

NATURAL HISTORY AND BIOMARKERS IN HEREDITARY SENSORY NEUROPATHY TYPE 1

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ABSTRACT: *Introduction:* Hereditary sensory and autonomic neuropathy type 1 (HSAN1) is most commonly caused by missense mutations in *SPTLC1*. In this study we mapped symptom progression and compared the utility of outcomes. *Methods:* We administered retrospective surveys of symptoms and analyzed results of nerve conduction, autonomic function testing (AFT), and PGP9.5-immunolabeled skin biopsies. *Results:* The first symptoms were universally sensory and occurred at a median age of 20 years (range 14–54 years). The onset of weakness, ulcers, pain, and balance problems followed sequentially. Skin biopsies revealed universally absent epidermal innervation at the distal leg with relative preservation in the thigh. Neurite density was highly correlated with total Charcot-Marie-Tooth Examination Score (CMTES; $r^2 = -0.8$) and median motor amplitude ($r^2 = -0.75$). *Conclusions:* These results confirm sensory loss as the initial symptom of HSAN1 and suggest that skin biopsy may be the most promising biomarker for future clinical trials.

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Small-fiber polyneuropathy (SFPN) is prominent in hereditary sensory and autonomic neuropathy type 1 (HSAN1) and contributes to many symptoms, including sensory loss, neuropathic pain, and tissue necrosis.^{1–3} Standard nerve conduction studies are reliable for assaying large-fiber function, but do not measure small-fiber function, as these fibers do not contribute to main components of compound motor action potentials and sensory nerve action potentials. Autopsy and surgical nerve biopsies from patients with HSAN1 have demonstrated marked cell loss in the dorsal root ganglia and sural

nerves.^{1–5} However, nerve biopsy is now considered too invasive for many research uses and is increasingly being supplanted by less-invasive methods. The best objective diagnostic tests for SFPN are distal-leg skin biopsy immunolabeled to reveal density of small-fiber epidermal innervation [Level C recommendation by the American Academy of Neurology (AAN), Level A recommendation by the European Federation of Neurological Societies],^{6,7} and autonomic function testing (AFT) of cardiovascular, adrenergic, and sudomotor small-fiber function (AAN Level B recommendation).⁶

The aims of our study were to characterize the natural history and disease progression of HSAN1 and to assess the utility of various outcome parameters to identify the best biomarkers for future clinical trials.

METHODS

Subject Recruitment. Patients with HSAN1 were recruited and enrolled between January 2011 and January 2012 at the Massachusetts General Hospital Neuromuscular Diagnostic Center and the University of Massachusetts Medical School. Inclusion criteria included age ≥ 18 years, ability to provide written informed consent, clinical diagnosis of HSAN1, and confirmation of a pathogenic mutation in the *SPTLC1* gene. Patients with other major risk factors for acquired neuropathy (e.g., diabetes) or severe medical comorbidities were excluded. All procedures and protocols had institutional review board approval, and subjects or relatives of deceased subjects provided written informed consent.

Phenotypic Characterization. Subject symptoms were assessed using a questionnaire created by HSAN1 patients plus a follow-up survey created by our group (refer to Supplementary Material, available online). The surveys were completed by affected patients or by relatives of deceased patients. In addition to demographic information, the age of onset and detailed characteristics of all sensory, autonomic, and motor symptoms were assessed. The presence, location, and time of onset of ulcers were documented, and respondents were asked to recall the timing of the onset of specific symptoms, including sensory loss, weakness in upper and lower extremities, onset of balance

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Abbreviations: AAN, American Academy of Neurology; AFT, autonomic function testing; CASS, Composite Autonomic Scoring Scale; CMAP, compound muscle action potential; HSAN1, hereditary sensory and autonomic neuropathy type 1; CMTES, Charcot-Marie-Tooth Examination Score; ENF, epidermal nerve fiber; IENFD, intraepidermal nerve fiber density; QSART, quantitative sudomotor axonal reflex test; SFPN, small-fiber polyneuropathy; SNAP, sensory nerve action potential

Key words: hereditary neuropathy; neurogenetics; peripheral neuropathy; skin biopsy; small fiber neuropathy

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difficulty, and onset of neuropathic pain. Information collected about effects on lifestyle included quality of sleep, pain levels, ability to perform activities of daily living, and need for walking aids. Tobacco and alcohol use were also assessed.

