Charcot-Marie-Tooth neuropathy score and ambulation index are both predictors of orthotic need for patients with CMT

Valeria Prada1,2 · Riccardo Zuccarino3 · Cristina Schenone1 · Giulia Mennella1 · Marina Grandis1,4 · Michael E. Shy2 · Angelo Schenone1,4

Received: 22 July 2021 / Accepted: 29 September 2021 / Published online: 6 October 2021 © The Author(s) 2021

Abstract
Charcot-Marie-Tooth (CMT) disease is the most common hereditary neuropathy with an estimated prevalence of 1 person affected on 2500. Frequent symptoms include distal weakness and muscle wasting, sensory loss, reduced deep tendon reflexes, and skeletal deformities, such as hammer toes and pes cavus. CMT is a progressive disease and patients’ needs change over their lifetime. In particular, ambulation aids are increasingly needed to maintain ambulation and reduce the risk of falls. We performed a retrospective analysis of medical records from 149 patients with confirmed CMT to evaluate patients ambulation needs related to the severity of their CMT as measured by the CMT Neuropathy Score (CMTNS) and Ambulation Index (AI). Most patients required some form of orthotics (86.6%). The CMTNS and AI scores both differed significantly between patients with no orthotics compared to those who wore insoles/inserts. The CMTNS and AI also differed significantly between patients wearing insoles and those with ankle foot orthotics (AFOs). CMTNS and the AI were valid predictors of the type and choice of the orthotics. Both the CMTNS and AI can be effective tools to aid in the correct choice of orthotics in patients affected by CMT.

Keywords Charcot-Marie-Tooth disease · CMT · Predictor · Ambulation · Braces · Orthotics

Introduction
Charcot-Marie-Tooth (CMT) disease is the most common hereditary neuropathy with an estimated prevalence of 1 in 2500 [1, 2]. CMT is caused by mutations in more than 80 different genes [3–5]. Despite the genetic heterogeneity, most forms have common symptoms though the rates of progression may differ [6]. These symptoms include distal weakness and muscle wasting, distal loss of proprioception, sensory loss, reduced deep tendon reflexes, and skeletal deformities, such as hammer toes and pes cavus [7]. Taken together, these symptoms usually result in an abnormal gait, foot drop, and difficulties with balance in patients [8, 9]. Progression of these symptoms is variable ranging from slow [10] to more rapid depending on the specific genetic cause and mutation [5].

However, in all these cases, patients need to have their gait analyzed over a lifetime because the need and type of correction will change as disability increases [11]. Moreover, foot deformities and surgeries are variables that can influence the needs of orthotics over time [12–14].

Patients with CMT typically use different kinds of orthotics devices, ranging from shoe inserts to ankle foot orthoses (AFOs) [15]. We separate AFOs into two types. The first are the low AFOs that extend just above the ankle and provide stability only at the ankle, without using the tibia bone as a lever. The second type are the AFOs which extend higher up the calf and use the tibia as a lever (like leaf spring or toe off orthoses) [16, 17]. Currently, we are unaware of consensus or guidelines as when to use any of these devices. We hypothesize that having more complete information of the relationship between disease severity and the type of...
orthotic used would provide useful information for health care providers to help provide the best choice in what they prescribe for patients [16].

To begin addressing this issue, we performed a retrospective analysis of a cohort of 149 patients with CMT followed in the outpatient unit of the Neurological Clinic of the San Martino Hospital (Italy). The aim of the study was to identify correlations between the use of different insoles/orthoses and clinical outcome assessments (COA) of CMT so that we could develop preliminary algorithms that would be useful for managing the ambulation needs of our patients.

Methods

A retrospective analysis was carried out on medical records maintained in the Inherited Neuropathy Facility of the Ospedale Policlinico San Martino IRCCS of Genova. Data was recorded anonymously and placed in a file excel, specifying the following: sex, date of birth, genetical diagnosis, date of the visit, type of orthosis, Ambulation Index (AI), and CMT Neuropathy Score version 2 (CMTNSv2) [18, 19]. We also recorded data for the AI, based on the time and needs of patients to walk 25 feet [20].

