Original research

Functional movement disorder gender, age and phenotype study: a systematic review and individual patient meta-analysis of 4905 cases

Sarah C. Lidstone, Michael Costa-Parke, Emily J. Robinson, Tommaso Ercoli, Jon Stone, FMD GAP Study Group

ABSTRACT

Functional movement disorder (FMD) is a common manifestation of functional neurological disorder presenting with diverse phenotypes such as tremor, weakness and gait disorder. Our current understanding of the basic epidemiological features of this condition is unclear. We aimed to describe and examine the relationship between age at onset, phenotype and gender in FMD in a large meta-analysis of published and unpublished individual patient cases. An electronic search of PubMed was conducted for studies from 1968 to 2019 according to Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines. Individual patient data were collected through a research network. We described the distribution of age of onset and how this varied by gender and motor phenotype. A one-stage meta-analysis was performed using multilevel mixed-effects linear regression, including random intercepts for country and data source. A total of 4905 individual cases were analysed (72.6% woman). The mean age at onset was 39.6 years (SD 16.1). Women had a significantly earlier age of onset than men (39.1 years vs 41.0 years). Mixed FMD (23.1%), tremor (21.6%) and weakness (18.1%) were the most common phenotypes. Compared with tremor (40.7 years), the mean ages at onset of dystonia (34.5 years) and weakness (36.4 years) were significantly younger, while gait disorders (43.2 years) had a significantly later age at onset. The interaction between gender and phenotype was not significant. FMD peaks in midlife with varying effects of gender on age at onset and phenotype. The data gives some support to ‘lumping’ FMD as a unitary disorder but also highlights the value in ‘splitting’ into individual phenotypes where relevant.

INTRODUCTION

Functional movement disorder (FMD) is a common presentation of functional neurological disorder (FND)1 with various symptoms of altered movement, including tremor, jerks, dystonia and gait disorders, often displaying combinations of phenotypes. Although heterogeneous, people with different forms of FMD share aetiological risk factors, experience overlapping symptoms such as pain or fatigue and exhibit common findings on physical examination, including variability and distractibility.1 FMD is more commonly seen in women (70% in the largest studies).2 Despite its prevalence, the basic epidemiology of FMD is unclear. With some exceptions,3–6 large-scale epidemiological data are limited due to presentation heterogeneity, as most studies are smaller case series describing specific phenotypes. It is unknown whether different FMD phenotypes are associated with differences in age at onset (eg, whether functional parkinsonism is more common in older individuals) or if certain phenotypes exhibit gender differences (eg, the reported higher incidence of functional axial myoclonus in men).7 There is evidence that gender ratios may vary by age at onset in other forms of FND. A large case series of 698 subjects with functional seizures reported the mean age at onset in women was significantly younger than in men, yet was approximately equal between genders at the extremes of life.8 Characterising basic data in larger samples can uncover clues to underlying risk factors for developing FMD or potential underlying neurobiological mechanisms.

We collected a large data set on FMD from (1) a systematic review and individual patient data meta-analysis using published data, and (2) unpublished data from individual researchers. Adding unpublished data permits reliable augmentation of the data set beyond published studies. We also planned a comparison with the reported higher incidence of functional axial myoclonus in men.7 There is evidence that gender ratios may vary by age at onset in other forms of FND. A large case series of 698 subjects with functional seizures reported the mean age at onset in women was significantly younger than in men, yet was approximately equal between genders at the extremes of life.8 Characterising basic data in larger samples can uncover clues to underlying risk factors for developing FMD or potential underlying neurobiological mechanisms.

We collected a large data set on FMD from (1) a systematic review and individual patient data meta-analysis using published data, and (2) unpublished data from individual researchers. Adding unpublished data permits reliable augmentation of the data set beyond published studies. We also planned a comparison with the reported higher incidence of functional axial myoclonus in men.7 There is evidence that gender ratios may vary by age at onset in other forms of FND. A large case series of 698 subjects with functional seizures reported the mean age at onset in women was significantly younger than in men, yet was approximately equal between genders at the extremes of life.8 Characterising basic data in larger samples can uncover clues to underlying risk factors for developing FMD or potential underlying neurobiological mechanisms.

We collected a large data set on FMD from (1) a systematic review and individual patient data meta-analysis using published data, and (2) unpublished data from individual researchers. Adding unpublished data permits reliable augmentation of the data set beyond published studies. We also planned a comparison with the reported higher incidence of functional axial myoclonus in men.7 There is evidence that gender ratios may vary by age at onset in other forms of FND. A large case series of 698 subjects with functional seizures reported the mean age at onset in women was significantly younger than in men, yet was approximately equal between genders at the extremes of life.8 Characterising basic data in larger samples can uncover clues to underlying risk factors for developing FMD or potential underlying neurobiological mechanisms.