Prospective Assessment of Biomarkers. A subset of patients also underwent prospective neurologic examination, nerve conduction studies, autonomic function testing, and skin biopsies. Patients were scored using the Charcot-Marie-Tooth Examination Score (CMTES), a 12-point functional rating scale that incorporates both patient symptoms and examination findings.^{8,9} Skin biopsy samples were anonymized for interpretation.

Electrophysiologic Assessments. All nerve conduction studies were performed in a single clinical electrodiagnostic laboratory by a board-certified (both American Board of Electrodiagnostic Medicine and American Board of Psychiatry and Neurology) physician using an electromyography system (Synergy; Natus). Percutaneous stimulation and surface recordings using non-gelled, self-adhesive surface recording electrodes (Neuroline 725; Ambu) were used. Limb temperatures were maintained at around 32°C using heating packs.

Median sensory conduction studies were recorded orthodromically after ring electrode stimulation of the second digit at a 13-cm distance. Radial (8-cm), sural (14-cm), and superficial fibular (14-cm) sensory conduction studies were performed antidromically according to standard techniques. For all sensory conduction studies, peak latencies and peak-to-peak amplitudes were recorded.

Fibular motor conduction studies were performed with surface recordings obtained from the extensor digitorum brevis and tibialis anterior after stimulation of the fibular nerve at the ankle (9-cm extensor digitorum brevis study), below and above the fibular head. Tibial motor conduction studies were performed with recordings from the adductor hallucis with stimulation at the ankle (10 cm) and popliteal fossa. Median motor conduction studies were obtained with surface recordings from the abductor pollicis brevis after stimulation at the wrist (6 cm) and antecubital fossa. For the motor conduction studies, baseline-to-peak amplitudes, distal motor latencies, and conduction velocities were recorded.

Autonomic Assessments and Testing. A comprehensive battery of autonomic tests was used to evaluate the cardiovascular, adrenergic, and sudomotor autonomic domains.¹⁰ Standardized tests included tilt-table testing, a deep breathing test, and Valsalva maneuver. Deep breathing was performed at the rate of 6 breaths/minute for 1 minute. The

Valsalva maneuver was performed with an expiratory pressure of 40 mm Hg for 15 seconds. Participants were tilted for 10 minutes. The quantitative sudomotor axonal reflex test (QSART) was performed with a simulation current of 2 mA for 5 minutes. Blood pressure was monitored continuously (Finometer; Finapres, Amsterdam, The Netherlands; and Dinamap ProCare 10; GE, Fairfield, Connecticut). The validated Composite Autonomic Scoring Scale (CASS) was used to quantitate and assess AFT results.^{10,11} The CASS stratifies autonomic dysfunction as mild (total score 1–3), moderate (total score 4–6), or severe (total score 7–10). CASS scores were calculated automatically,¹⁰ and values from HSAN1 patients were compared with normative values of age-matched controls.¹⁰

Neurodiagnostic Skin Biopsies. Skin biopsies were performed by trained neurologists using local anesthesia and standard practices.^{6,7,12–14} Three-millimeter-diameter punch biopsies were performed in the distal calf (10 cm above the lateral malleolus) and in the upper thigh. The sites were treated with antibiotic ointment, and protective dressings were applied. All biopsies were processed and analyzed by the clinical diagnostic skin biopsy laboratory at Massachusetts General Hospital according to consensus standards.^{6,15} Free-floating 50- μ m vertical sections were labeled immunohistochemically against PGP9.5, a pan-neuronal marker (Chemicon, Temecula, California) to reveal epidermal nerve fibers (ENF) and to permit standard nerve density. Almost all PGP9.5-immunoreactive epidermal neurites are nociceptive small-fiber endings, and axonal localization of epidermal PGP9.5 immunolabeling has been verified ultrastructurally. One skilled morphometrist, blinded to group allocation, measured ENF density. Our laboratory reports ENF densities per square millimeter of skin surface area to control for varying skin section thickness between laboratories.