A total of 221 charts of confirmed diagnosis of CMT were examined. We excluded 72 patients because they lacked a CMTNSv2 or specifications about insoles or orthoses. We therefore included in the study all the patients with (1) a specified and genetical diagnosis of CMT; (2) a CMTNSv2 score; (3) an AI score; and (4) a prescription of an insole or orthoses. All data came from the same visit.

Statistical analyses

An ANOVA test with Tukey’s multiple comparisons test was made for comparison of different classes. A Spearman r test was used for correlation between scales. Analysis of the difference between two groups was carried out with a Mann–Whitney t test. Multiple comparisons were analyzed with the Tukey multiple comparisons test. Significance was considered for \( p < 0.05 \).

Results

One hundred forty-nine patients (74 males and 75 females) were included in the study and met all the inclusion criteria. The mean age was 53.2 (±16.7) with an age range between 18 and 89 years. Sixty-eight patients (45.6%) had CMT1A, 6 (4.0%) patients had CMT1B, 24 patients (16.1%) had CMT2 subtypes, and 3 (2.0%) had recessive CMT4 forms. Thirty patients (20.0%) were diagnosed with CMTX (29 CMTX1 and 1 CMTX5) and the remaining 18 patients (12.1%) had HNPP.

Most of the patients required some form of foot or ankle support; only 13.4% (\( n = 20 \)) were not wearing any type of orthotic. Forty-eight percent (47.7; \( n = 71 \)) wore just inserts or insoles. Low AFOs were used by 12.8% (\( n = 19 \)), and higher AFOs by 17.4% (\( n = 26 \)). Seven patients (4.7%) also required a cane to walk and 6 patients (4.0%) needed a walker or a wheelchair for ambulation (Table 1). The mean and the range of the scores from the CMTNS and ambulation are shown in Table 2 for the different CMT subtypes. We identified strong correlations between the CMTNS and AI scales (\( r = 0.75; r^2 = 0.49; p < 0.0001; \) Supplementary Fig. 1).

We next compared the type of orthoses used with the CMTNSv2 and found that CMTNS scores were significantly different between patients wearing no orthotics compared to those with insoles alone (patients without orthotics: \( 2.5 \pm 2.6 \) CMTNS; patients with insoles: \( 9.3 \pm 4.1 \) CMTNS; Table 1).
There were also significant differences between patients wearing insoles compared to those wearing low and high AFOs (patients with low AFO: 15.2 ± 4.9 CMTNS; high AFO: 17.1 ± 5.4; \(p < 0.0001\) for both). Although there were slight differences between the group wearing low and high AFOs, these differences did not reach significance (\(p = 0.45\); Fig. 1A).

Similar results were obtained with correlations to the AI (Fig. 1B). The mean AI for the three groups was as follows: no orthotics worn 0.05 ± 0.2; insoles 0.93 ± 1.0; low AFO 2.2 ± 0.8; AFO 2.6 ± 1.1. There is a significant difference between the AI of people who do not wear orthotics and people who use insoles (\(p = 0.01\)), between people who wear insoles and people who wear low AFO (\(p < 0.0001\)), and between people who use insoles and people who wear AFO (\(p < 0.0001\)). No differences have been found in the group who wears the low AFO and the one who uses AFO (\(p = 0.27\)).

We then evaluated patients who were using canes, walkers, or wheelchairs, though these numbers were small. Patients who needed a cane for walking had significantly lower CMTNS compared to those who needed a walker or a wheelchair (one cane: 13.1 ± 4.1; chair or walker 24.6 ± 1.5; \(p = 0.001\); Fig. 2).

