |
| USA1 | 41 | R | 98 | 52 | 3.4 | 57 | ND | ND | 0.08 | 40 | 27.2 | 58 | 8.4 | 63 | 164 | 34 | + | ND |
| USA1 | 47 | R | 12.4 | 50 | 3.9 | 57 | ND | ND | ND | ND | 19.5 | 61 | 9.0 | 50 | 104 | 41 | + | ND |

Age (y), age at examination in years; R, right; L, left; CMAP, complex motor amplitude potential; CV, conduction velocity; NA, not available; ND, not detected (no response); SNAP, sensory nerve action potential; Sp. Act., spontaneous activity; N. dis., neuromyotonic discharges; –, not measured; +, present.
strong, constitutive promotor, we over-expressed human HINT1 transgenes carrying different variants into a HNT1 KO yeast strain [3]. In contrast to the p.Arg37Pro alteration, which has a detrimental effect on HINT1 protein stability both in yeast and in human cells [1], the p.Glu100Gly variant resulted in substantial HINT1 protein expression in yeast (Fig. 2A). Importantly, this mutated protein appears to retain activity, because it showed (partial) rescue of the restrictive growth deficiency associated with loss of yeast HNT1 (Fig. 2B). Notably, previous study demonstrated that the overexpression system in yeast can lead to enhanced protein expression compared to the endogenous situation [13], therefore increasing artificially the residual activity of the mutant protein, prompting us to interpret the observed rescue effect with caution. Therefore, we analyzed the HINT1 protein levels in peripheral blood mononuclear cells extracted from patients carrying homozygous (Lit2.4) or compound heterozygous (Lit3.3) p.Glu100Gly variants. Immunoblotting analysis showed no detectable HINT1 protein expression in both subjects (Fig. 2C), while the HINT1 transcript was expressed (Fig. 2D), indicating that the p.Glu100Gly substitution causes severe protein instability in patient cells. The findings were in line with known causal variants (e.g. p.Arg37Pro) where we previously demonstrated that the lack of protein is a result of a post-translational event [1, 13]. In combination with the genetic findings, these functional results provide convincing evidence for pathogenicity of the p.Glu100Gly variant.

Discussion
This is the first systematic assessment of NMAN in Lithuania, where we identified a total of eight patients from 46 families carrying biallelic missense variations in the HINT1 gene: the known p.Arg37Pro and/or the novel p.Glu100Gly. The same compound heterozygous variants were identified in a patient from the USA. In line with previous studies, patients had motor impairment predominating in the distal lower limbs and starting from the first decade of life. Neuromyotonia was reported in some but not all of them. Mild sensory symptoms were only present in a minority of the cases. In addition, some patients displayed atypical features that have been described before, such as developmental delay and intellectual disability [19, 20], speech delay [21], and mood disorder [13, 22]. However, bulbar weakness in the

Fig. 2 Functional characterization of the identified HINT1 variants. A Western blot analysis of protein extract from HNT1-deleted yeast strain expressing human HINT1, either wildtype (hWT) or the p.Arg37Pro or p.Glu100Gly alleles. Equal loading was validated with mouse monoclonal anti-PGK1 antibody and relative HINT1 expression was normalized to hWT. The graph represents relative quantification of band intensities of four independent replicates. Note the severe reduction of HINT1 in the p.Arg37Pro-expressing yeast, and the modest reduction of HINT1 in the p.Glu100Gly expressing yeast. B Genetic complementation analysis in HNT1-deleted yeast strain performed by spot assay. Serial dilutions of the different yeast strains were spotted on minimal media without leucine, supplemented with either 2% glucose or 2% galactose, and incubated at 39 °C for 3 days. Note the reduction of growth of yeast expressing p.Arg37Pro or p.Glu100Gly compared to hWT under the restrictive conditions. C Western blot analysis of total protein extracts from HINT1 patient Lit2.4 (p.Glu100Gly/p.Glu100Gly) and patient Lit3.3 (p.Glu100Gly/p.Arg37Pro) cells or control lymphoblasts. Membranes were immunoblotted with polyclonal rabbit anti-human HINT1 antibody. Equal loading was validated with mouse monoclonal anti-β-actin antibody. The bar charts represent the means with standard error of the mean (s.e.m.) of the relative quantification of band intensities of three independent replicates. Note the severe reduction of HINT1 from the patient samples. Statistical one-way ANOVA analysis was performed. ns = not significant; ***p<0.001; ****p<0.0001. D Sanger sequencing traces of HINT1 cDNA isolated from peripheral blood mononuclear cells of patient Lit2.4 (p.Glu100Gly/p.Glu100Gly) and patient Lit3.3 (p.Glu100Gly/p.Arg37Pro) or a control individual. The c.299A>G transition is framed. Note the comparable intensity of the peaks at the c.299 position in the compound heterozygous patient.
form of dysphagia, rhinophonia, dysphonia, dysarthria has never been reported before.

