Dynamometer-based measure of spasticity confirms limited association between plantarflexor spasticity and walking function in persons with multiple sclerosis

Theodore R. Kremer  
Saint Louis University

Linda R. Van Dillen  
Washington University School of Medicine in St. Louis

Joanne M. Wagner  
Saint Louis University

Follow this and additional works at: http://digitalcommons.wustl.edu/open_access_pubs

Recommended Citation
http://digitalcommons.wustl.edu/open_access_pubs/3881

This Open Access Publication is brought to you for free and open access by Digital Commons@Becker. It has been accepted for inclusion in Open Access Publications by an authorized administrator of Digital Commons@Becker. For more information, please contact engeszer@wustl.edu.
Dynamometer-based measure of spasticity confirms limited association between plantarflexor spasticity and walking function in persons with multiple sclerosis

Theodore R. Kremer, BS;1 Linda R. Van Dillen, PT, PhD;2 Joanne M. Wagner, PT, PhD3*
1School of Medicine, Saint Louis University, St. Louis, MO; 2Program in Physical Therapy, Washington University School of Medicine, St. Louis, MO; 3Department of Physical Therapy and Athletic Training, Doisy College of Health Sciences, Saint Louis University, St. Louis, MO

Abstract—The literature shows inconsistent evidence regarding the association between clinically assessed plantarflexor (PF) spasticity and walking function in ambulatory persons with multiple sclerosis (pwMS). The use of a dynamometer-based spasticity measure (DSM) may help to clarify this association. Our cohort included 42 pwMS (27 female, 15 male; age: 42.9 +/- 10.2 yr) with mild clinical disability (Expanded Disability Status Scale score: 3.6 +/- 1.6). PF spasticity was assessed using a clinical measure, the modified Ashworth Scale (MAS), and an instrumented measure, the DSM. Walking function was assessed by the timed 25-foot walk test (T25FWT), the 6-minute walk test (6MWT), and the 12-item Multiple Sclerosis Walking Scale (MSWS-12). Spearman rho correlations were used to evaluate relationships between spasticity measures, measures of walking speed and endurance, and self-perceived limitations in walking. The correlation was small between PF spasticity and the T25FWT (PF maximum [Max] MAS rho = 0.27, PF Max DSM rho = 0.26), the 6MWT (PF Max MAS rho = -0.20, PF Max DSM rho = -0.21), and the MSWS-12 (PF Max MAS rho = 0.11, PF Max DSM rho = 0.26). Our results are similar to reports in other neurologic clinical populations, wherein spasticity has a limited association with walking dysfunction.

Key words: 12-item Multiple Sclerosis Walking Scale, ambulation, dynamometer, modified Ashworth Scale, multiple sclerosis, self-perceived limitations, spasticity, walking dysfunction, walking endurance, walking speed.

INTRODUCTION

Multiple sclerosis (MS) is a chronic, inflammatory disease of the central nervous system (CNS) characterized by demyelination and loss of motor and sensory axons within the brain and spinal cord [1]. Spasticity, defined as a motor disorder characterized by a velocity-dependent increase in tonic stretch reflexes and increased tendon jerks resulting from disinhibition of the stretch reflex [2], is a typical consequence of pathological changes within the corticospinal system. Lower-limb spasticity is believed to be a key contributor to mobility disability in persons with MS (pwMS). As such, lower-limb spasticity is routinely assessed in clinical practice to

Abbreviations: 6MWT = 6-minute walk test, CNS = central nervous system, DF = dorsiflexion, DSM = dynamometer-based spasticity measure, EDSS = Expanded Disability Status Scale, EMG = electromyography, MAS = modified Ashworth Scale, Max = maximum, MS = multiple sclerosis, MSWS-12 = 12-item Multiple Sclerosis Walking Scale, PF = plantarflexor, pwMS = persons with multiple sclerosis, ROM = range of motion, RRMS = relapsing-remitting multiple sclerosis, T25FWT = timed 25-foot walk test.

*Address all correspondence to Joanne M. Wagner, PT, PhD; Program in Physical Therapy, 3437 Caroline Mall, Suite 1026, St. Louis, MO 63104; 314-977-8532; fax: 314-977-8513. Email: jwagne34@slu.edu

http://dx.doi.org/10.1682/JRRD.2013.08.0186
document clinical disability and monitor disease progression, and therapies aimed at reducing spasticity are employed to promote better walking in pwMS [3–4].