We collected a large data set on FMD from (1) a systematic review and individual patient data meta-analysis using published data, and (2) unpublished data from individual researchers. Adding unpublished data permits reliable augmentation of the data set beyond published studies. We also planned a comparison with the reported higher incidence of functional axial myoclonus in men.7 There is evidence that gender ratios may vary by age at onset in other forms of FND. A large case series of 698 subjects with functional seizures reported the mean age at onset in women was significantly younger than in men, yet was approximately equal between genders at the extremes of life.8 Characterising basic data in larger samples can uncover clues to underlying risk factors for developing FMD or potential underlying neurobiological mechanisms.
there would be gender differences among FMD phenotypes, which may be partly explained by differing age at onset.

METHODS
Study design
An individual patient data (IPD) meta-analysis was performed including published and unpublished FMD cases. The University of Toronto coordinated, collected and managed data, according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines.8 The systematic review protocol was not registered.

Eligibility criteria and phenotypic characterisation
Inclusion criteria for published data were applied at the study level. Studies were considered eligible if they contained individual patient data satisfying three minimal criteria: (1) diagnosis of FMD classified as either ‘documented,’ ‘clinically established’ or ‘probable;’10,11 (2) FMD age at onset (either explicitly stated or derivable from comparing age and symptom duration entries); and (3) sex/gender were reported. Only reports written in English were included. We carefully considered whether to use the term ‘gender’ or ‘sex’ for this IPD meta-analysis. Since most of the data set relied on patient/clinician self-report of identified gender, we chose to use this term. Accordingly, we use the terms ‘women’ and ‘men’ throughout this report and recognise that transgender patients would need to be included separately, which is noted further in the discussion. We could not collect both gender and sex in this retrospective study.

Inclusion criteria for unpublished data were used to ensure that the individual data aligned with the published data set as much as possible and included: (1) patients with FMD as a primary diagnosis including any hyperkinetic or hypokinetic movement disorder and limb weakness; (2) adults and children; (3) diagnosed by neurologists with movement disorders training and derived from neurological settings. Exclusion criteria were: (1) other FND types (eg, functional seizures); (2) Parkinson disease; (3) patients with comorbid additional movement disorders (eg, Parkinson disease); (3) patients with both FMD and functional seizures; (4) paroxysmal FMD (risk of overlap between episodic FMD and functional seizures).12

FMD phenotypes were classified into the following 10 categories, based on the most common FMD presentations: tremor, dystonia, gait disorder, weakness, jerks/myoclonus, mixed, facial symptoms, parkinsonism, other and unknown/not documented. ‘Mixed’ was included as a phenotype for cases with multiple phenotypes present at onset (eg, tremor and a gait disorder). ‘Facial symptoms’ encompassed facial dystonic spasms. ‘Other’ was included for other distinct FMD phenotypes. ‘Unknown/not documented’ accommodated cases which lacked phenotype characterisation. Adults and children were considered eligible. Harmonisation of criteria for published and unpublished data are described further in the online supplemental material.

Study identification and search strategy
Published data were collected from an advanced PubMed search from 1968 to 2019 using the following key words: ‘functional’, ‘psychogenic’ or ‘conversion’; in combination with any of the following: ‘motor’, ‘movement’, ‘gait’, ‘tremor’, ‘dystonia’, ‘weakness’, ‘myoclonus’, ‘tic’, ‘facial’, ‘paralysis’, ‘paroxysmal’, ‘jerks’ and ‘parkinsonism.’ Titles that included ‘seizure’, ‘epilepsy’ or ‘epileptic’ were excluded to remove reports of dissociative/psychogenic seizures. All titles using the term ‘functional’ unrelated to FMD or movement disorders were removed. The search was expanded by including all relevant references. Furthermore, additional records were identified through hand search, using the above-mentioned keywords, by one of the investigators. The results from the literature review and the hand search were combined, removing all duplicates. Papers were then excluded if they were missing the requisite variables of interest (sex/gender or age at onset), only group-level data were available and not IPD, or any cases where a ‘possible’ level of diagnostic certainty was recorded. Corresponding authors of papers only reporting group-level data were contacted and invited to share their published data in IPD format. Details of the systematic review can be found in the online supplemental methods.