Haplotype analysis confirmed that the c.299A>G transition in the six identified patients and their relatives originated from a single founder event. The variant is extremely rare, and a search of public genetic variation databases resulted in only a single heterozygous carrier. This individual is (non-Finnish) European, but no further details about ethnicity or nationality are known. As opposed to the most common c.110G>C variant, that was found in 1 out of 98 Lithuanian control individuals, the novel c.299A>G variation was not detected in WGS (n = 50) (unpublished data) and genotype (n = 399) data [23], further exemplifying the rarity of this allele. The American patient carried two recurrent disease-causing HINT1 variants that cluster in Europe (c.110G>C/ c.299A>G) and shared the same disease haplotype as the Lithuanians for both variants. He has Eastern-European heritage, as both his maternal and paternal grandparents emigrated to the US from Poland, a neighboring country of Lithuania and of the Czech Republic, which has among the highest known carrier rates of c.110G>C in Europe (1:182) [14]. Taken together, our results confirm the existence of another pathogenic founder allele in the HINT1 gene, p.Glu100Gly, that may have originated in the Baltic region. Moreover, our findings expand the geographical distribution of the p.Arg37Pro disease haplotype to the Baltic region [2].

Functional characterization of the novel p.Glu100Gly variant revealed that it causes severe protein degradation in patient cells, providing strong evidence for pathogenicity following a loss-of-function disease mechanism. These results are in line with previous studies showing over 80% of pathogenic HINT1 substitutions to trigger proteasome-mediated protein degradation [1, 13]. Contrastingly, this degradation did not occur to the same extent in the yeast overexpression model, enabling functionality testing of the residual protein. Notably, this experiment proved that the p.Glu100Gly variant gives rise to a HINT1 protein that retains its activity. Therefore, it fits into the same category as other HINT1 variants like p.Cys84Arg, which also renders a protein that remains catalytically active, as seen in an in vitro enzymatic assay, but is degraded in patient cells [1, 24]. This is in contrast to other NMAN-causing mutations that lead to stable but enzymatically dead (e.g. p.His112Asn [1]) or unstable and enzymatically dead (e.g. p.Arg95Gln [13]) protein. To this end, the results of this study have important implications in light of future therapy development. Affected individuals carrying variations like p.Glu100Gly, belong to a subgroup of patients who would benefit from treatment with a pharmacological chaperone that stabilizes the affected yet still catalytically active HINT1 enzyme. Similar approach has been developed for other recessive disorders like cystic fibrosis [25].

On the HINT1 protein structure, p.Glu100 is positioned at the far edge of the dimer interface, a region where multiple NMAN-causing variations cluster (p.Gly93Asp, p.Tyr94Cys, p.Arg95Gln, p.Val97Met). It has been shown before that HINT1 dimerization is crucial to retain enzymatic activity [24]. Through genetic complementation testing in yeast we established that, despite its localization at the dimer interface, the p.Glu100Gly substitution does not seem to abolish the capability of the HINT1 enzyme to form dimers, because the overexpressed protein is functionally active. More likely, the loss of the glutamate side chain at this position could disturb the internal structure of the monomer, similar to other NMAN-variations that preserve dimerization (e.g. p.Cys84Arg, p.Gly89Val), which show reduced thermal stability compared to the wildtype protein [24].