Finally, we studied the distribution of the different orthoses between the different CMT subtypes. In CMT1A and CMT1B, the most frequently used orthotics were insoles (CMT1A 70.6%, \(n = 48\); CMT1B 50.0%, \(n = 3\)). Patients with CMT2 had a different distribution: 29.2% (\(n = 7\)) wore insoles, 20.8% (\(n = 5\)) wore low AFO, 29.2% (\(n = 7\)) wore high AFOs, 16.7% (\(n = 4\)) used one cane, and only 4.2% (\(n = 1\)) did not use orthoses. The majority of our CMT4 patients used wheelchairs or walkers (66.7%, \(n = 2\)), and only 1 patient wore an AFO (33.33%) and used insoles. 11.1% (\(n = 2\)) needed AFOs and 1 patient (5.6%) walked with one cane (Supplementary Table 1; Fig. 3A).

### Table 2 Ambulation Index and CMTNS of the different clusters of patients

| Cluster   | Mean age | Range age | Mean Ambulation Index | Range Ambulation Index | Mean CMTNS | Range CMTNS |
|-----------|----------|-----------|-----------------------|------------------------|------------|-------------|
| Overall   | 53.2 ± 16.7 | 18–89     | 1.5 ± 1.5             | 0–7                    | 11.3 ± 6.7 | 1–28        |
| CMT1A     | 51.0 ± 17.4 | 18–89     | 1.3 ± 1.4             | 0–5                    | 11.4 ± 5.8 | 1–27        |
| CMT1B     | 49.3 ± 15.1 | 34–73     | 2.2 ± 2.5             | 0–7                    | 16.2 ± 5.2 | 10–25       |
| CMTX      | 52.1 ± 16.5 | 24–84     | 1.9 ± 1.5             | 0–6                    | 13.2 ± 6.6 | 3–25        |
| CMT2      | 66.3 ± 12.9 | 34–81     | 1.9 ± 1.2             | 0–4                    | 11.4 ± 4.8 | 3–21        |
| CMT4      | 40.7 ± 17.6 | 21–55     | 4.7 ± 1.5             | 3–6                    | 24.3 ± 3.5 | 21–28       |
| HNPP      | 49.7 ± 11.8 | 31–73     | 0.5 ± 1.0             | 0–3                    | 4.3 ± 6.7  | 0–26        |
We paid particular attention to the patients with CMTX (29 CMTX1 and 1 CMTX5) because we had a large group of these patients and because males are often more severely affected than females. Two patients, both females, did not wear any orthotic (6.7%). Nine patients with CMTX (30.0%) used inserts (6 males, 3 females), low AFO were worn by 8 (26.7%) patients (7 males, 1 female), and high AFOs were used by 9 patients (30.0%, 7 males, 2 females). One with CMTX (3.3%) used a cane to walk and an additional patient used a wheelchair. Both of these patients were male (Fig. 3B).

Discussion

We have retrospectively evaluated orthotic use in 149 patient subjects with different confirmed types of CMT. Consistent with the literature, our largest subgroup had CMT1A [21, 22]. Both genders were equally represented. Most of our patients (86.6%) required orthotics of some type highlighting the importance of inserts or AFOs in managing CMT. This is in keeping with our concept that the use of orthotics and braces is important for CMT patients because it facilitates a better quality of gait and balance [23, 24]. In addition, the choice of an appropriate orthotic is important in progressive disorders like CMT in which ambulation needs change with disease impairment, as well as with alterations in foot structure and foot and ankle surgeries. A poor choice in the type of orthotics/orthoses used will likely result in exacerbating difficulties in ambulation [14], decreased likelihood that patients will wear the orthotic, and increased risk in tripping and falling during daily activities [8].

The literature suggests that most patients are satisfied overall with the orthotics they are wearing [17]. However, even in this group there were often concerns from patients about how their orthotics could be improved [17], and this is important because these devices are fundamental for many activities of daily living [25]. Specifically, we found that correlations with the CMTNS and AI accurately both predicted the type of orthotic worn by patients, consistent with prior studies that the CMTNS and AI correlate with each other [18]. The correlations with the CMTNS and AI
are also consistent with what was predicted in the original CMTN5 report in which neuropathies were characterized as mild (CMTNS ≤ 10), moderate (CMTNS 11–20), and severe (≥ 21) [18]. We found in our study people wearing no orthotics typically fell into the mild neuropathy scoring. Among people who used orthotics, 47.7% used only insoles and 30.2% utilized low or high AFOs. Those who had CMTNS between 10 and 15 usually wore only insoles, those with scores up to 20 most often wore low AFOs, and those with scores above 20 either wore high AFOs, like the leaf spring, or rarely needed walkers or wheelchairs. We recognize that the “motor symptoms legs” component of the CMTN5v2 includes orthotics as part of the scoring [19]. However, this represents just one of the nine items in the CMTNS so we believe this one item cannot be responsible for all the correlation with orthotic use and the CMTN5v2. The AI also predicted well the type of orthotic worn by patients as correlations with these scores and the different orthotics were also significant, even, in this case, the results could strongly depend on the fact that, in this scale, orthotics are one on the criteria to consider in the scoring.