Unpublished data sharing
Unpublished data were shared from experts in the field via an initial invitation to the Movement Disorders Society FMD Study Group. Members were invited to contribute de-identified, unpublished data from their own personal or research databases in compliance with their own institutional research ethics boards. See online supplemental material for details of the included individual data sets.

Outcomes
The outcomes of interest were the distribution of the age at onset and whether this exhibited gender differences and/or differed by FMD phenotype, and whether gender moderated any differences between phenotypes. We additionally compared our data set to a similar recently published data set of functional seizures,8 another well-defined subtype of FND.

Statistical analysis
Descriptive statistics were reported using means (SD) or frequencies (%), where appropriate. Histograms of age at onset were generated for the entire sample, for each of the six main FMD phenotypes, and split by gender. A Doornik-Hansen omnibus test was used to assess multivariate normality of age at onset, which was repeated for men and women separately. To formally evaluate the association with gender and phenotype, a one-stage IPD meta-analysis was performed with age at onset as the dependent variable in a multilevel mixed-effects linear regression model. Gender (men vs women) and phenotype (tremor; dystonia; gait; weakness; jerks/myoclonus; mixed; other/unknown) were included as fixed effects, whereas country (28 levels) and data source (111 levels) were included as random effects, to account for between-source heterogeneity. Parkinsonism and facial symptoms were not included in the regression analysis due to small numbers. The model was rerun using standardised age of onset (subtracting the mean and dividing by the SD to force the mean as 0 and SD as 1 to report results on both the original and SD scale) to help interpret the magnitude of effect sizes. An interaction term between gender and phenotype was included in a second model to test for potential moderating effects, that is, to detect if the effect of gender on age at onset differs by phenotype. We also planned: (1) a statistical comparison between individuals with age at onset less than 16 and those with age at onset over 60 and (2) a visual comparison of age at onset and gender with the largest sample of dissociative seizures, a distinct subtype of functional neurological disorder. All statistical analyses were performed using Stata V.16.1 (StataCorp). An assessment of risk of bias was waived given the minimal data set of basic demographic data.

RESULTS
The IPD selection process is summarised in figure 1. The systematic review is presented in the online supplemental material. The total data set consisted of 4905 individual FMD cases gathered from the systematic review and hand-search (n=878), shared published group-level data (n=570) and shared unpublished databases (n=3457). In total, there were 1448 individual FMD cases from all published sources included in the analysis. Unpublished IPD was collected...
from 33 separate international groups from 28 countries involving 70 authors (online supplemental figure 2 and online supplemental table 3).

**Data characteristics**

A summary of the descriptive statistics is provided in Table 1. The cohort comprised 3558 (72.5%) female patients. The mean age at onset was 39.7 years (median 39.8). Three hundred and eighty-three cases (7.8%) of the sample were paediatric onset (age ≤16), and 10.7% (526 cases) of the sample had age at onset over age 60. The most prevalent phenotypes were mixed FMD (23.0%) and tremor (21.6%), and the least prevalent were facial symptoms and parkinsonism. There were 479 (9.8%) individuals classified either as ‘other’ or ‘unknown’ phenotype (online supplemental table 4). Almost half

| Table 1 | Descriptive summary statistics and one-stage individual patient data meta-analysis mixed-effects linear regression model for age of symptom onset, with ‘tremor’ and ‘female’ as the reference categories |