Conclusions
This study represents the first analysis of HINT1 neuropathy in Lithuania, where we identified a rare novel pathogenic allele (p.Glu100Gly). Functional characterization in yeast and patient cells provided mechanistic insights on how the newly reported substitution leads to loss of HINT1 function. The patients displayed typical symptoms associated with HINT1 neuropathy, including motor impairment in distal lower limbs predominant from the first decade of life, but also some atypical features such as developmental delay and mood problems. Our findings expand the genetic epidemiology of HINT1-related disorders.

Methods
Patients and evaluation
Patients underwent a routine neurological examination. The family history was taken in all cases. Age of onset was determined by asking about the first neuropathy related symptoms. Nerve conduction studies (NCS) were performed using standard techniques. Clinical data and biological samples were collected for all the patients and their relatives whenever possible.

Sequencing analysis
Genomic DNA of the Lithuanian patients and their relatives was extracted from peripheral leukocytes using standard methods. Next generation sequencing (NGS) was performed using a custom-designed gene panel of 150 genes associated with Charcot-Marie-Tooth disease and related hereditary neuropathies for one affected person from each family. The panel was designed for Ion AmpliSeq™ technology (Ion Torrent, Thermo Fisher Scientific). The DNA libraries were sequenced on an
Ion PGM™ Sequencer (Life Technologies). Bioinformatic analysis was performed including alignment of raw sequence reads to a reference human genome and variant calling on the Ion Torrent Suite™ Server. For identification of disease-causing variants, annotation and filtration of identified sequence variants was performed, using ANNOVAR software [26]. Variants with population frequency over 1% in the Single Nucleotide Polymorphism (dbSNP v137), Genomes Aggregation (gnomAD[15]) and 1000 Genomes Project [27] databases were filtered out. Only variants predicted to affect the coding regions (including non-synonymous, predicted missense, nonsense, splice acceptor and donor site, and insertions or deletions) were selected for further analysis. Several in silico prediction programs (PolyPhen-2 [16], Mutation-Taster [17], SIFT [18]) were used to predict the functional effect as well as the genomic evolutionary rate profiling (GERP) [28] score. Segregation analysis was performed by Sanger sequencing [29].

Genomic DNA was extracted from peripheral blood sample of the US patient using standard procedures. All three exons as well as the 5’ and 3’ UTR of HINT1 were screened for variants by Sanger Sequencing. PCR products were purified using ExoSAP-IT™ (Thermo Fisher Scientific, Massachusetts, USA) and sequenced in both directions on a 3730xl DNA Analyzer (Applied Biosystems, Foster City, CA, USA). SNP genotyping was done by Sanger sequencing of the purified plasmid.

Yeast strain and transformation
*S. cerevisiae* strain BY8-5c (MATα *ura3*Δ52 *his3*Δ200 *trplΔ901* *lys2*-801 *suc2*-Δ9 *leu2*-3,112 *hnt1Δ::URA3*) [3] was provided by Dr. Brenner, University of Iowa, USA. Yeast cells were cultured in rich medium (YPD). Transformation of BY8-5c with the pAG415GPD expression plasmids carrying one of the HINT1 variants or the human wild-type was done with the LiAc/SS carrier DNA/PEG method [30]. Positive clones were selected in minimal medium without Leucine (SD-Leu) supplemented with 2% glucose.

Spot assay in yeast
Pre-cultures of the different yeast clones were grown overnight in SD-Leu supplemented with glucose. Absorbance was measured and adjusted to an optical density of OD_{600nm} = 5. Serial dilutions of each culture were spotted in 5ul drops on SD-Leu agar plates supplemented either with 2% glucose or 2% galactose. Plates were incubated for three days at 39°C.