We were not able to predict the type of orthotic used based on CMT subtype, probably because there was variability in severity of neuropathy within the different subtypes including CMT1A, CMT1B, CMTX, and CMT2. The one potential exception was in patients with recessive forms of CMT such as CMT4, who were usually more severely affected and were likely to use high AFOs or even walkers and wheelchairs.

We did pay special attention to differences between men and women with CMTX (mostly of them affected by CMTX1), since women are often more mildly affected than men due to the presence of a normal X-chromosome (mostly affected by CMTX1), since women are often more mildly affected than men due to the presence of a normal X-chromosome [26]. In agreement, men with CMTX were more likely to use high AFOs or even a cane/walker compared to women.

In conclusion, we find that the choice of the correct AFO for patients really depends on the severity of their neuropathy, which can be predicted by their CMTNS or AI. We recognize potential contributions to orthotic needs by anatomical foot structure and/or previous surgical interventions. However, our data demonstrate that the CMTNS and AI are both good predictors of orthotic needs for patients with CMT.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s10072-021-05646-9.

Data availability The datasets generated and analyzed during the current study are not publicized but are available from the corresponding author on reasonable request.

Declarations

Ethical approval The study was approved by the Local Ethics Committee (approval number #4/2012).

Consent to participate Written informed consent was obtained from each participant.

Consent for publication Not applicable.

Conflict of interest None.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

References

1. Barreto LCLS, Oliveira FS, Nunes PS et al (2016) Epidemiologic study of Charcot-Marie-Tooth disease: a systematic review. Neuroepidemiology 46:157–165
2. Grandis M, Shy ME (2005) Current therapy for Charcot-Marie-Tooth disease. Curr Treat Options Neurol 7:23–31
3. Braatnen GJ (2012) Genetic epidemiology of Charcot-Marie-Tooth disease. Acta Neurol Scand. https://doi.org/10.1111/ane.12013
4. Frasquet M, Rojas-García R, Argente-Escrig H et al (2020) Distal hereditary motor neuropathies: mutation spectrum and genotype-phenotype correlation. Eur J Neurol. https://doi.org/10.1111/ene.14700
5. Fridman V, Reilly MM (2015) Inherited neuropathies. Semin Neurol 35(4):407–423. https://doi.org/10.1055/s-0035-1558981
6. Reilly MM, Murphy M, Laura M (2011) 2009 Peripheral nerve society meeting Charcot-Marie-Tooth disease. J Peripher Nerv Syst 14:1–14
7. Pareyson D, Marchesi C (2009) Diagnosis, natural history, and management of Charcot-Marie-Tooth disease. Lancet Neurol 8(7):654–667. https://doi.org/10.1016/S1474-4422(09)70110-3
8. Ong CF, Geijtenbeek T, Hicks JL, Delp SL (2019) Predicting gait adaptations due to ankle plantar flexor muscle weakness and contracture using physics-based musculoskeletal simulations. PLoS Comput Biol. https://doi.org/10.1371/journal.pcbi.1006993
9. Sackley C, Disler PB, Turner-Stokes L et al (2009) Rehabilitation interventions for foot drop in neuromuscular disease. Cochrane Database Syst Rev (3):CD003908. https://doi.org/10.1002/14651858.CD003908.pub3
10. Rossor AM, Shy ME, Reilly MM (2020) Are we prepared for clinical trials in Charcot-Marie-Tooth disease? Brain Res 1729:146625. https://doi.org/10.1016/j.brainres.2019.146625

