Prevalence and characteristics of tremor in the NARCOMS multiple sclerosis registry: a cross-sectional survey

John R Rinker II,1,2 Amber R Salter,3 Harrison Walker,1 Amy Amara,1 William Meador,1 Gary R Cutter3

ABSTRACT

Objectives: (1) To describe the prevalence and severity of tremor in patients with multiple sclerosis (MS) registered within a large North American MS registry; (2) to provide detailed descriptions on the characteristics and severity of tremor in a subset of registants and (3) to compare several measures of tremor severity for strength of agreement.

Setting: The North American Research Committee on MS (NARCOMS) registry.

Participants: Registrants of NARCOMS reporting mild or greater tremor severity.

Outcome measures: We determined the cross-sectional prevalence of tremor in the NARCOMS registry over three semiannual updates between fall 2010 and fall 2011. A subset of registrants (n=552) completed a supplemental survey providing detailed descriptions of their tremor. Outcomes included descriptive characteristics of their tremors and correlations between outcome measures to determine the strength of agreement in assessing tremor severity.

Results: The estimated prevalence of tremor in NARCOMS ranged from 45% to 46.8%, with severe tremor affecting 5.5–5.9% of respondents. In the subset completing the supplemental survey, mild tremor severity was associated with younger age of MS diagnosis and tremor onset than those with moderate or severe tremor. However, tremor severity did not differ by duration of disease or tremor. Respondents provided descriptions of tremor symptoms on the Clinical Ataxia Rating Scale, which had a moderate to good (ρ=0.595) correlation with the Tremor Related Activities of Daily Living (TRADL) scale. Objectively scored Archimedes’ spirals had a weaker (ρ=0.358) correlation with the TRADL. Rates of unemployment, disability and symptomatic medication use increased with tremor severity, but were high even among those with mild tremor.

Conclusions: Tremor is common among NARCOMS registrants and severely disabling for some. Both ADL-based and symptom-descriptive measures of tremor severity can be used to stratify patients.

BACKGROUND

Associated with multiple sclerosis (MS) since the earliest descriptions of the disease, tremor is noteworthy for its strong association with disability1 and resistance to symptomatic treatment.2 The exact prevalence of tremor in the MS population is unknown; however, estimates from two prior studies suggest a prevalence range of 25–60%, with severe tremor in 3–15% of patients.3 4 Yet, the modest size of these studies (100 and 200 patients) and the populations they describe (a clinic-based cohort, and a community-based cohort with overall mild disease course5) allows for the possibility that their prevalence estimates might not generalise to the wider MS population.

A major challenge to accurately estimating the prevalence and severity of MS tremor arises from the difficulty in obtaining a large, representative cross-section of the MS population. The North American Research Committee on MS (NARCOMS) registry is a unique resource, containing over 36000 participants with MS, of whom more than 15 000 submit semiannual updates on various aspects of their disease,6 including the NARCOMS Tremor and Coordination Scale (TACS), which has been validated against clinically assessed measures of tremor.7 The size of the NARCOMS registry

Strengths and limitations of this study

This cohort represents the largest descriptive study of multiple sclerosis (MS)-associated tremor to date.

The data presented are mostly derived from patient-reported scales, which have been used in previous descriptive studies of MS tremor.

Results of the study could not be objectively validated; however, findings are similar to previous descriptive studies which did include objective validation.

These data reinforce previous smaller studies describing the disproportionate effect of tremor on employment, function and overall quality of life for patients affected with MS.


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allows for the identification of large subsets of patients with less common manifestations, such as severe tremor.

This work represents the largest descriptive study of tremor in MS to date, estimating its prevalence and severity in the NARCOMS registry and providing a detailed, patient-reported summary of the effects of tremor on daily function and quality of life (QOL). While there is no gold standard for the measurement of tremor severity in patients with MS, better normative data about how tremor severity relates to QOL in patients with MS could be applied in trial design for potential therapeutic interventions.