<table>
<thead>
<tr>
<th>Gender</th>
<th>n (%)</th>
<th>Age of onset (Mean (SD))</th>
<th>Gender</th>
<th>Coefficient (95% CI)</th>
<th>Standardised effect size (95% CI)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Women</td>
<td>3558</td>
<td>39.1 (15.9)</td>
<td>Women</td>
<td>0.32 (−1.05 to 1.69)</td>
<td>0.02 (−0.07 to 0.10)</td>
<td>0.649</td>
</tr>
<tr>
<td>Men</td>
<td>1347</td>
<td>41.0 (16.5)</td>
<td></td>
<td>−1.66 (0.71 to 2.62)</td>
<td>0.10 (0.04 to 0.16)</td>
<td>&lt;0.001*</td>
</tr>
<tr>
<td>Phenotype</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mixed</td>
<td>1127</td>
<td>42.1 (16.3)</td>
<td>Women</td>
<td>−3.74 (−5.35 to 2.14)</td>
<td>−0.23 (−0.33 to 0.13)</td>
<td>&lt;0.001*</td>
</tr>
<tr>
<td>Tremor</td>
<td>1056</td>
<td>40.7 (16.6)</td>
<td></td>
<td>−4.31 (−5.98 to 2.65)</td>
<td>−0.27 (−0.37 to 0.16)</td>
<td>&lt;0.001*</td>
</tr>
<tr>
<td>Weakness</td>
<td>887</td>
<td>36.4 (13.4)</td>
<td></td>
<td>−5.31 (3.19 to 5.03)</td>
<td>0.20 (0.09 to 0.31)</td>
<td>0.001*</td>
</tr>
<tr>
<td>Dystonia</td>
<td>578</td>
<td>34.5 (14.8)</td>
<td></td>
<td>−5.98 (2.65)</td>
<td>0.06 (−0.07 to 0.20)</td>
<td>0.363</td>
</tr>
<tr>
<td>Gait</td>
<td>405</td>
<td>43.2 (18.4)</td>
<td></td>
<td>−3.19 (3.25)</td>
<td>0.32 (0.09 to 0.31)</td>
<td>0.001*</td>
</tr>
<tr>
<td>Jerks/myoclonus</td>
<td>223</td>
<td>39.8 (18.7)</td>
<td></td>
<td>0.20 (0.09 to 0.31)</td>
<td>0.363</td>
<td></td>
</tr>
<tr>
<td>Parkinsonism†</td>
<td>83</td>
<td>44.6 (12.9)</td>
<td></td>
<td>−1.19 (3.25)</td>
<td>0.32 (0.09 to 0.31)</td>
<td>0.001*</td>
</tr>
<tr>
<td>Facial symptoms†</td>
<td>67</td>
<td>37.3 (12.6)</td>
<td></td>
<td>−1.20 (3.25)</td>
<td>0.32 (0.09 to 0.31)</td>
<td>0.001*</td>
</tr>
<tr>
<td>Other/unknown</td>
<td>479</td>
<td>40.3 (15.8)</td>
<td></td>
<td>−1.11 (3.19 to 0.98)</td>
<td>0.07 (−0.20 to 0.06)</td>
<td>0.298</td>
</tr>
<tr>
<td>Total sample</td>
<td>4905</td>
<td>39.7 (16.1)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Coefficients are to be interpreted in terms of age at onset in years; and standardised coefficients are interpreted in SD. Positive coefficient means that the variable was associated with a later age of onset and negative coefficient with an earlier age of onset.

*Statistically significant at the p<0.05 level.
†Not inputted into linear regression model.
(42%) of the data set came from the USA. The remaining cases were from the Europe, Australia, Mexico and Canada. Countries in Africa, Asia and South America were under-represented (online supplemental material).

**Whole sample**
The distribution of the age at onset of the entire sample is presented in figure 2A. The data approached a bimodal distribution with peaks between the ages of 16–22 and 35–45. The Doornik-Hansen test for multivariate normality indicated the data were significantly different to the normal distribution, which holds for men and women separately despite there being less skewness in the male group.

**Gender differences**
When controlling for FMD phenotype (and heterogeneity between data sources within countries), men demonstrated a significantly older mean age at onset as compared with women (41.0 years vs 39.1 years) (table 1). Women represented greater than 64% of the sample across all age groups, with a tendency for more cases in the younger age groups (age 10–50) (figure 3A,C). By comparison, the age at onset of functional seizures in women, analysed from the Cognitive Behavioural Therapy versus Standardised Medical Care for Adults with Dissociative Non-Epileptic Seizures trial database of 698 individuals, produced a different distribution with a clear early peak in late adolescence, tapering into adulthood. This is in contrast to a similar frequency across the lifespan in men (figure 3B).

**Phenotype differences**
The mean age at onset was significantly different across phenotypes when controlling for gender and data source within country random effects (table 1). Compared with functional tremor, the mean age at onset of functional dystonia was younger by 4.3 years (95% CI 2.7 to 6.0 years, p<0.001), functional weakness was younger by 3.7 years (95% CI 2.1 to 5.3 years, p<0.001), while functional gait disorder presented with a significantly older mean age at onset, by 3.2 years (95% CI 1.4 to 5.0, p<0.05). The distribution of phenotypes (including ‘other’) followed roughly the same trajectory (ie, non-normal distribution with a peak in midlife) regardless despite there being less skewness in the male group.

**DISCUSSION**
This large individual patient data meta-analysis primarily provides detail on age of onset and gender for FMD and its phenotypes with unprecedented resolution compared with previous studies. We acknowledge that age of onset curves and gender form a limited data set on which to draw nosological conclusions. Nonetheless we think this data does offer some support for considering FMD both as a single entity but also one in which distinguishable phenotypes have validity. FMD is defined clinically through common historical features and positive signs on examination including variability and distractibility, but phenomenological differences have contributed in part to segregating FMD into separate disorders. Combining a large data set across phenotypes shows that FMD subtypes also share common characteristics (eg, similar shaped curves of age at onset) and have different epidemiological features (eg, 10-year difference between the mean age at onset of functional dystonia and functional gait disorder).