Data Analysis. Demographic data and the frequency of each specific symptom and age of onset of that symptom were extracted and analyzed. Summary statistics included group median \pm standard error. To assess objective outcome data, patients were rank ordered according to severity for each individual outcome measure, and the ranks were compared. CMTES scores were correlated with electrophysiologic and skin biopsy results. The prevalence of abnormal results was compared between groups by chi-square analysis.

RESULTS

Phenotypic Characterization. Twenty-three patients with HSAN1 (10 men, 13 women) completed the HSAN1 survey (Table 1). Twenty were from an

Table 1. Characteristics of patients with HSAN1 ($n = 23$).

Demographic characteristics	Number (proportion) or median (range)
Gender	
Men	10 (43%)
Women	13 (57%)
Age (years)	52.2 (range 28–79)
Mutation	
C133Y	20 (87%)
C133W	3 (13%)
Symptoms	
Loss of sensation ($n = 23$)	23 (100%)
Weakness ($n = 23$)	21 (91%)
Balance abnormality ($n = 23$)	21 (91%)
Shooting pain ($n = 23$)	23 (100%)
Ulcers ($n = 23$)	17 (74%)
Amputations ($n = 23$)	8 (35%)
Changes in sweating ($n = 23$)	8 (35%)
Orthostasis ($n = 23$)	3 (13%)
Symptom age of onset (years)	
First symptom ($n = 23$)	20 (14–54)
Sensory loss feet ($n = 23$)	20 (14–54)
Sensory loss hands ($n = 15$)	32 (22–59)
Weakness feet ($n = 18$)	27 (20–60)
Weakness hands ($n = 15$)	39 (25–70)
First ulcer ($n = 17$)	29 (15–70)
Shooting pain ($n = 19$)	30 (15–70)
Balance difficulty ($n = 19$)	30 (21–70)

Symptom onset in men vs. women

	Men (mean age, in years)	Women (mean age, in years)	
First symptom	21.2 ($n = 10$, SD = 9.6)	29.5 ($n = 13$, SD = 14.9)	$P = 0.0410$
Onset weakness	30.3 ($n = 6$, SD = 10.4)	35.4 ($n = 12$, SD = 15.4)	$P = 0.4777$
First ulcer	27.2 ($n = 9$, SD = 12)	39 ($n = 8$, SD = 16.2)	$P = 0.1059$
Balance difficulty	31.9 ($n = 7$, SD = 11.8)	39.9 ($n = 12$, SD = 16.4)	$P = 0.2758$

HSAN1, hereditary sensory autonomic neuropathy type 1; SD, standard deviation.

extended family with the C133Y mutation in *SPTLC1*, and 3 were from a family with the C133W mutation. The median age at first symptom onset was 20 years (range 14–54 years). However, many patients noted that their symptoms probably began in the early teens but were unappreciated at the time. The mean age at symptom onset was less in men than it was in women (21 years vs. 30 years, $P = 0.04$).

Sensory problems were universal and appeared first. Twenty of 23 (87%) subjects reported reduced sensation as their initial symptom, and the others reported positive sensory phenomena (paresthesias or foot discomfort) as their initial symptom. Among those with sensory loss, all described a stocking-glove distribution, and more reported impaired temperature perception than light touch or pressure. One patient (patient 18) reported profound sensory loss that spared only the face and parts of the back. See Table 2 for selected detailed patient responses. Of the patients who reported reduced sensation in both the hands and feet, the mean latency between onset of foot and hand symptoms was 10 years (SD = 7.4, range 3–27 years).

The onset of weakness, ulcers, pain, and balance problems followed sensory problems sequentially on average 7.6, 8.1, 9.5, and 11.4 years later, respectively (Table 1 and Fig. 1). With regard to painful symptoms, 10 of 23 (43%) patients reported tingling, and 5 of 23 (22%) reported burning. Patients reported using multiple pain medications, including gabapentin, venlafaxine, amitriptyline, sertraline, oxcarbazepine, pramipexole, duloxetine, acetylsalicylic acid, acetaminophen, meperidine, and oxycodone. Two patients reported that gabapentin and venlafaxine improved their symptoms significantly, but many patients reported that their pain was refractory to all medications.