Cell line establishment and culture
Peripheral blood lymphocytes were isolated using a Ficol Paque gradient and subsequently transformed with Epstein-Barr virus. After a two-hour incubation at 37°C, cells were centrifuged and re-suspended in RPMI complete medium (Invitrogen, Carlsbad, CA, USA) supplemented with 1% phytohaemagglutinin. Cells were seeded on a 24-well plate and incubated at 37°C and 5% CO₂ for three days. After establishment, lymphoblastoid cells were grown in RPMI complete medium containing 15% fetal bovine serum (FBS, Gibco, Waltham, MA, USA), 1% sodium pyruvate, 1% L-Glutamine (Gibco, Waltham, MA, USA) and 1% penicillin/streptomycin (Gibco, Waltham, MA, USA).

cDNA analysis
Total RNA was isolated from peripheral blood mononuclear cells using the Universal RNA kit (Roboklon GnmG) according to the manufacturer’s instructions and was subsequently treated with DNase (TURBO DNA-free kit, Applied Biosystems). cDNA was synthesized by RT-PCR with random hexamers using the iScript Advanced cDNA Synthesis Kit (Bio-Rad Laboratories). Full length *HINT1* cDNA was amplified by PCR and the amplicons were Sanger sequenced and analyzed as described above. The sequences of the cDNA primers are available upon request.
Immunoblotting

Human cells were lysed in RIPA lysis buffer (20 mM Tris–HCl pH = 7.4; 150 mM NaCl; 0.1% Nonidet P-40; 0.5% sodium deoxycholate; 0.1% sodium dodecyl sulfate) supplemented with Halt™ Protease Inhibitor Cocktail (ThermoFisher Scientific, Waltham, MA, USA). Protein concentration was determined with the Pierce BCA protein assay kit (ThermoFisher Scientific, Waltham, MA, USA) and adjusted to 20 μg per sample. Lysates were boiled for five minutes in reducing Laemmli sample buffer (Bio-Rad, Hercules, CA, USA) supplemented with 100 mM 1,4-Dithiothreitol (DTT).

Yeasts proteins were extracted following a previously published protocol [31]. Briefly, cells were collected before stationary phase (OD600nm = 1) by centrifugation. Then cells were washed first with 2.0 M LiAc and then 0.4 M NaOH for 5 min on ice. Cells were finally boiled for five min in Laemmli sample buffer (Bio-Rad, Hercules, CA, USA) supplemented with 100 mM DTT.

Proteins were separated in 4–15% Mini-PROTEAN® TGX Stain Free™ Protein gels (Bio-Rad, Hercules, CA, USA) and transferred to a nitrocellulose membrane (Hybond™-P, GE Healthcare, Chicago, IL, USA) using the semi-dry Trans-Blot™ Turbo™ Transfer System (Bio-Rad, Hercules, CA, USA). Membranes were blocked for an hour at room temperature with 5% milk powder diluted in PBS supplemented with 0.1% Tween-20 and then incubated with primary antibody overnight at 4°C and one hour with a secondary horseradish peroxidase-conjugated antibody at room temperature. Blots were developed with Enhanced Chemiluminescence ECL Plus™ (ThermoFisher Scientific, Waltham, MA, USA) and imaged with ImageQuant™ LAS 4000 (GE Healthcare, Chicago, IL, USA).

The antibodies used in this study were: polyclonal rabbit anti-human HINT1 antibody (Sigma, San Luis, MO, USA), and to demonstrate equal loading: monoclonal mouse anti-PGK1 antibody (ThermoFisher Scientific, Waltham, MA, USA) or monoclonal mouse anti-β-actin antibody (Sigma, San Luis, MO, USA).

Abbreviations

AMP: Adenosine monophosphate; BCA: Bicinchoninic acid; CMAP: Compound muscle action potential; FBS: Fetal bovine serum; GERP: Genomic evolutionary rate profiling; GJB1: Gap gene beta-1 protein; gnomAD: Genome Aggregation Database; HINT1: Histidine Triad Nucleotide Binding protein 1; HPRT: Hit family protein 1; KO: Knockout; LiAc: Lithium acetate; MFN2: Mitofusin-2; MPZ: Myelin protein zero; NCS: Nerve conduction studies; NGS: Next-generation Sequencing; NMAN: Neuromyotonia, axonal neuropathy; PBS: Phosphate-buffered saline; PEG: Polyethylene glycol; PGK1: Phosphoglycerate kinase 1; PMP22: Peripheral myelin protein 22; SD-Leu: Synthetic defined medium with-leucine; SNP: Single nucleotide polymorphism; STR: Single tandem repeat; SUMO1: Small ubiquitin-like modifier 1; VUS: Variant of unknown significance; YPD: Yeast extract peptone dextrose.