These results confirm previous much smaller studies demonstrating that FMD is more common in women (73%–75%) and symptom onset occurs most commonly in midlife. Given the probable under-representation of paediatric and elderly cases in the sample, we cannot be confident that this ratio is stable across the lifespan in FMD, although the same gender ratio (71%–74%) has been reported in paediatric case series. Prior studies have found gender differences among FMD phenotypes, including higher rates...
Figure 3  Histograms showing age of onset distribution by gender in (A) the whole sample of this study (n=4905) compared with (B) the previously published UK sample of functional seizure patients (n=698).3 C shows the comparison of FMD (red) and functional seizure (FS, green) patients displayed as percentage women with CIs. Figure adapted from Goldstein et al.8 FMD, Functional movement disorder.
of dystonia in women (84%). Of note, our study did not differentiate between axial and other types of myoclonus. Women are well-established to be more common than men in fibromyalgia, chronic fatigue syndrome and other functional disorders including irritable bowel syndrome. This gender difference has been attributed to several factors including the higher reported incidence of anxiety, depression and trauma in women, and that women are more likely to present to health services. Gender influences on the behaviour of the community, clinicians and patients is a social and psychological modifier of disease presentation, and a factor in determining how, when and why a person accesses medical care as well as what diagnoses they receive. However, autoimmune disorders such as multiple sclerosis and systemic lupus erythematosus are also more frequent in women highlighting the important role of biological sex differences, alongside gender differences.

Mixed FMD was the most common phenotype, followed by tremor, weakness and dystonia. Patients with FMD commonly display multiple phenotypes, both at a cross-sectional point in time (eg, at onset) and over time. Importantly, our results show that all phenotypes share a similar distribution (figure 4B) which, together with the frequency of ‘mixed’ phenotype, further supports FMD as a unitary disorder. Indeed, distinguishing between FMD phenotypes using clinical and demographic factors is challenging. Paroxysmal FMD and functional seizures can overlap. Our study excluded these cases for practical purposes, but it would be important to include these, as well as FMD occurring in comorbid neurological disease, in subsequent efforts to distinguish FND subtypes.

The onset of FMD occurs in midlife, ranging from 35 to 45 years, although this probably underestimates paediatric and elderly cases. The clustering of FMD mean age at onset in midlife and a second peak in late adolescence is a new finding (ie, a non-normal distribution). Our results confirm that functional dystonia has a younger age at onset and is more often seen in women, and a relatively higher proportion of men develop functional jerks and parkinsonism than other phenotypes (figure 5), although both still more commonly occur in women. A gender difference between different types of FMD was recently reported in a series of 196 patients with similar results.

Another advantage of the large data set is the ability to specifically look at paediatric and elderly onset cohorts. The most common phenotype in both groups was tremor followed by dystonia in younger patients and gait disorders in older patients, reflecting previous paediatric and elderly case series from which a portion of our data are drawn. Our analysis found no association between phenotype and age at onset or gender. This may reflect stability of FMD across the lifespan, supporting the view of a unitary diagnosis rather than a phenotype-based diagnosis.

We found different distributions of age at onset of FMD and functional seizures, two well-described FND phenotypes, which can overlap phenomenologically but tend to be clinically managed in segregated pathways. In women, functional seizures have a younger modal age at onset in the late teens and a mean in the late 20s, whereas men with functional seizures show a stable frequency of age at onset across the lifespan. This comparison also raises the possibility that, despite shared predisposing and precipitating factors in FMD and functional seizures, there may be important differences. Women may experience disproportionate trauma or adverse life events at younger ages than men, predisposing them to dissociation and developing functional seizures earlier in life, whereas physical injury or illness that may precipitate FMD may be more skewed towards older age groups.