Seventeen of 23 patients (74%) reported ulcers. These developed at an average of 8.1 years after initial neuropathic symptoms ($n = 17$, range 0–20 years; Table 1). The ulcers were restricted to the lower extremities in 17 patients and affected both the upper and lower extremities in 4 patients. Approximately one-third of patients required distal limb amputations. There was a trend toward earlier ulceration in men (mean age of 27 vs. 40 years, $P = 0.28$).

Table 2. Patient-reported outcomes in HSN1 (*n* = 23).

Gender	Age (years)	Mutation	Sensory loss in feet	Sensory loss in hands	Weakness of feet	Weakness of hands	Ulcers	Amputations	Balance difficulty	Shooting pain
M	38	C133Y	18	32	27	36	N	N	26	25
M	38	C133Y	14	31	32	32	29	Y	32	28
W	32	C133Y	20	25	29	N	N	N	24	30
M	59	C133Y	22	25	26	29	23	N	28	30
W	61	C133Y	45	?	45	55	N	N	45	46
W	82	C133Y	54	59	54	59	54	Y	60	70
W	56	C133Y	14	36	25	46	30	Y	40	34
M	63	C133Y	47	55	50	58	57	N	58	49
W	85	C133Y	50	Y (? age)	Y (? age)	Y (? age)	70	Y	50	Y (? age)
W	61	C133Y	28	39	50	55	44	N	54	42
W	63	C133Y	20	25	20	39	34	N	40	30
M	35	C133Y	18	Y (? age)	N	N	21	N	23	22
W	34	C133Y	16	Y (? age)	20	Y (? age)	20	N	22	21
W	40	C133Y	21	32	23	34	N	N	23	15
W	28	C133Y	20	24	22	25	N	N	21	22
M	?	C133Y	25	Y (? age)	N	N	26	Y	?	Y (? age)
M	D	C133Y	17	Y (? age)	Y (? age)	Y (? age)	28	Y	?	Y (? age)
M	D	C133Y	15	Y (? age)	Y (? age)	Y (? age)	15	Y	Y (? age)	Y (? age)
W	79	C133Y	23	50	60	70	30	Y	70	45
M	62	C133Y	19	25	20	25	26	Y	28	45
M	33	C133W	19	22	27	29	20	Y	28	28
W	36	C133W	23	N	25	N	30	N	30	32
W	59	C133W	52	55	52	55	N	N	N	55

Numbers refer to the age of onset of various symptoms. D, deceased; W, woman; M, man; N, no; Y, yes.

Concerning autonomic symptoms, 8 patients (35%) reported altered sweating, and 3 (13%) had light-headedness. Two reported low blood pressure and light-headedness starting in the fifth decade, whereas 1 recalled these symptoms in the teenage years. Three patients reported early satiety developing after age 50 years. For motor symptoms, the mean latency between awareness of the first sensory symptoms and awareness of weakness averaged 8.1 years (range 0–38 years). Foot weakness came first, with hand weakness following, on average, 7.6 years later (SD = 6.0, range 0–21 years). Difficulty with balance was noticed, on average, 11 years after initial symptoms (SD = 11.9, range 0–48 years). Nine patients (39%) reported sleep difficulty, largely related to discomfort.

Concerning activities of daily living, 85% of patients required help with dressing, beginning at a median age of 40 years (range 26–70 years). Several specifically reported difficulty buttoning clothes. Regarding ambulation, 12 of 20 (60%) required ankle orthotics, beginning at a median age of 32.5 years (range 24–60 years). Half of the patients used a cane, beginning at a median age of 40 years (range 34–70 years). Only 3 required a walker at a median age of first use at 56 years (range 55–75 years). Five of 20 patients (25%) used a wheelchair, starting at a median age of 49 years (range 40–78 years), but usually only for long distances. Several patients noted that, in retro-

spect, they would have benefited from walking aids long before they began using them. Eight of 23 patients (35%) reported ever smoking tobacco. Nine reported alcohol use, and 4 reported periods of heavy consumption. One patient reported aggravation of symptoms during each of her 2 pregnancies. Another patient reported worsened symptoms in the setting of sleep deprivation and after eating.