**METHODS**

**Tremor prevalence**

NARCOMS distributes semiannual surveys to all of its registered members asking a wide range of MS-related and QOL-related questions. Tremor and ataxia are assessed in these surveys by the TACS, which has been shown to have construct and criterion validity. On the scale, respondents score their tremor severity from 0 to 5 indicating absent, minimal, mild, moderate, severe or totally disabling tremor. Minimal indicates the presence of tremor without functional impairment, while the other levels indicate increasing impact of tremor on function.

In order to estimate tremor prevalence within NARCOMS, any TACS score of 1 or greater was counted towards the tremor prevalence. To approximate the methods used in prior prevalence studies of MS tremor, the prevalence of severe tremor (including totally disabling tremor) was also determined.

**Surveyed cohort**

With prior approval from the local IRB, we designed a supplemental printed survey that was mailed to a subset of NARCOMS registrants who were using MS disease modifying drugs (DMDs) and who indicated at least a mild effect of tremor on function on a NARCOMS update. While some demographic information and details of respondents’ MS was extracted from the master NARCOMS data set, the survey requested additional information specific to the experience of tremor, including handedness, distribution of tremor in the body, duration and family history of tremor, use of symptomatic medications to suppress tremor, and several scales designed to assess tremor severity (table 1). As part of the tremor severity assessments, respondents also drew Archimedes’ spirals with each hand using a pen provided with the survey. The pens were provided to minimise variability between respondents that might be introduced by differing writing instruments. Respondents were also asked to indicate current use of DMDs.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Instruments used in the study</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Name</strong></td>
<td><strong>Items (n)</strong></td>
</tr>
<tr>
<td>Tremor and Coordination Scale</td>
<td>1</td>
</tr>
<tr>
<td>Tremor Related Activities of Daily Living</td>
<td>25</td>
</tr>
<tr>
<td>Visual analog scale</td>
<td>1</td>
</tr>
<tr>
<td>Clinical Ataxia Rating Scale</td>
<td>4</td>
</tr>
<tr>
<td>Archimedes’ spirals</td>
<td>2</td>
</tr>
<tr>
<td>Tremor-related handicaps</td>
<td>9</td>
</tr>
<tr>
<td>Patient Determined Disease Steps</td>
<td>1</td>
</tr>
</tbody>
</table>

*MS, multiple sclerosis.*
the Patient Determined Disease Steps (PDDS), and general health-related QOL is gauged by the Short Form (SF)-12 physical and mental composite scores.

Archimedes’ spirals were scored by three raters (two primary, one secondary) with experience in the treatment of movement disorders. Each Archimedes spiral was assigned a score between 0 and 10, with 0 representing no evidence of tremor and 10 indicating inability to complete the spiral. Scores were assigned according to a scoring system published by Bain and Alusi. When the primary reviewer scores were identical or differed by 1 point, a mean of the two primary reviewer scores were used for that spiral. When scores differed by >1 point, the secondary reviewer’s score was incorporated into the mean score. A subset of spirals were duplicated and randomly distributed within the scored spirals to test intra-rater reliability. Each respondent’s left-handed and right-handed mean spiral scores were summed to generate a sum spiral score which was used for analysis. In order to account for spiral differences related to handedness, respondents were asked in the survey to self-identify their dominant hand.

Statistical methods

Statistical analysis was conducted using JMP 10.0.0 (SAS Institute Inc., Cary, North Carolina, USA). To determine the prevalence of tremor in the NARCOMS registry, we calculated the proportion of patients indicating a TACS score ≥1 relative to the total number of completed surveys across three consecutive semiannual survey periods: fall 2010, spring 2011 and fall 2011 (the ‘Tremor Prevalence Cohort’). Three periods were included in order to maximise the number of eligible respondents to whom we could send the paper tremor survey, which then defined the ‘Surveyed Cohort’. The cohort was stratified by tremor severity as measured by the TACS, as this scale alone is used to identify tremor on the semiannual NARCOMS updates distributed to active participants. Owing to the small number of respondents indicating totally disabling tremor on the TACS, these respondents were combined with those reporting severe tremor for the purpose of determining severe tremor prevalence.

Descriptive characteristics of the cohort were reported as frequencies, means with SDs, or medians with IQRs. Comparisons between groups were performed using analysis of variance for continuous outcomes (eg, age), and χ² tests were used to compare nominal data (eg, sex). Intra-rater and inter-rater reliability of the spiral ratings were determined by calculating κ scores and Spearman correlations for the three raters. Spearman’s correlations were calculated for the various tremor and disease severity scales, to determine strength of agreement. Missing data were handled through listwise deletion.