The most significant limitation of this study is that two-thirds of the IPD was derived from unpublished databases. This approach,
however, enabled us to augment the data set by over 3000 cases. Harmonising published and unpublished data can result in underestimation of both eligible studies and individual cases and required efforts to try to ensure that true FMD cases were captured as reliably as possible. The clinical diagnostic criteria are shifting in the field toward the Diagnostic and Statistical Manual of Mental Disorders Fifth Edition DSM-5, which does not reflect the older diagnostic criteria we specified in our literature search. However, as long as we could reasonably certain that all cases included were FMD, given the bare minimal data set requested, there is value in this analysis, and in particular, because we collapsed across phenotypes. We relied on contributors to ensure that their data were accurate and that diagnostic clinical criteria were applied correctly, recognising that newer diagnostic criteria are routedly used in the clinical and may have also been used. We also relied on contributors to follow the inclusion criteria and ensure there were no duplications in their data. In addition, we did not request the timeframe for data collection if cases were consecutive as we felt these requirements would have substantially reduced the size of the data set. Although these cases were drawn from movement disorder specialists with a clinical interest in FMD, phenotypic characterisation can be challenging in this population. However, ‘mixed’, ‘other’ and ‘unknown’ categories, behaved as the more readily diagnosable phenotypes (eg, tremor), suggesting that diagnosis in these categories was as reliable.

These data are drawn from subspecialty clinic-based samples and do not reflect population-level epidemiological data. Furthermore, movement disorders clinics often see more complex or persistent FMD cases, underestimating mild or remitting FMD, thus recruitment bias may also influence the data. However, studying the epidemiology of FMD at a population level is inherently problematic since it normally requires secondary care expertise to make the diagnosis. FMD has high rates of comorbidity with other functional symptoms (including functional seizures) and this may have led to distortions in the data. There was an under-representation of paediatric and elderly cases, and data in these groups is less certain. We compared our case series data to the published functional seizures data set drawn from an interventional study. The systematic review was conducted in a single database, thus potentially under-representing published cases. The geographical distribution of this sample was heavily weighted to North America and Europe, however transcultural studies have not shown significant phenotypic differences in FMD.26 27 Factors determining age at onset and gender could potentially vary geographically. A post-hoc analysis using developed/developing country as a covariate in the main regression model did not impact the outcome.

Since we relied on retrospective data, we could only describe reported gender (from both published and unpublished sources) rather than both gender and biological sex. Most studies did not clarify whether sex or gender are reported so our study almost certainly conflates the two. Non-binary or transgender patients were not considered separately in our study. We are aware of four transgender patients in the data set and although unlikely to alter the conclusions, it is an important distinction that needs to be properly accounted for in any prospective data collection.

In summary, our data gives unprecedented resolution on the basic epidemiology of FMD which occurs across the lifespan but peaks in midlife with varying effects of gender on age at onset and phenotype. We do not wish to claim that our limited variable data set necessarily provides unique or novel insights into to the disorder. We do not think it can resolve, for example, whether this is a unitary disorder. Nonetheless we do see some support for ‘lumping’ FMD as a unitary disorder, at least in certain situations, on the ground that the FMD subtypes age of onset and gender profiles appear to have more in common with each other than with the ‘comparable’ movement disorder, for example, parkinsonism, tremor, and dystonia. However, with respect to the concept of ‘functional neurological disorder’ we have shown, visually at least, some potentially important differences between the basic epidemiology of FMD and functional seizures which generally appear larger than those between individual movement disorder phenotypes. Finally, there are statistical signals that not all phenotypes are equal with dystonia, myoclonus and gait disorder especially standing out.

This study provides a precedent for synthesising data across subphenotypes in functional disorders, which may reveal common, relevant underpinnings of FMD worthy of hypothesis testing. Future research should follow patients longitudinally to observe shifting phenotypes over time and determine the association between treatment outcomes and phenotypes and other biomarkers. Finally, our study also offers utility for health services planning and sets a new benchmark for international collaboration in FND.

Correction notice This article has been corrected since it first published. Affiliations have been added to the collaborator section. In addition, it has been made open access under a CC BY licence. 19th May 2023.

Twitter Sarah C. Lidstone @sarahlidstone, Tommaso Ercoli @ercolitommaso and Jon Stone @jonstoneneuro