Outcome Measures. All procedures were well-tolerated and without adverse effects. Eight patients (5 with the C133Y gene mutation and 3 with the C133W mutation) were examined using the CMTES and underwent nerve conduction studies (Table 3)

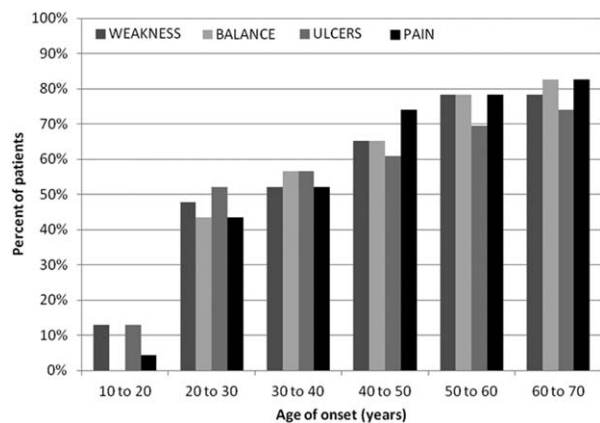


FIGURE 1. Cumulative incidence of symptoms in HSN1 (*n* = 23).

Table 3. Nerve conduction studies and skin biopsies (IENFD) (*n* = 8).

Age/Sex	57/F	35/F	30/F	59/F	32/M	26/F	33/F	55/M
Dorsum Foot	NA	NA	0	0	0	0	0	NA
Distal Leg	NA	NA	0	0	0	0	0	0
Thigh	NA	NA	249	88	45	17	15	12
IENFD Rank	NA	NA	3	4	5	6	7	8
Motor	Med (7.8),	Med (12.6),	Med (6.8),	Med (5.5),	Med (4.6),	Med (3.5),	Med (0.1),	Absent
CMAPS	F (2.4),	F (0.1),	Absent	Absent	Absent	Absent	Absent	(Med/F)
(mv)	T (0.3)	T(0.3)	(F/T)	(F/T)	(F/T)	(F/T)	(F/T)	
Sensory	Med (17.1),	Med (2.4),	R (3.8),	Absent	Absent	Absent	Absent	Absent
NCS	R (25.2),	R (6.4),	Absent	(Med/R/	(Med/R/	(Med/R/	(Med/R/	(Med/R/
(μv)	S (2.9),	Absent	(Med/S/SF)	S/SF)	S/SF)	S/SF)	S/SF)	S/SF)
	Absent	(S/SF)						
	(SF)							
Motor	Med-APB	Med-APB	Med-APB	Med-APB	Med-APB	Med-APB	NA	NA
CV (m/s)	(58.3),	(50)	(49.4)	(48.8)	(41.7)	(30.1)		
	P-EDB							
	(49)							
NCS Rank	1	2	3	4	5	6	7	8
CMTES	2	7	7	16	16	12	16	23

APB, abductor pollicis brevis muscle; EDB, extensor digitorum brevis muscle; F, female; IENFD, intraepidermal nerve fiber density; M, male; median, median nerve; NA, not applicable; NCS, nerve conduction studies; F, fibular nerve; R, radial nerve; S, sural nerve; SF, superficial fibular nerve; T, tibial nerve.

at a median age of 34 years (range 26–59 years). The median age of symptom onset was 19 years (range 15–54 years). Median CMTES score was 14 (range 2–23). Electrophysiological studies revealed inability to elicit any sensory nerve action potentials (SNAPs) in the arms or legs in 5 of 8 patients. Lower extremity SNAPs were absent in 7 of 8 patients and were abnormal in the eighth individual; this least-affected subject had an absent superficial fibular SNAP and a low-amplitude sural SNAP. Compound muscle action potentials (CMAPs) in the lower extremities were absent in 6 of 8 patients (Table 3). Among patients with measurable CMAPs, 4 had normal conduction velocities. Two had slowing of the median motor conduction velocities (41.7 and 30.1 m/s) without clear prolongation of the median distal motor latencies to suggest entrapment at the wrist. In aggregate, nerve conduction studies confirmed the presence of significant large-fiber involvement of both sensory and motor fibers.