Acknowledgements

We are grateful to the patients and their families for the kind cooperation. We wish to thank the Neuromics Support Facility at VIB-UAntwerp Center of Molecular Neurology for genotyping support.

Author contributions

MM, SAB, AJ, KP: conception and design of the study; MM, SAB, CAR, AE, EDV, AJ, KP, BB, RV, AV, LA, KG, AM, JB, AL, SSP: acquisition and analysis of the data; AJ, KP: BB: drafting the text. All authors read and approved the final manuscript.

Funding

This study was funded by the Research Foundation-Flanders (FWO): Research grants #G.04921.17N and #G0A2122N to A.J.; Postdoctoral fellowship to K.P.; Pre-doctoral fellowships to S.A.B and A.E; Senior Clinical Researcher mandate 1805021N to J.B. The AFMTELETHON: research grant 23708 (to A.J.), Trampoline grant to K.P. The Vilnius University Faculty of Medicine: pre-doctoral fellowship to B.B. SSS’s work was supported by the Judy Setzer Levenson Memorial Fund for CMT Research, the INC (US4NS065712), which is a part of the NCATS Rare Diseases Clinical Research Network (RDCRN), an initiative of the Office of Rare Diseases Research (ORDR), NCATS, funded through a collaboration between NCATS and the NINDS. JB is a member of the European Reference Network for Rare Neuromuscular Diseases (ERN EURO-NMDD) and of the µNEURO Research Centre of Excellence of the University of Antwerp. None of the funding agencies were directly involved in this study.

Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Declarations

Ethics approval and consent to participate

This study was approved by the Vilnius Regional Ethical Committee and the Ethical Committee of the University of Antwerp and the University Hospital Antwerp. Written informed consent was obtained from all participants or their legal guardians before participation.

Consent for publication

Written informed consent was obtained from all participants or their legal guardians before participation.

Competing interests

The authors declare that they have no competing interest.

Author details

1 Molecular Neurogenomics Group, VIB Center for Molecular Neurology, VIB, Antwerp, Belgium. 2 Molecular Neurogenomics Group, Department of Biomedical Sciences, University of Antwerp, Antwerp, Belgium. 3 Department of Human and Medical Genetics, Institute of Biomedical Sciences, Faculty of Medicine, Vilnius University, Vilnius, Lithuania. 4 Institute of Clinical Medicine, Faculty of Medicine, Vilnius University, Vilnius, Lithuania. 5 Translational Neurosciences, Faculty of Medicine and Health Sciences, University of Antwerp, Antwerp, Belgium. 6 Laboratory of Neuromuscular Pathology, Institute Born-Bunge, University of Antwerp, Antwerp, Belgium. 7 Neuromuscular Reference Center, Department of Neurology, Antwerp University Hospital, Antwerp, Belgium. 8 Department of Neurology, The Perelman School of Medicine at the University of Pennsylvania, Philadelphia, PA, USA. 9 Department of Medical Chemistry and Biochemistry, Medical University - Sofia, Sofia, Bulgaria.