RESULTS

Tremor prevalence

The mean prevalence of tremor among NARCOMS respondents was approximately 45% for each of the three update periods (the ‘Tremor Prevalence Cohort’, table 2). Approximately one quarter of respondents experienced tremor with at least some impact on function (mild or worse tremor). In table 3, the NARCOMS Prevalence Cohort is compared to the London, UK and Olmsted County, USA prevalence studies on MS tremor. Among the approximately 3600 NARCOMS respondents with functionally significant tremor (TACS≥2), 777 were mailed supplemental surveys, and of those returned, 552 were included in the final database (the
Surveyed Cohort (figure 1). The Surveyed Cohort constitutes approximately 16% of NARCOMS registrants with mild or worse tremor. Respondents who completed the survey and attributed their tremor entirely to a cause other than MS (2.7% of respondents) were excluded from the final data set.

The distribution of tremor severity within the Surveyed Cohort differed somewhat from the Tremor Prevalence Cohort. Respondents in the Surveyed Cohort were less likely to characterise their tremor as mild (35.5% vs 42.7–45.8%), and somewhat more likely to identify themselves as having moderate (38% vs 31.8–32.7%) or severe (22.8% vs 16.3–18.9%) tremor. The ranges presented for the Prevalence Cohort derive from the three separate survey periods. PDSS Scores, reflecting global MS disability, were the same by TACS level between the Prevalence and Surveyed Cohorts (data not shown).

Demographic characteristics of the Surveyed Cohort are reported in table 4. Women comprised a greater proportion of those with mild tremor; while men were more likely to report severe or totally disabling tremor. Tremor severity did not differ between white and non-white respondents; however, NARCOMS is known to underrepresent minority ethnic groups. Current age, MS onset age, and tremor onset age all differed between TACS severity groups; however, neither duration of tremor nor duration of MS were significantly different between tremor severity groups.

‘Surveyed Cohort’, figure 1). The Surveyed Cohort constitutes approximately 16% of NARCOMS registrants with mild or worse tremor. Respondents who completed the survey and attributed their tremor entirely to a cause other than MS (2.7% of respondents) were excluded from the final data set.

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Demographics and distribution of tremor

Demographic characteristics of the Surveyed Cohort are reported in table 4. Women comprised a greater proportion of those with mild tremor; while men were more likely to report severe or totally disabling tremor. Tremor severity did not differ between white and non-white respondents; however, NARCOMS is known to underrepresent minority ethnic groups. Current age, MS onset age, and tremor onset age all differed between TACS severity groups; however, neither duration of tremor nor duration of MS were significantly different between tremor severity groups.

Figure 2 summarises the patient-reported descriptions of body regions affected by tremor. Similar to previous reports, the upper extremities were most commonly involved, and among those who reported unilateral tremor (n=208), respondents were twice as likely to report tremor in their dominant arm (n=140, 67.3%) as compared with their non-dominant arm (n=68, 32.7%).

Tremor severity measures

Both subjective and objective instruments have been developed to measure tremor severity. Subjective instruments generally relate tremor severity to QOL and interference with routine activities, while objective instruments rely on direct observation of tremor by a clinician expert or indirect rating of the effects of tremor, such as accelerometers, Archimedes’ spirals or handwriting samples. At present, there is no widely accepted ‘gold standard’ for assessing tremor severity in MS. As such, we chose to include a range of instruments including QOL-based measures (TACS, Tremor Related Activities of Daily Living (TRADL), SF-12 Health Survey), objectively scored Archimedes’ spirals and the descriptive (but not objective) CARS.