PwMS perceive spasticity to be related to self-reported mobility disability [5]. Nonetheless, there is inconsistent evidence regarding the association between lower-limb spasticity, measured using the modified Ashworth Scale (MAS) [6], and walking speed, walking endurance, and self-perceived walking limitations. Sosnoff et al. reported on a group of 34 pwMS, 15 of whom exhibited plantarflexor (PF) spasticity [7]. In this cohort of pwMS, the group with PF spasticity had slower walking speed ($p = 0.02$), reduced walking endurance ($p < 0.01$), and greater self-perceived walking limitations ($p = 0.04$) than the group without spasticity. In contrast, we recently reported a small relationship between PF spasticity measured using the MAS and walking speed ($\rho = 0.27, p = 0.08$), walking endurance ($\rho = -0.20, p = 0.20$), and self-perceived walking limitations ($\rho = 0.11, p = 0.48$) [8]. There also were no differences in walking speed ($p = 0.34$), walking endurance ($p = 0.72$), or self-perceived walking limitations ($p = 0.64$) in pwMS with PF MAS scores greater than 0 ($n = 29$) compared with pwMS with PF MAS equal to 0 ($n = 13$).

The MAS is the most commonly reported clinical measure of spasticity; however, the objectivity of the MAS has been questioned [9]. Dynamometer-based spasticity measures (DSMs), which objectively quantify velocity-dependent resistance to passive movement, have been developed to address the subjectivity of the MAS [10–14]. Dynamometer-based estimates of lower-limb spasticity have confirmed a limited association between lower-limb spasticity, as measured using clinical scales, and walking dysfunction in other neurologic clinical populations [10–14]. Quantifying spasticity using a DSM in addition to the MAS may clarify the ambiguous evidence regarding the effect of lower-limb spasticity on walking dysfunction in pwMS. A better understanding of the association between lower-limb spasticity and walking dysfunction in pwMS is required for the development of targeted therapeutic interventions aimed at improving walking.

The purpose of this study was to determine the effect of PF spasticity on walking function in pwMS by examining the relationships between clinically assessed PF spasticity; a DSM of PF spasticity; and walking function assessed by standardized measures of walking speed, walking endurance, and self-perceived walking limitations. We hypothesized based on our previous work in this cohort [8], and based on studies of walking dysfunction in people with other CNS conditions, that there is a limited association between PF spasticity, whether measured clinically or using a dynamometer, and walking speed, walking endurance, and self-perceived walking limitations in pwMS.

### METHODS

#### Participants

Study participants included 45 ambulatory pwMS. All participants were recruited through the Multiple Sclerosis Clinic at Saint Louis University, The John L. Trotter Multiple Sclerosis Clinic at Washington University School of Medicine, and the Gateway chapter of the National Multiple Sclerosis Society. Participants were included (1) if they had a diagnosis of MS from a board-certified neurologist, including relapsing-remitting MS (RRMS), secondary progressive MS, and primary progressive MS; (2) if they were 18 to 65 yr old; (3) if they had minimal to moderate clinical disability as evidenced by an Expanded Disability Status Scale (EDSS) score ranging from 0.0 to 6.0, as determined by a board-certified neurologist; and (4) if they had RRMS but were at least 6 mo following their last clinical exacerbation of MS. Participants were excluded if they (1) had lower-limb orthopedic conditions that limited ambulation, (2) were pregnant, or (3) were unable to provide informed consent.

#### Data Collection

PF spasticity was measured bilaterally using the MAS [6]. The MAS provides a rating of spasticity on a 0 to 4 scale, where 0 = no increase in muscle tone and 4 = affected part(s) are rigid in flexion or extension. The MAS has been found to be a reliable measure in pwMS [15].

Walking function was assessed by standardized measures of walking speed, walking endurance, and self-perceived walking limitations. The timed 25-foot walk test (T25FWT), a component of the Multiple Sclerosis Functional Composite, is a standardized clinical measure of short-distance maximal walking speed [16]. The 6-minute walk test (6MWT) is a standardized clinical measure of walking endurance [17]. The 12-item Multiple Sclerosis Walking Scale (MSWS-12) is a self-report measure of the effect of MS on walking ability [18]. The MSWS-12
provides a score from 0 to 100, with larger values indicating a greater perception of walking difficulty due to MS. These tests were administered following standardized procedures by a physical therapist.