Collaborators The FMD GAP Study Group includes the following authors: Omar Ahmad, Royal North Shore Hospital, Sydney, Australia; Sepidah Akbariapanahi, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, Maryland, USA; Alberto Albanese, Department of Neurology, IRCCS Humanitas Research Hospital, Rozzano Milano, Italy; Selma Aybek, University Inselspital Bern, Switzerland; Valek F. Bhatia, FRCP Radcliffe Infirmary Oxford, Oxford, UK; Carolina Bolognese, IRCCS Ospedale Maggiore Policlinico Milano, Milan, Italy; Riccardo Bolognini, Department of Clinical and Movement Neuroscience, Queen Square, London, UK; Verónica Cabrera, Department of Neurology, Centro Hospitalar Universitario de São João and Faculty of Medicine University of Porto, Porto, Portugal; Alan J Carson, Center for Clinical Brain Sciences, University of Edinburgh, Edinburgh, UK; Agnese Castagnola, IRCCS Fondazione Don Carlo Gnocchi Onlus, Milano, Italy; Russell C. Dale, Kids Neuroscience Centre, University of Sydney, Sydney, Australia; Carla Dallochio, Neurology Unit, Department of Medical Area, Pavia, Italy; Giovanni Defazio, Department of Medical Sciences and Public Health, University of Cagliari, Italy; Bertrand Degos, Department of Neurology, Avicenne University Hospital, Sorbonne Paris Nord University, Bobigny, France; Benedetta Demartini, Dipartimento di Scienze della Salute, Università degli Studi di Milano, Milano, Italy; Günter Desuchi, Department of Neurology, Christian-Albrechts Universität Kiel, Germany; Galina Diukova,Loginov Moscow Motor and Scientific Center, Moscow, Russia; Kevin R. Duda, Department of Neurology and Rehabilitation Medicine, Gardner Family Center for Parkinson’s Disease and Movement Disorders, University of Cincinnati, Cincinnati, Ohio, USA; Mark J Edwards, Neuroscience Research Centre, St George’s University of London, Cramer Terrace, London, UK; Steven A. Epstein, Georgetown University School of Medicine and MedStar Health, Washington DC, USA; Alberto J. Espay, Department of Neurology and Rehabilitation Medicine, Gardner Family Center for Parkinson’s Disease and Movement Disorders, University of Cincinnati, Cincinnati, Ohio, USA; Stewart A Factor, Jean and Paul Amos Parkinson’s Disease and Movement Disorders Program, Emory University, Atlanta, Georgia, USA; Beatrice Garbin, Department of Neurology, Avicenne University Hospital, Sorbonne Paris Nord University, Bobigny, France; ChristianGeroin, Neurology Unit, Movement Disorders Division, Department of Neurosciences, Biomedicine and Movement Sciences, University of Verona, Verona, Italy; Muriel Hagenauer, Université d’Antwerpen, Antwerpen, Belgium; Yvonne Hallett, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, Maryland, USA; Thomas Hassa, Lorjia Institute for Rehabilitation and Health Sciences, Allensbach, Germany; Anhar Hassan, Mayo Clinic, Rochester, Minnesota, USA; Lorena D. Herbert, University of Texas Southwestern, Dallas, Texas, USA; Samantha K. Holden, University of Colorado Anschutz Medical Campus, Denver, Colorado, USA; Joseph Jankovic, Parkinson’s Disease Center and Movement Disorders Clinic, Baylor College of Medicine, Houston, Texas, USA; Richard A. Kanaan, Department of Psychiatry, University of Melbourne, Austin Health, Heidelberg, Victoria, Australia; C.A. (Lianne) Kempe, Department of Mood Disorders, PsyG, Parramiss Groep, The Hague, The Netherlands; Maja Kojovic, Department of Neurology, University Clinical Centre Ljubljana, Slovenia; Kate Kompoliti, Department of Neurological Sciences, Rush University Medical Center, Chicago, Illinois, USA; Vedat Kostić, School of Medicine, University of Belgrade, Belgrade, Serbia; Christian Krüger, Department of Neurology and Psychiatry, Massachusetts General Hospital, Harvard Medical School, Boston, MA, USA; Kathrin La Faver, Feinberg School of Medicine, Northwestern University, Chicago, IL, USA; Anthony E. Lang, Edmond J.
Movement disorders