For the scoring of Archimedes’ spirals, κ and Spearman statistics were determined for inter-rater and intra-rater reliability. The κ scores reflect the consistency with which raters assigned identical scores to individual spirals, while the Spearman scores reflect how consistently spirals were ranked in order of severity. Intra-rater κ scores ranged from 0.48 to 0.56 (moderate agreement) and Spearman coefficients ranged from 0.79 to 0.89 (good agreement),
while inter-rater $\kappa$ scores ranged from 0.30 to 0.38 (fair agreement) and Spearman coefficients ranged from 0.81 to 0.85 (good agreement). While the $\kappa$ scores indicate broad inconsistency in assigning the same score to the same spiral, the Spearman scores suggest good consistency in ranking spirals by order of severity.

### Table 4 Demographics and descriptive characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Total cohort (n=550)</th>
<th>TACS=2 (n=196)</th>
<th>TACS=3 (n=209)</th>
<th>TACS=4 or 5 (n=145)</th>
<th>$p$ Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex, n (%)</td>
<td></td>
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<td></td>
<td></td>
<td>0.004*</td>
</tr>
<tr>
<td>Female</td>
<td>429 (78.0)</td>
<td>162 (82.7)</td>
<td>168 (80.4)</td>
<td>99 (68.3)</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>121 (22.0)</td>
<td>34 (17.3)</td>
<td>41 (19.6)</td>
<td>46 (31.7)</td>
<td></td>
</tr>
<tr>
<td>Race, n (%)</td>
<td></td>
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<td></td>
<td></td>
<td>0.821</td>
</tr>
<tr>
<td>White</td>
<td>485 (96.0)</td>
<td>175 (95.6)</td>
<td>179 (96.8)</td>
<td>131 (95.6)</td>
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</tr>
<tr>
<td>Non-white</td>
<td>20 (4.0)</td>
<td>8 (4.4)</td>
<td>6 (3.2)</td>
<td>6 (4.4)</td>
<td></td>
</tr>
<tr>
<td>MS subtype, n (%)</td>
<td></td>
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<td></td>
<td></td>
<td>0.005*</td>
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<tr>
<td>Relapsing–remitting</td>
<td>230 (48.9)</td>
<td>102 (59.3)</td>
<td>80 (47.3)</td>
<td>48 (36.9)</td>
<td></td>
</tr>
<tr>
<td>2° Progressive</td>
<td>145 (30.8)</td>
<td>48 (27.9)</td>
<td>50 (29.6)</td>
<td>47 (36.2)</td>
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</tr>
<tr>
<td>1° Progressive</td>
<td>27 (5.7)</td>
<td>5 (2.9)</td>
<td>8 (4.7)</td>
<td>14 (10.7)</td>
<td></td>
</tr>
<tr>
<td>Progressive-relapsing</td>
<td>30 (6.4)</td>
<td>6 (3.5)</td>
<td>14 (8.3)</td>
<td>10 (7.7)</td>
<td></td>
</tr>
<tr>
<td>Unsure</td>
<td>33 (7.0)</td>
<td>11 (6.4)</td>
<td>13 (7.7)</td>
<td>9 (6.9)</td>
<td></td>
</tr>
<tr>
<td>Current age, mean (SD)</td>
<td>55.3 (9.2)</td>
<td>53.4 (9.6)</td>
<td>56.0 (9.1)</td>
<td>57.0 (8.4)</td>
<td>&lt;0.001*</td>
</tr>
<tr>
<td>Age of MS diagnosis, mean (SD)</td>
<td>38.1 (9.5)</td>
<td>36.6 (9.4)</td>
<td>39.2 (9.6)</td>
<td>38.7 (9.3)</td>
<td>0.018*</td>
</tr>
<tr>
<td>Age of tremor onset, mean (SD)</td>
<td>43.6 (11.4)</td>
<td>41.8 (11.6)</td>
<td>44.4 (11.0)</td>
<td>45.0 (11.3)</td>
<td>0.021*</td>
</tr>
<tr>
<td>Duration of MS, mean (SD)</td>
<td>17.1 (8.4)</td>
<td>16.7 (8.2)</td>
<td>16.8 (7.9)</td>
<td>18.3 (9.2)</td>
<td>0.146</td>
</tr>
<tr>
<td>Duration of tremor, mean (SD)</td>
<td>11.6 (8.8)</td>
<td>11.2 (8.3)</td>
<td>11.6 (8.9)</td>
<td>12.1 (9.5)</td>
<td>0.702</td>
</tr>
<tr>
<td>Family history of tremor, n (%)</td>
<td>84 (15.5)</td>
<td>30 (15.7)</td>
<td>34 (16.4)</td>
<td>20 (14.1)</td>
<td>0.845</td>
</tr>
<tr>
<td>Patient Determined Disease Step (IQR)</td>
<td>5 (3, 6)</td>
<td>4 (2, 5)</td>
<td>5 (3, 6)</td>
<td>6 (4, 6)</td>
<td>&lt;0.001*</td>
</tr>
</tbody>
</table>

*p<0.05.