PF spasticity was also quantified using a Biodex System 4 dynamometer (Biodex Medical Systems, Inc; Shirley, New York). Both ankles were tested in a random order. Each participant was placed on the dynamometer in a semisupine position. For the tested limb, the leg was positioned so that the knee was in full extension and supported by the dynamometer chair. The foot was placed in the dynamometer dorsiflexion (DF)/plantar flexion attachment, and the ankle joint center was aligned with the center of rotation of the dynamometer. All participants were tested wearing shoes. Ankle range of motion (ROM) limits were preestablished to allow for a total of 40° of ROM, ranging from 30° PF to 10° DF. Waist and knee straps were used to stabilize the pelvis and lower limb, and a foot rest provided support for the nontested limb. For the test, the ankle was passively rotated from PF to DF 3 times at an angular velocity of 5 °/s and 10 times at angular velocities of 20, 60, and 90 °/s [19]. Participants were instructed not to help move the lever arm and remain relaxed while spasticity measurements were being taken. DSMs have moderate reliability [10,12]. Bipolar surface electromyography (EMG) electrodes (MA300 EMG system, Motion Lab Systems Inc; Baton Rouge, Louisiana) placed on the gastroc-soleus and anterior tibialis were used to visually monitor agonist and antagonist muscle activity, respectively, to ensure the movement was passive. Analog torque (foot-pounds), position (degrees), and velocity (degrees per second) signals were sampled directly from the dynamometer at a rate of 1 kHz using a PowerLab 16/30 (ADInstruments; Dunedin, New Zealand) and LabChart Pro version 7 software (ADInstruments) and saved directly to disk for subsequent analysis.

Data Analysis

Because the MAS includes a score of 1+, the raw PF MAS scores were transformed to a 0 to 5 scale [20]. The PF maximum (Max) MAS score (PF Max MAS) for each participant was determined and used for subsequent analysis. PwMS with PF MAS scores 1.0 or more for either leg were classified as having spasticity, whereas pwMS with PF MAS scores less than 1.0 on both legs were classified as not having spasticity [7].

For the DSM, data analysis was performed using LabChart Pro version 7 and MATLAB R2011b (The MathWorks Inc; Natick, Massachusetts). Torque angle data were low-pass filtered (50 Hz) then processed to correct for the effects of limb weight and gravity using an anthropomorphic method [21]. The weight of the foot was assumed to be 1.45 percent of the entire body weight [22]. The first trial of each velocity was excluded from analyses. To determine the average resistance to stretch for an entire trial, the work done on the dynamometer was calculated by determining the area under the gravity-corrected torque versus angle curve. To exclude end effects caused by the dynamometer, the boundaries of the area were set at −5°, −30°, the torque curve, and the zero torque line. Work can be calculated using the formula

\[
W = \int_{-5°}^{-30°} T \times d\theta
\]

where \( T \) = the torque produced against the dynamometer and \( d\theta \) = a small change in the angle (Figure 1) [10].

![Figure 1](image)

Figure 1.
Work was calculated by determining area between gravity-corrected torque versus angle curve and zero torque. To exclude end effects, boundaries of area were set at −5°, −30°, torque curve, and zero torque line. Work can be calculated using the equation, where \( T \) = torque produced against dynamometer and \( d\theta \) = small change in angle.
average work was calculated for each individual at each velocity tested. Linear regression was used to determine the slope of the line of best fit for the average work as a function of velocity (Figure 2). This slope has been shown to be a quantitative measure of spasticity in that it represents a velocity-dependent resistance to passive stretch [10]. The maximum slope value calculated for each participant was used for subsequent analysis (PF Max DSM).

**Statistical Analysis**

Statistical analyses were performed in SPSS version 20 (IBM Corporation; Armonk, New York). Significance was accepted at $p < 0.05$. Since distributions were not nor-
nal, Spearman rho correlations were used to evaluate relationships between spasticity measures (PF Max MAS, PF Max DSM) and measures of walking speed (T25FWT), walking endurance (6MWT), and self-perceived limitations in walking (MSWS-12). Interpretation of the magnitude of the correlation coefficients were as follows: weak ($r < 0.30$), moderate ($0.30 \leq r < 0.70$), and strong ($r \geq 0.70$) [23]. Based on our sample size, correlation coefficients greater than $0.30$ and $0.49$ were statistically significant at the $p < 0.05$ and $p < 0.001$ levels, respectively [24]. Mann-Whitney U tests assessed differences in DSM between pwMS with a MAS $\geq 1$ and pwMS with a MAS $= 0$. The magnitude of the difference between groups (pwMS with a MAS $\geq 1$ vs pwMS with a MAS $= 0$) was estimated by calculating the Cohen $d$ effect size [25].