None of the 6 patients who underwent skin biopsies of the distal leg and thigh had any epidermal innervation remaining at the distal leg site, whereas some epidermal innervation always remained at the thigh site, consistent with the known length-dependent pattern of the neuropathy (Table 3 and Fig. 2). The correlation coefficient between the CMTES and the density of epidermal innervation was -0.8 and between CMTES and median CMAP amplitudes was -0.75 . Six patients underwent autonomic function testing (Table 4) on 2 occasions at 1-year intervals. Total CASS scores averaged 1.5 at the first visit, indicating mild autonomic dysfunction. No significant changes were detected 1 year

later, with scores averaging 1.2. Only 1 patient had a moderate CASS score of 6. This man, who was 55 years old at study onset, also had the highest CMTES, the most severe neuropathy on nerve conduction studies (absent SNAPs and CMAPs throughout), and the least remaining epidermal innervation at the thigh site.

DISCUSSION

This study was a comprehensive characterization of HSAN1-associated symptoms with a focus on the timing of onset of specific symptoms. Symptom onset in our patients was usually in adolescence or early adulthood, and men were more severely affected than women. This study demonstrated that HSAN1 is a misnomer, at least for the most common *SPTLC1* mutations. Motor axonopathy was severe as assessed by symptoms, signs, and electrophysiological study, whereas autonomic neuropathy was scant. We also performed and analyzed potential objective neurophysiologic and pathologic outcome markers in a subset of patients, and we conclude that IENFD in the thigh is the most promising among them for research use.

The phenotype of our patient population was similar to that described by Houlden *et al.* in 6 families with the C133W mutation in *SPTLC1*.² Much like their patients, many of ours developed early weakness and required walking aids. Motor involvement is thus an important feature of both the C133Y and C133W mutations. Other *SPTLC1* mutations, such as the S331F and the more recently described S331Y mutation, can cause

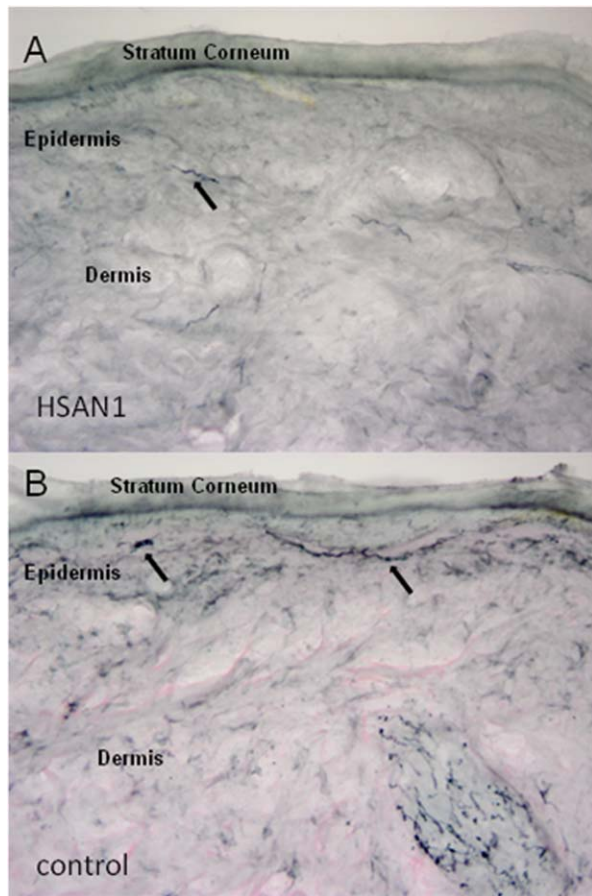


FIGURE 2. Skin biopsy section from the thigh showing loss of intraepidermal nerve fibers (black arrows) in a representative HSAN1 patient (A) as compared with an age-matched control (B) with normal distribution of dermal and intraepidermal nerve fibers.

severe weakness and hypotonia plus additional features not reported by our patients, including scoliosis, respiratory failure, and early cataracts.¹⁶

Because there is increasing awareness of the need to include patient-defined outcomes in clinical research, we included a questionnaire developed by patients and collected information on the impact of HSAN1 on lifestyle and patient-centered

outcome measures. The questionnaire included detailed questions regarding the physical symptoms of HSAN1, as well as quality-of-life measures such as sleep quality, stress levels, alcohol use, ability to perform activities of daily living, and the impact of pain. The findings highlight a relatively predictable sequence of symptom onset in HSAN1. Many patients reported subtle sensory symptoms in early- to mid-teenage years that did not rise to physician awareness. Sensory loss was universally the first physician-recorded symptom, followed by weakness, with neuropathic pain and skin ulcers developing, most often in the second and third decades of life. Balance difficulty was common but absent until the third decade and became gradually more common over time. Thus, as is true of many hereditary neuropathies, HSAN1 is actually a pediatric-onset disease, and the optimal timing for therapeutic intervention is likely to be in childhood.