MS, multiple sclerosis; TACS, Tremor and Coordination Scale.

Figure 2 Bodily distribution of tremor among surveyed registrants. The figure depicts the number (%) of patients reporting tremor in each of 7 bodily regions. The table reports the proportion of respondents indicating tremor in either or both upper or lower extremities, and the occurrence of tremor in the dominant versus non-dominant arm. The histogram depicts how the tremor was distributed throughout the body for respondents with at least mild tremor.
The correlation between spiral sum scores and the CARS was weak ($p=0.288$, $p<0.001$). A summary of the Spearman correlations between the objective/descriptive instruments (spiral scores and CARS) and the subjective/QOL instruments is depicted in table 5. For all of the subjective QOL measures the correlation with the CARS was stronger than with spirography scores. The strongest correlation was between the CARS and the TRADL. The strength of this correlation between ADL impairments and specific abnormalities in body movement suggests patient-provided descriptions of tremor (on the CARS) correspond reasonably well to tremor impacts on activities of daily living (the TRADL). Correlations were weakest with the physical SF-12, which is neither MS-specific nor tremor-specific.

In an effort to improve the correlation strength between spirography and TRADL, we limited the correlation to those respondents indicating involvement of the dominant upper extremity. Even using this limited data set, the strength of the correlation only improved to 0.381.

Figure 3 depicts the association between tremor severity and unemployment, disability, and use of symptomatic medications (to suppress tremor severity). The intention was to ‘anchor’ the inherently subjective measures of severity to outcomes less prone to responder opinion. For comparison purposes, we included the TACS (since it is regularly included in NARCOMS updates) and the TRADL (owing to its detailed inventory of tremor-related QOL scenarios). As the level of tremor severity increased, an increasing proportion of those in each TACS level reported being unemployed and disabled. With respect to the use of symptomatic medications, the TRADL quartiles appeared to better correspond with use of tremor-suppressing medications, although the most severely affected patients on both scales appeared less likely to use symptomatic medications.

**DISCUSSION**

This study describes the largest cohort of patients with MS-associated tremor to date. In our study, the prevalence of both overall tremor and severe tremor fall between that described in the London clinic-based prevalence study and the Olmsted County population-based study. By utilising patient-reported data, our study reflects the characteristics of a large, heterogeneous patient population across a wide geographic area. While registry-based studies by nature are limited by the inability to objectively confirm the data being provided, the size of our cohort should alleviate some concerns about whether these results are indicative of the general MS population. Indeed, an advantage to our approach is that we are less prone to the bias towards greater disease severity often encountered in studies based in specialised academic centres, and less prone to the population homogeneity of a single-region population-based study.

Another aim of this study was to introduce an objectively scored measurement of tremor (the Archimedes’ spirals) into an otherwise subjective battery of outcomes. However, the correlation of spirography scores with subjective measures of tremor severity were at best only fair, suggesting the task of drawing spirals does not correspond with daily function. Likewise, drawing spirals would not be expected to capture resting or postural tremors, or tremors affecting parts of the body other than the distal upper extremities. We were also unable to control for patients bracing their drawing hands to improve the spiral quality, although the surveys did include instructions not to do so.

In contrast to the Archimedes’ spirals, the modified CARS, which asked respondents to put themselves in the place of an observant clinician and rate the severity of their tremor based on manoeuvres commonly performed in a neurological examination, yielded the strongest correlations with the tremor-specific (TRADL, TACS, visual analog scale) and the MS-specific (PDDS) disability measures. These results suggest patient-report measures of tremor need not be limited to daily function and QOL-anchored scales, although formal validation would further strengthen this conclusion.