11. Ramdharry G (2018) Peripheral nerve disease. In: Handbook of clinical neurology. 2018;159:403–415. https://doi.org/10.1016/B978-0-444-63916-5.00026-4

12. Laurá M, Singh D, Ramdharry G et al (2018) Prevalence and orthopedic management of foot and ankle deformities in Charcot–Marie–Tooth disease. Muscle Nerve 57:255–259. https://doi.org/10.1002/mus.25724

13. Crosbie J, Burns J (2007) Predicting outcomes in the orthotic management of painful, idiopathic pes cavus. Clin J Sport Med 17:337–342. https://doi.org/10.1097/JSM.0b013e31814c3e9e

14. Waterval NFJ, Nollet F, Harlaar J, Brehm MA (2017) Precision orthotics: optimising ankle foot orthoses to improve gait in patients with neuromuscular diseases; Protocol of the PROOF-AFO study, a prospective intervention study. BMJ Open. https://doi.org/10.1136/bmjopen-2016-013342

15. Ramdharry GM, Pollard A, Anderson C et al (2014) A pilot study of proximal strength training in Charcot-Marie-Tooth disease. J Peripher Nerv Syst 19:328–332. https://doi.org/10.1111/jnns.12100

16. Vinci P, Gargiulo P (2008) Poor compliance with ankle-foot orthoses in Charcot-Marie-Tooth disease. Eur J Phys Rehabil Med 44:27–31

17. Zuccarino R, Anderson KM, Shy ME, Wilken JM (2021) Satisfaction with ankle foot orthoses in individuals with Charcot-Marie-Tooth disease. Muscle Nerve. https://doi.org/10.1002/mus.27027

18. Shy ME, Blake J, Krajewski K et al (2005) Reliability and validity of the CMT neuropathy score as a measure of disability. Neurology. https://doi.org/10.1212/01.WNL.0000156517.00615.A3

19. Murphy SM, Herrmann DN, McDermott MP et al (2011) Reliability of the CMT neuropathy score (second version) in Charcot-Marie-Tooth disease. J Peripher Nerv Syst. https://doi.org/10.1111/j.1529-8027.2011.00350.x

20. Hauser SL, Dawson DM, Lehrich JR et al (1983) Intensive immunosuppression in progressive multiple sclerosis. A randomized, three- arm study of high-dose intravenous cyclophosphamide, plasma exchange, and ACTH. N Engl J Med 308:173–180

21. Liu P, Gelowani V, Zhang F et al (2014) Mechanism, prevalence, and more severe neuropathy phenotype of the Charcot-Marie-Tooth type 1A triplication. Am J Hum Genet. https://doi.org/10.1016/j.ajhg.2014.01.017

22. Shy ME (2004) Charcot-Marie-Tooth disease: an update. Curr Opin Neurol 17(5):579–585. https://doi.org/10.1097/00019052-200410000-00008

23. Sautreuil P, Mane M, Missaoui B et al (2016) Premiers chaussages orthétiques dans la maladie de Charcot-Marie-Tooth. Medecine/Sciences. https://doi.org/10.1051/medsci/201632s206

24. Guillebastre B, Calmels P, Rougier P (2013) Effects of muscular deficiency on postural and gait capacities in patients with Charcot-Marie-Tooth disease. J Rehabil Med. https://doi.org/10.2340/16501977-1113

25. Hachisuka A, Hubenig L, Chan KM (2021) Patient compliance with orthotic use- can we do better? An editorial for Zuccarino et al. “Satisfaction with ankle foot orthoses in individuals with Charcot–Marie-Tooth.” Muscle Nerve 63(1):3–4. https://doi.org/10.1002/mus.27098

26. Panosyan FB, Laura M, Rossor AM et al (2017) Cross-sectional analysis of a large cohort with X-linked Charcot-Marie-Tooth disease (CMTX1). Neurology 89:927–935. https://doi.org/10.1212/WNL.0000000000004296

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