Received: 20 May 2022 Accepted: 4 October 2022

Published online: 14 October 2022

References

1. Zimon M, Baets J, Almeida-Souza L, De Vriendt E, Nikodinovic J, Parman Y, et al. Loss-of-function mutations in HINT1 cause axonal neuropathy with neuromyotonia. Nat Genet. 2012;44(10):1080–3.
2. Peeters K, Charnava T, Tournev I, Jordanova A. Axonal neuropathy with neuromyotonia: there is a Hint. Brain. 2017;140(4):868–77.
3. Bieganowski P, Garrison PN, Hodawadekar SC, Faye G, Barnes LD, Brenner C. Adenosine monophosphorohydrolase activity of Hint and Hint1 supports function of K102, CD101, and TFB3. J Biol Chem. 2002;277(13):10852–60.
4. Cortes-Montero E, Rodriguez-Munoz M, Sanchez-Blazquez P, Garzon J. The axonal motor neuropathy-related HINT1 protein is a zinc- and calmodulin-regulated cysteine SUMO protease. Antioxid Redox Signal. 2019;31(7):503–20.
5. Weske J, Huber O. The histidine triad protein Hint1 triggers apoptosis independent of its enzymatic activity. J Biol Chem. 2006;281(37):27356–66.
6. Weske J, Huber O. The histidine triad protein Hint1 interacts with Ponnin and Repitin and inhibits TCF-beta-catenin-mediated transcription. J Cell Sci. 2005;188(14):3117–29.
7. Rodriguez-Munoz M, Cortes-Montero E, Pozo-Rodrigalvarez A, Sanchez-Blazquez P, Garzon-Alvina J. The ON:OFF switch, sigma1R-HINT1 protein, controls GPCR-NMDA receptor cross-regulation: implications in neurological disorders. Oncotarget. 2015;6(34):35458–77.
8. Linde C, Feng B, Wang JB, Golovina VA. Histidine triad nucleotide-binding protein 1 (HINT1) regulates Ca(2+)-signaling in mouse fibroblasts and neuronal cells via store-operated Ca(2+)-entry pathway. Am J Physiol Cell Physiol. 2013;304(11):C1098–104.
9. Meng L, Fu J, H Z, Zhang W, Wang Z, Yuan Y. Novel mutations in HINT1 gene cause autosomal recessive axonal neuropathy with neuromyotonia in two cases of sensorimotor neuropathy and one case of motor neuropathy. Neuromuscul Disord. 2018;28(8):646–51.
10. Schiagina OA, Milovidova TB, Murzatina AF, Rudenskaya GE, Nikitin SS, Dadali EL, et al. HINT1 gene pathogenic variants: the most common cause of recessive hereditary motor and sensory neuropathies in Russian patients. Mol Biol Rep. 2020;47:1331–6.
11. Wang Z, Lin J, Qiao K, Cai S, Zhang W, Zhao C, et al. Novel mutations in HINT1 gene cause the autosomal recessive axonal neuropathy with neuromyotonia. Eur J Med Genet. 2019;62(3):190–4.
12. Xu L, Wang G, Lv X, Zhang D, Yan C, Lin P. A novel mutation in HINT1 gene causes autosomal recessive axonal neuropathy with neuromyotonia, effective treatment with carbamazepine and review of the literature. Acta Neurol Belg. 2022;122:1305–12.
13. Amor-Barris S, Hoyer H, Brauteset LV, De Vriendt E, Strand L, Jordanova A, et al. HINT1 neuropathy in Norway: clinical, genetic and functional characterization. Orphanet J Rare Dis. 2021;16(1):116.
14. Lassuthova P, Brozikova DS, Krutova M, Neupauerova J, Haberlova J, Mazanec R, et al. Mutations in HINT1 gene are one of the most frequent causes of hereditary neuropathy among Czech patients and neuromyotonia is rather an underdiagnosed symptom. Neurogenetics. 2015;16(1):43–54.
15. Karczewski KJ, Francioli LC, Tao G, Cummings BB, Affoldi J, Wang Q, et al. Variation across 141,456 human exomes and genomes reveals the spectrum of loss-of-function intolerance across human genome sequences. Nature. 2010;467(7319):1061–73.
16. Cooper GM, Stone EA, Asimenos G, Program NCS, Green ED, Batzoglou S, et al. Distribution and intensity of constraint in mammalian genomic sequence. Genome Res. 2005;15(7):901–13.
17. Siavniene E, Petraitatie G, Bumyte B, Morkuene A, Miskiene V, Rancelis T, et al. Compound heterozygous c.998 +61del and c.1746+20C > G CAPN3 genotype cause autosomal recessive limb-girdle muscular dystrophy-1. A case report. BMCMusculoskeletDisord. 2021;12(1):1020.
18. Gotz RD, Schiestl RH. High-efficiency yeast transformation using the LiAc/SS carrier DNA/PEG method. Nat Protoc. 2007;2(1):31–4.
19. Zhang T, Lei J, Yang H, Xiu K, Wang R, Zhang Z. An improved method for whole protein extraction from yeast Saccharomyces cerevisiae: Yeast. 2011;28(11):795–8.

Publisher's Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.