Another finding was great phenotypic variability within both families, most evident as a gender dichotomy and in the timing and severity of weakness and the timing of upper limb involvement. Although most patients noted weakness within 5–10 years after onset, 1 reported a 38-year lag; this is consistent with earlier reports of significant intrafamilial variability in disease severity.^{1,2} It is unknown whether this reflects environmental modifiers or underlying pathophysiologic variability, but it should be considered in designing measures to analyze progression of HSAN1 and suggests that assessing responses to disease-modifying therapy will require individual rather than standardized baselines. In addition, careful assessment of lifestyle information will be important to detect environmental or hormonal modifiers. The repercussions of such chronic disability on mood, daytime function, and overall quality of life have not been well characterized, and it will be necessary to develop outcome measures that can assess the full impact of potential new therapies.

Table 4. Autonomic function testing at baseline and 1 year ($n = 6$).

M/W	Age	Baseline				1 year				
		CMTES	CV	ADR	SUD	CASS	CV	ADR	SUD	CASS
W	30	7	0	0	0	0	0	0	0	0
W	59	16	0	1	0	1	0	1	1	2
M	32	16	0	0	0	0	0	0	0	0
W	26	12	0	0	1	1	1	0	0	1
W	33	16	0	0	1	1	1	0	0	1
M	55	23	3	3	6	6	0	3	1	4

ADR, adrenergic score; W, woman; CV, cardiovascular score; SUD, sudomotor score; CASS, Composite Autonomic Severity Score. Key: 1—Cardiovascular score: based on deep breathing and Valsalva ratios. 2—ADR score: based on Valsalva and tilt testing. 3—SUD score: based on quantitative sudomotor axon test (QSART). 4—CASS (total score). 0 = normal; 13 = mild generalized autonomic failure; 46 = moderate generalized autonomic failure; 710 = severe generalized autonomic failure.

We evaluated skin biopsies, nerve conduction studies, and autonomic testing as potential outcome measures in a small number of patients. The composite nerve conduction study data correlated well with clinical severity. However, the marked reduction (or absence) of recordable SNAPs and CMAPs using standard nerve conduction study techniques limits the utility of NCS for monitoring disease progression. In contrast, the scarcity of autonomic symptoms and mild objective dysfunction without progression over a 1-year period^{2,3,17} suggest that autonomic testing may not suitably monitor disease severity either. Houlden *et al.* also observed relatively few autonomic abnormalities in HSN1.² The skin biopsy findings, which have not been described previously in HSN1, demonstrated universal epidermal denervation at the distal-leg site in adults. Even patients with severe disease had some remaining innervation at the thigh, and these densities correlated well with both nerve conduction studies and CMTES scores. This suggests that skin biopsies from the thigh may be a useful measure of disease severity and possibly progression in HSN1.

This study has several limitations in addition to its small sample size. Importantly, the retrospective survey allows for recall bias concerning the timing and nature of symptoms. Patient identification of new symptoms is affected by the timing of diagnosis, because patients with a diagnosis and receiving care in a Neurology clinic are more likely to become aware of new symptoms. The timing of skin ulcers, which varied greatly, may also reflect the timing of diagnosis and initiation of appropriate foot care. It is also noteworthy that we did not use the complete CMTNS, which is a combined scoring system incorporating neuropathy symptoms, signs, and electrophysiologic measurements. Although a subset of patients did undergo electrophysiologic evaluation, the specific measures required for calculating the CMTNS (radial SNAP and ulnar CMAP) were not obtained in most patients.

In conclusion, we have characterized HSN1 as a debilitating sensorimotor neurologic disease with broad impact on activities of daily living and quality of life. We reported patient-centered characterization of frequency and relative timing of the key clinical features of this disease. In addition, this study highlights that, among objective disease markers, measurement of epidermal innervation in

skin biopsies from the lower limb may be the most promising outcome measure for use in clinical trials.

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