By including multiple measures of tremor severity (eg, TRADL, TACS) we were also able to compare measures of tremor severity to end points such as employment and disability. Despite the limited scoring range of the TACS, it performed similar to the TRADL in stratifying respondents by tremor severity.

Additional interesting observations also bear mentioning. In all three prevalence studies on MS tremor, upper extremity tremor was by far the most common (>90% in London and Olmsted County studies, 77.1% in this study). In contrast, NARCOMS respondents were

**Table 5** Spearman correlations between descriptive and quality of life tremor assessment instruments

<table>
<thead>
<tr>
<th>Subjective measures of impairment or quality of life</th>
<th>Objective/descriptive measures of tremor</th>
<th>Spiral Sum Score</th>
<th>Clinical Ataxia Rating Scale</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tremor and Coordination Scale</td>
<td></td>
<td>0.203</td>
<td>0.368</td>
</tr>
<tr>
<td>Tremor Related Activities of Daily Living</td>
<td></td>
<td>0.358</td>
<td>0.595</td>
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<tr>
<td>Visual analog scale</td>
<td></td>
<td>0.246</td>
<td>0.485</td>
</tr>
<tr>
<td>Patient Determined Disease Steps</td>
<td></td>
<td>0.312</td>
<td>0.534</td>
</tr>
<tr>
<td>SF-12 Physical Health QOL</td>
<td></td>
<td>−0.145</td>
<td>−0.275</td>
</tr>
</tbody>
</table>

SF-12, Short Form 12; QOL, quality of life.
much more likely to report tremor in the lower extremities (68.5%) than in either of the two earlier studies (17–24%). The rates of head and truncal tremor were similar across all three studies. Since we know from our results that respondents were more likely to report tremor in a dominant upper extremity than in a non-dominant upper extremity, suggesting a reporting bias, they may likewise misidentify symptoms such as ataxia, clonus or spasticity in the lower extremities as tremor.

This study also found patients with MS with tremor report a high rate of tremor in their family history (15.5%), which was also observed by Bain and Alusi, who reported a rate of 7%. In comparison, the most common form of tremor among adults, essential tremor, affects approximately 0.9% of the adult population, and between 4.6% and 6.3% of the population older than 65 years. Whether this represents a bias towards increased recognition of tremor among affected families versus a biological predisposition towards tremor is unclear.

While our results in large part confirm or expand on prior observations regarding tremor in MS, our study does have limitations. As a registry-based study, our reports of tremor severity and other aspects of patients’ illness cannot be independently verified. However, past validation studies on the NARCOMS registry have largely
supported the validity and generalisability of its findings.17–19 Another potential criticism of this study is that participant selection for the Surveyed Cohort was not conducted completely at random. The selection criteria for the cohort were developed to answer a question related to the effects of DMDs on tremor over time, thus all patients who were not using DMDs were excluded, and only those with mild or worse tremor (indicating at least some interference with function) were invited to complete the detailed survey. By limiting the cohort to those patients with functionally relevant tremor, the correlations in this study are weakened by the absence of those patients with absent or very mild tremor. This excluded part of the tremor population also makes the correlations presented in this paper unsuitable for comparison to similar prior studies using the same instruments in a population containing the full range of tremor severity. In addition, since our population oversampled patients taking natalizumab as their DMD, the tremor severity. In addition, since our population oversampled patients taking natalizumab as their DMD, the sample may be biased towards patients with more severe disease, since natalizumab is commonly reserved for those patients with very active or severe MS.

In summary, this study confirms in a large patient registry that tremor is common among patients with MS, and that tremor can markedly detract from patient QOL and daily function. Better understanding the prevalence and severity of tremor in MS and how it impacts daily function could inform trial design for potential therapeutic interventions.

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Contributors JRR conceived and designed the study, performed statistical analysis, and authored and revised the manuscript. ARS participated in study design and statistical analysis, and assisted in drafting and revising the manuscript. HW, AA and WM each contributed to study design, performed blinded spiral ratings and critically revised the manuscript. GRC contributed to the design and statistical analysis of the study, and provided critical review of the manuscript. All authors provided final approval of the submitted manuscript.

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Competing interests None.

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Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

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