**RESULTS**

Three participants were excluded from data analysis because their dynamometer data could not be processed: one participant due to an extremely limited DF ROM and two participants because the ankle was unable to be passively rotated due to ankle clonus or leg spasms confirmed by visual inspection of EMG. Table 1 provides demographics and clinical characteristics of the remaining participants ($n = 42$). Based on median EDSS scores, our sample of pwMS had mild clinical disability [26]. Of the participants, 12 (29%) reported using a straight cane for community ambulation. Seven (17%) pwMS used a cane during the T25FWT. Nine (21%) pwMS used a cane during the 6MWT. Three of these participants wore a unilateral ankle-foot orthosis during testing. Six (14%) pwMS reported the use of oral antispasticity medications.

Our sample of pwMS had mild PF spasticity based on the PF Max MAS and PF Max DSM values (Table 2) [27]. Of the participants, 69 percent (29/42) were classified as having clinically assessed spasticity based on a PF MAS score $\geq 1$ on either leg. The PF Max MAS and the PF Max DSM were measured in the same limb in 90 percent (38/42) of participants. PF Max DSM values did not differ between pwMS with a MAS $\geq 1$ and pwMS with a MAS $= 0$ (0.0040 $\pm$ 0.0015 vs 0.0037 $\pm$ 0.0012, $p = 0.38$, $d = 0.23$). There was a weak relationship between the clinically assessed measures and DSMs of PF spasticity (PF Max MAS vs PF Max DSM: $\rho = 0.260$, $p = 0.10$).

**DISCUSSION**

Our sample of pwMS had mild to moderate limitations in walking speed and endurance [17,28] and self-perceived limitations in walking (Table 2). Regardless of the method, PF spasticity weakly correlated with the T25FWT (PF Max MAS: $\rho = 0.27$, $p = 0.08$; PF Max DSM: $\rho = 0.26$, $p = 0.10$), 6MWT (PF Max MAS: $\rho = -0.20$, $p = 0.20$; PF Max DSM: $\rho = -0.21$, $p = 0.18$), and MSWS12 (PF Max MAS: $\rho = 0.11$, $p = 0.48$; PF Max DSM: $\rho = 0.26$, $p = 0.09$).
Self-reported walking difficulties become more common and more pronounced in pwMS as the severity of self-reported spasticity increases [5,35]. It is possible that the weak association found between PF spasticity and self-reported walking dysfunction in our cohort of pwMS is due to the low levels of spasticity in our cohort of pwMS. A stronger association might exist between PF spasticity, quantified using the MAS or DSM, and self-reported walking dysfunction in pwMS with more severe PF spasticity or in pwMS exhibiting clonus.

Our results differ from a report on pwMS where those with PF spasticity were found to have greater mobility impairment than those without PF spasticity [7]. In Sosnoff et al., the spasticity group had considerably greater clinical disability than the no spasticity group [7]. The association between PF spasticity and mobility impairment was not reported. In our cohort, clinical disability was similar between the spasticity and no spasticity groups. We have previously reported no differences in walking between the groups [8]. Because the level of PF spasticity was similar in both studies, it is possible that additional impairments contributing to overall disability (i.e., weakness, sensory loss, and ataxia) may have contributed to the group differences in mobility reported by Sosnoff et al. [7].

The weak association between the MAS and DSM in our cohort of ambulatory pwMS is congruent with reports in other neurologic clinical populations, demonstrating a limited association between clinical and instrumented measures of upper- and lower-limb spasticity [10,13–14,36]. Collectively, these results suggest that the MAS and DSM may quantify different components of spasticity. The MAS quantifies “increase in muscle tone,” which may be due to both reflex and nonreflex components of resistance to passive movement [6]. The DSM quantifies the change in work at different velocities. The nonreflex component (e.g., stiffness) of resistance to passive movement should be similar at different velocities. Therefore, we believe the DSM primarily quantifies the reflex component of resistance to passive stretch. Our study design does not allow us to determine the unique contribution of the nonreflex component of spasticity [37]. Differences in participant positioning for the two measures may also contribute to the weak association between clinical and instrumented measures. Additionally, the limited range of MAS scores in our cohort of ambulatory pwMS may have attenuated the relationship between the MAS and DSM [14].