Safa Program in Parkinson’s Disease, the Morton and Gloria Shulman Movement Disorders Clinic, Toronto Western Hospital and the Division of Neurology, University of Toronto Lindsay MacGillivray PhD, MD Department of Psychiatry, University Health Network, University of Toronto, Canada; Davide Martino, Department of Clinical Neuroscience, Cumming School of Medicine, University of Calgary, Calgary, Canada; Júlia Massano, Department of Neurology, Centro Hospitalar Universitário de São João and Faculty of Medicine University of Porto, Porto, Portugal; Carine W. Maurer, Stony Brook University School of Medicine, Stony Brook, New York, USA; Laura McWherter, Centre for Clinical Brain Sciences, University of Edinburgh, Edinburgh, UK; Raja Mehanna, University of Texas Health Science Center at Houston and McGovern Medical School, Houston, Texas, USA; Francine Mesriti, Neuropsychology Department, Hospital Pitíà Sallêpêtrê, AP-HP, Paris, France; Jiong C. Morris, Washington University School of Medicine, St Louis, Missouri, USA; Glenn Nielsen, Neuroscience Research Centre, Institute of Molecular and Clinical Sciences, St Georges University of London, UK, UK; Anastasia Obukhova, I.M. Schenon First Moscow State Medical University, Moscow, Russia; Sanjay Pandey, Govind Ballabh Pant Institute of Postgraduate Medical Education and Research, JLN Marg, New Delhi, India; David L. Perez, Departments of Neurology and Psychology, Massachusetts General Hospital, Harvard Medical School, Boston, Massachusetts, USA; Igor Petrovic, Medical Faculty, University of Belgrade, Serbia; Seth L. Pullman, Columbia University Irving Medical Center, New York, New York, USA; Angelo Quartoare, Department of Biomedical Science and Morphological and Functional Images, University of Messina, Messina, Italy; Karin Roolofs, Radboud University, Behavioural Science Institute and Donders Institute for Brain, Cognition and Behaviour, Nijmegen, The Netherlands; Anette Schrag, Queen Square Institute of Neurology, University College London, London, UK; Yuri Selivestrov, Research Center of Neurology, Moscow, Russia; Tereza Sereanová, Charles University in Prague, 1st Faculty of Medicine and General University Hospital, Prague, Czech Republic; Ulf Sægård, Psychiatric Research Unit, Slagelse, Denmark; Petr Sojka, Department of Psychiatry, Faculty of Medicine, Masaryk University Brno and University Hospital Brno, Brno, Czech Republic; Maria Stamolou, Parkinson’s Disease and Movement Disorders Department, Hipogea Hospital, Athens, Greece. European University of Cyprus, Nicosia, Cyprus; Philipps University Marburg, Germany; Christopher D. Stephen, Movement Disorders Unit, Massachusetts General Hospital and Harvard Medical School, Boston, Massachusetts, USA; John F. Stins, Department of Human Movement Sciences, Faculty of Behavioural and Movement Sciences, Vrije Universiteit Amsterdam, Amsterdam, The Netherlands; Michele Tinazzi, Department of Clinical Neurosciences, Biomedical and Movement, University of Verona, Verona, Italy; Aleksandra Tomić, Neurology Clinic, Clinical Center of Serbia, and Faculty of Medicine, University of Belgrade, Belgrade, Serbia; Anaabela Valadas, Servico de Neurologia, Centro Hospitalar Universitário Lisboa Norte, Lisbon, Portugal; Valerie Voon, Department of Psychiatry, University of Cambridge, Cambridge, UK; Jeff L. Vaugh, University of Texas Southwestern, Dallas, Texas, USA; Allan D Wu, University of California Los Angeles, Los Angeles, California, USA.

Contributors SL—guarantor of overall content, conceptualisation, data curation, methodology, project administration, supervision, data validation, writing (original draft, review and editing). MC-P—data curation, project administration, data validation, writing (original draft, review and editing). TE—data curation, project administration, writing (original draft, review and editing). EJR—formal analysis, writing (original draft, review and editing). JS—conceptualisation, methodology, supervision, data validation, writing (original draft, review and editing). FMD-GAP Study Group—data sharing, feedback on initial manuscript, editorial, critical appraisal.

Funding SL and MC-P were supported by an anonymous donation to the Toronto Western Hospital Movement Disorders Clinic. JS was supported by an NHS Scotland NRS Career Fellowship.

Map disclaimer The inclusion of any map (including the depiction of any boundaries therein), or of any geographical or locational reference, does not imply the expression of any opinion whatsoever on the part of BMI concerning the legal status of any country, territory, jurisdiction or area or of its authorities. Any such expression remains solely that of the relevant source and is not endorsed by BMI. Maps are provided without any warranty of any kind, either express or implied.

Competing interests JS reports royalties from UpToDate for articles on FND and runs a free self-help website for people with FND. He carries out independent expert medicolegal work including in relation to FND. He is secretary of the FND society and on the medical advisory board for FND Hope and FND Action.

Patient consent for publication Not applicable.

Ethics approval This study involves human participants and was approved by University Health Network Research Ethics Board reference number #19-5901. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMI Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMI. BMI disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMI does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and it is not responsible for any error arising from omission or adaptation from translation or adaptation otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/licenses/by/4.0/.

ORCID iDs
Sarah C. Lidstone http://orcid.org/0000-0002-0147-0202
Tommaso Ercoli http://orcid.org/0000-0002-3260-1892
Jon Stone http://orcid.org/0000-0001-9829-8092

REFERENCES