The current study has limitations. First, pwMS in this investigation had low levels of clinical spasticity; therefore, the results may not generalize to ambulatory pwMS with more pronounced clinical spasticity. Second, pwMS exhibiting clonus or muscle spasms during the DSM were excluded from analyses because it invalidated our algorithm for processing velocity-dependent resistance to passive stretch. Third, the ROM during the DSM was truncated due to the deceleration artifact of the dynamometer. Thus, the DSM did not evaluate spasticity over the entire ROM. Fourth, because reflexes depend on the

<table>
<thead>
<tr>
<th>Measure</th>
<th>Mean</th>
<th>Standard Deviation</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>PF Max MAS (score)</td>
<td>1.43</td>
<td>1.33</td>
<td>0–4</td>
</tr>
<tr>
<td>PF Max DSM (J/[°/s])</td>
<td>0.0039</td>
<td>0.0014</td>
<td>0.0022–0.0096</td>
</tr>
<tr>
<td>T25FWT (s)</td>
<td>5.84</td>
<td>2.30</td>
<td>3.68–14.8</td>
</tr>
<tr>
<td>6MWT (m)</td>
<td>466.7</td>
<td>133.9</td>
<td>157.9–677.0</td>
</tr>
<tr>
<td>MSWS-12 (score)</td>
<td>42.9</td>
<td>22.3</td>
<td>0–90.5</td>
</tr>
<tr>
<td>PF Max MAS (score)</td>
<td></td>
<td></td>
<td>Median</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Interquartile Range</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>0–2</td>
</tr>
</tbody>
</table>

6MWT = 6-minute walk test, DSM = dynamometer-based spasticity measure, MAS = modified Ashworth Scale, Max = maximum, MSWS-12 = 12-item Multiple Sclerosis Walking Scale, PF = plantarflexor, T25FWT = timed 25-foot walk test.

Table 2. Spasticity and walking measures for persons with multiple sclerosis (n = 42).
specific task and limb positioning [38], spasticity measured in the semisupine position may not be equivalent to spasticity measured during ambulation. Fifth, spasticity was only evaluated at the ankle; we are unable to determine whether, and to what extent, spasticity of other muscle groups contribute to walking dysfunction in pwMS. Finally, our study was not designed to investigate the contribution of other lower-limb impairments to walking dysfunction.

CONCLUSIONS

PF spasticity has a limited association with walking speed, walking endurance, and self-perceived walking limitations in pwMS with mild clinical disability, regardless of whether spasticity was measured clinically using the MAS or using a dynamometer. The weak association between the DSM and the MAS suggests that the two tests may quantify different components of spasticity.

ACKNOWLEDGMENTS

Author Contributions:

Study concept and design: L. R. Van Dillen, J. M. Wagner.

Acquisition of data: J. M. Wagner.


Drafting of manuscript: T. R. Kremer, L. R. Van Dillen, J. M. Wagner.

Final approval of manuscript: T. R. Kremer, L. R. Van Dillen, J. M. Wagner.

Obtained funding: J. M. Wagner.

Financial Disclosures: Dr. Wagner serves on the speaker’s bureau of Acorda Therapeutics Inc, Ardsley, New York.

Funding/Support: This material was based on work supported by the National Institutes of Health (grant K12 HD055931, primary investigator: J. M. Wagner).

Additional Contributions: We thank Anne H. Cross, MD, for her assistance with participant recruitment. We thank all participants for their involvement in the study; Elissa Held Bradford, MSPT, NCS, MSCS, for assistance with data collection; and the Gateway chapter of the National Multiple Sclerosis Society for assistance with recruitment of participants.

Institutional Review: The study was approved by the Saint Louis University Institutional Review Board and the Washington University Human Research Protection Office, and all participants provided informed consent prior to participation.

Participant Follow-Up: The authors have no plans to notify the study subjects of the publication of this article because approval to contact participants about publication was not included in the informed consent process.

REFERENCES


Submitted for publication August 30, 2013. Accepted in revised form March 18, 2014.

This article and any supplementary material should be cited as follows: