Hand grip strength and fatigability: correlation with clinical parameters and diagnostic suitability in ME/CFS

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Abstract

Background: Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (ME/CFS) is a complex and debilitating disease accompanied by muscular fatigue and pain. A functional measure to assess muscle fatigability of ME/CFS patients is, however, not established in clinical routine. The aim of this study is to evaluate by assessing repeat maximum hand-grip strength (HGS), muscle fatigability as a diagnostic tool and its correlation with clinical parameters.

Methods: We assessed the HGS of 105 patients with ME/CFS, 18 patients with Cancer related fatigue (CRF) and 66 healthy controls (HC) using an electric dynamometer assessing maximal (Fmax) and mean force (Fmean) of ten repetitive measurements. Results were correlated with clinical parameters, creatinine kinase (CK) and lactate dehydrogenase (LDH). Further, maximum isometric quadriceps strength measurement was conducted in eight ME/CFS patients and eight HC.

Results: ME/CFS patients have a significantly lower Fmax and Fmean HGS compared to HC (p < 0.0001). Further, Fatigue Ratio assessing decline in strength during repeat maximal HGS measurement (Fmax/Fmean) was higher (p ≤ 0.0012). The Recovery Ratio after an identical second testing 60 min later was significantly lower in ME/CFS compared to HC (Fmean2/Fmean1; p ≤ 0.0020). Lower HGS parameters correlated with severity of disease, post-exertional malaise and muscle pain and with higher CK and LDH levels after exertion.

Conclusion: Repeat HGS assessment is a sensitive diagnostic test to assess muscular fatigue and fatigability and an objective measure to assess disease severity in ME/CFS.

Keywords: Myalgic encephalomyelitis/chronic fatigue syndrome, Diagnostic, Handgrip, Muscular fatigue, Muscle strength, Muscular recovery

Background

Chronic Fatigue Syndrome (also known as Myalgic Encephalomyelitis, ME/CFS), is a complex disease with persistent mental and physical fatigue causing severe impairment of quality of life. The cardinal symptom is exertional intolerance with post-exertional malaise (PEM), which describes a disproportionate intensification of symptoms and a prolonged regeneration phase after physical or mental effort [1–3]. Muscle fatigability is another important hallmark in which muscles become weaker after exertion and it can last for days before full muscle strength is restored.

To date, the etiology and pathophysiology of ME/CFS is still unresolved, but there is ample evidence for a disturbed vascular regulation [4]. Diagnosing
of ME/CFS is challenging for patients and physicians due to many unspecific symptoms, the broad differential diagnosis of chronic fatigue and the lack of an established biomarker. Currently, ME/CFS is a clinical diagnosis using comprehensive clinical evaluation and diagnostic criteria with the international Canadian Consensus Criteria (CCC) formulated by Carruthers et al. in 2003 as the most accepted [5]. However, to date ME/CFS patients are often un- or misdiagnosed as depression or burn-out, leading probably to a marked underestimation of prevalence [1, 6].

As objective measure to assess exertion intolerance in ME/CFS a repeat cardiopulmonary exercise test (CPET) is recommended by the National Institute of Health (NIH). Studies showed the maximum oxygen uptake and workload is significantly reduced at the day two of CPET [7]. However, the repeated exercise tests frequently lead to severe PEM and cannot be recommended for most patients [8].

Assessment of disease severity is mostly relied on questionnaires. Fatigue is a complex symptom and can be both mental and physical related to impaired muscle performance. Physical fatigue can be assessed objectively. Assessment of hand grip strength (HGS) is an established and highly reproducible tool to assess muscular force and provides information about the person’s physical function and state of health [9]. First studies showed that HGS is impaired in ME/CFS [10–12]. A recent study showed that maximal handgrip strength was significantly correlated with peak oxygen uptake and can predict maximal physical performance in CPET [13]. In the study by Meeus et al. also the HGS recovery after 45 min was analyzed and found to be impaired, while it was normal in HC [10]. Another approach to assess muscular force is the measurement of the quadriceps strength (QS). In an earlier study, ME/CFS patients showed lower QS and a prolonged recovery compared to the control group when performing consecutive contractions [14], another study reported normal quadriceps muscle force in ME/CFS [15].

Chronic fatigue occurs in many other diseases including cancer. About 30% of cancer survivors suffer from cancer-related fatigue (CRF) years after treatment severely impairing patients’ quality of life [16]. In order to assess specificity of HGS assessment for ME/CFS we included a cohort of CRF patients in our study.

In this study we evaluate repeat HGS measurement for assessing muscle fatigue in ME/CFS. By performing repeat hand grip testing fatigability and recovery of muscle strength could be assessed as diagnostic test.

### Methods

#### Patients and controls

105 patients with ME/CFS and 18 patients with CRF who presented at the outpatient clinic for fatigue at the Institute for Medical Immunology at the Charité Berlin from December 2018 to January 2021 were included in the study. ME/CFS patients were diagnosed based on the Canadian Consensus Criteria (CCC) and exclusion of other medical or neurological diseases which may cause fatigue [5]. Further inclusion criteria were age ≤18 years or participation in an interventional study. CRF patients were diagnosed according to Cella criteria [16] and had to be in complete remission for at least six months. 66 age and sex–matched HC with self-reported healthy status served as control group. For QS measurement we included eight female ME/CFS patients and eight female healthy controls.

#### Hand grip measurement

We measured the HGS with a digital hand dynamometer (CAMRY, model: SCACAM-EH101) in two separate sessions with a recovery break of 60 min between the sessions. The participant had to sit in an upright position and place the forearm of the dominant hand on a standard table in full supination. Before the start of the measurement, all participants had the opportunity to pull the handle twice to become familiar with the device. The handle was pulled with maximum force for three seconds followed by a five second relaxation phase under supervision by the study nurse. Within one session this procedure was repeated ten times with the dominant hand. After 60 min without any strenuous physical activity, a second session was conducted. Participants were verbally motivated during the measurement to continue using their maximum strength and to perform all repetitions. The dynamometer measures the highest value reached within the three seconds (force measurement in kg). The attempt with the highest measurement out of the ten repetitions was recorded as maximum strength.

#### Assessment of Fmax and Fmean strength, fatigability and recovery

To estimate strength, fatigability and recovery we examined several parameters and ratios listed in Table 1.

#### Quadriceps strength measurement

Maximal isometric muscle strength of the quadriceps muscle (expressed in Newton, N) was measured as described in previous publications [17]. Briefly, the freely hanging leg of the sitting participant was connected at the ankle with a pressure transducer.
Baseline maximal isometric strength was assessed from the best of three contractions on each leg, with a resting period of at least 60 s in between.

Afterwards the quadriceps muscle fatigue protocol was performed on the stronger leg as described previously [18]. In brief, participant performed repeated contractions at 30–40% of the maximum strength for one second, followed by one second of relaxations, using an acoustic signal as a guide. This cycle was performed for 40 s, followed by 20 s of rest. The 60 s cycle was repeated for 20 min. Participants were asked to undertake a maximal contraction at 5, 10, 15 and 20 min.

**Assessment of symptoms by scores**

Each participating ME/CFS patient filled in the following questionnaires.

**Bell score**

Assessment of disease severity and everyday restrictions ranging from zero (total loss of self-dependence) to 100 (without restrictions) [19].

**SF-36—Health status questionnaire**

Assessment of physical function ranging from zero (greatest possible health restrictions) to 100 (no health restrictions) [20].

**Chalder Fatigue Scale**

Assessment of the severity of fatigue on a scale from zero (no fatigue) to 33 (heavy fatigue) [21].

**COMPASS-31**

Assessment of autonomic dysfunction including vaso-motor, orthostatic, ocular, bladder and gastrointestinal symptoms ranging from zero (without symptoms) to 100 (strong autonomic dysfunction) [22] as was shown previously in ME/CFS patients [23].

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**Table 1** Parameters of muscle strength assessed by handgrip measurements

| Parameter/Ratio | Formula | Explanation |
|-----------------|---------|-------------|
| Fmax            | Fmax [kg] | Maximum grip strength within one session (ten repeat trials) |
| Fmean           | Σ10pulls/10 | Mean grip strength of all ten trials |
| Fatigue ratio   | Fmax/Fmean | Higher values indicate stronger decrease of force during one session |
| Recovery ratio  | Fmean2/Fmean1 | Low values indicate impaired recovery |

(Multitrace 2, Lectromed, Jersey, Channel Islands). Baseline maximal isometric strength was assessed from the best of three contractions on each leg, with a resting period of at least 60 s in between.

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**PEM criteria**

Assessment of frequency, severity and duration of post-exertional malaise (PEM) symptoms, such as muscle weakness, pain or mental tiredness, ranging from 0 (no PEM) to 46 (frequent, severe and long PEM) [24].

**Symptom Score**

Quantification of symptoms of the Canadian Consensus Criteria (one = no symptoms to ten = extreme symptoms) [25, 26].

**Assessment of CK and LDH**

Creatine kinase (CK, immunological UV-test) and lactate dehydrogenase (LDH, photometric test) were determined from blood samples drawn after the second HGS assessment at the Charité diagnostics laboratory (Labor Berlin GmbH, Berlin, Germany).

**Statistical analysis**

We performed the statistical analysis with the software GraphPad Prism 6.0. To exclude physiological, sex-related differences in strength, we studied male and female subjects separately. Gaussian distribution of data was examined by D’Agostino and Pearson test. Accordingly, we used unpaired t-tests and Mann–Whitney test to analyze differences between the groups. For comparison of the first and the second session we used paired t-test. For detection of linear and non-linear correlations we calculated Pearson and Spearman correlation coefficients, respectively. For analysis of the diagnostic ability of handgrip measurements and determination of discrimination thresholds between HC and ME/CFS, we performed receiver operator characteristics (ROC) analyses and computed sensitivity and specificity. A p-value of < 0.05 was considered as statistically significant. Due to multiple testing p-values are considered descriptive.

**Results**

**Study population**

105 ME/CFS patients were enrolled for HGS measurement. Control groups were 66 age- and sex-matched healthy controls (HC) and 18 female patients with cancer related fatigue (CRF). Table 2 provides further characteristics of the participants. Fmax and Fmean HGS

All 189 participants completed ten consecutive maximal HGS measurements which were repeated after 60 min. Both male and female ME/CFS patients showed strongly reduced maximal and mean HGS in the first session (Fmax1, Fmean1) and the second session after
one hour (Fmax2, Fmean2) compared to HC (Table 3 and Fig. 1, all \( p < 0.0001 \)). In female CRF patients Fmax and Fmean in both sessions were also significantly diminished compared to female HC (Table 3, Fig. 1, both \( p < 0.0001 \)).

Further, ME/CFS patients had a significantly lower Fmax and Fmean HGS in the second session compared to the first (Table 3 and Fig. 1). In female CRF patients, Fmax and Fmean were significantly reduced in the second session. In contrast female HC showed a similar Fmax and Fmean after one-hour break and male HC even a significant increase in Fmax and Fmean.

**Fatigability and recovery of HGS**

The Fatigue Ratio (Fmax/Fmean) was assessed as correlate of decrease of HGS during repeat measurements. ME/CFS patients had a stronger decrease in HGS resulting in higher Fatigue Ratios in comparison to HC in both first and second measurement (Table 3 and Fig. 2a). Further, Recovery Ratio of HGS (Fmean2/Fmean1) was diminished in the ME/CFS patients, as correlate of lower HGS during the second measurement, while HC had Recovery Ratios values of about one (Fig. 2b).

In female CRF patients, similar to ME/CFS, higher Fatigue Ratios 1 and 2 and reduced Recovery Ratios were observed compared to HC (Fig. 2 and Table 3).

**Correlation of HGS with symptom severity in ME/CFS patients**

We next correlated HGS with parameters of symptom severity and disability (Table 4 and Additional file 1: Figures S1, Additional file 2: Figure S2 and Additional file 4: Table S1).

Lower HGS parameters correlated with more disability (Bell Score), post-exertional malaise and

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### Table 2 Characteristics of the study population

|                  | HC           | ME/CFS       | CRF |
|------------------|--------------|--------------|-----|
| Sample size (n)  |              | 105          | 18  |
| Female           | 36           | 61           |     |
| Male             | 30           | 44           | 18  |
| Age (years)      | 42 (22–62)   | 49 (21–76)   | 44.5 (19–63) |
| Bell Score       | –            | –            | 30 (10–65) |
| Duration of disease (years) | – | 4 (1–32) | 4 (1–42) |
| Post-infectious ME/CFS | – | n = 52 | n = 40 |

Median values with range in brackets

HC healthy controls, ME/CFS myalgic encephalomyelitis/chronic fatigue syndrome, CRF cancer related fatigue, Bell Bell Score

### Table 3 Results of handgrip strength measurement

|                  | Females |               | Males |                 |
|------------------|---------|---------------|-------|-----------------|
|                  | HC, n = 36 | ME/CFS, n = 61 | CRF, n = 18 | HC, n = 30 | ME/CFS, n = 44 |
| Fmax1            | 28.2 (6.25) | 18.1 (6.68) | \( p < 0.0001 \) | 18.9 (6.73) | \( p < 0.0001 \) | 45.8 (10.2) | 31.2 (10.4) | \( p < 0.0001 \) |
| Fmax2            | 28.7 (5.73) | 16.0 (7.14) | \( p < 0.0001 \) | 17.2 (6.00) | \( p < 0.0001 \) | 48.3 (9.58) | 29.2 (10.9) | \( p < 0.0001 \) |
| Fmean1           | 25.6 (5.87) | 14.6 (6.09) | \( p < 0.0001 \) | 15.5 (6.34) | \( p < 0.0001 \) | 41.8 (9.66) | 26.3 (9.91) | \( p < 0.0001 \) |
| Fmean2           | 25.9 (5.46) | 12.9 (6.24) | \( p < 0.0001 \) | 14.3 (5.74) | \( p < 0.0001 \) | 44.2 (9.32) | 24.5 (10.4) | \( p < 0.0001 \) |
| Fatigue Ratio 1  | 1.105 (0.09) | 1.300 (0.297) | \( p < 0.0001 \) | 1.261 (0.21) | \( p = 0.0001 \) | 1.097 (0.04) | 1.213 (0.18) | \( p = 0.0006 \) |
| Fatigue Ratio 2  | 1.113 (0.07) | 1.299 (0.25) | \( p < 0.0001 \) | 1.223 (0.13) | \( p = 0.0004 \) | 1.097 (0.05) | 1.216 (0.17) | \( p = 0.0012 \) |
| Recovery Ratio   | 1.0 (0.077) | 0.87 (0.19) | \( p < 0.0001 \) | 0.96 (0.26) | \( p = 0.0020 \) | 1.1 (0.14) | 0.92 (0.14) | \( p < 0.0001 \) |

Comparison of first and second measurement

|                  | Females |               | Males |                 |
|------------------|---------|---------------|-------|-----------------|
|                  |         |               |       |                 |
| Fmax1            | 28.2     | 18.1          | \( p < 0.0001 \) | 18.9 | \( p = 0.0053 \) | 45.8 | 32.2 | \( p = 0.0019 \) |
| Fmax2            | 28.7     | 16.0          | \( p < 0.0001 \) | 17.2 | 48.3 | 29.2 |
| Fmean1           | 25.6     | 14.6          | \( p < 0.0001 \) | 15.5 | 41.8 | 26.3 | \( p = 0.0057 \) | 26.3 | \( p = 0.0022 \) |
| Fmean2           | 25.9     | 12.9          | \( p < 0.0001 \) | 14.3 | 44.2 | 24.5 |

Mean value of handgrip strength in kg and ratios with standard deviation in brackets; \( p \)-value refers to comparison with healthy controls (Mann–Whitney-Test) or, in the second compartment of the table, to comparison of both sessions (Paired t-test), respectively.
muscle pain. Specifically, the Bell score correlated with Fmax1 in females and with Fmean and Fatigue Ratios in males. Higher PEM scores correlated with lower Fmean2 in females and higher Fatigue Ratio2 in both sexes and more muscle pain with lower Fmean2 and Recovery Ratio in females. Strikingly, the COMPASS score for vasomotor dysregulation assessing changes of skin color on hands/feet correlated with Fmax2 and Fmean1,2 in both males and females and in Fmax2, Fatigue Ratio2 and Recovery Ratio in males, too. Physical functioning assessed by SF-36 questionnaire correlated with Fmax2 and Fmean2 in males.

No correlations were found with total COMPASS score and severity of fatigue assessed by Chalder Fatigue Scale (Additional file 4: Table S1).

Correlation of HGS with CK and LDH post exertion in ME/CFS patients

Finally, we correlated the muscle enzymes CK and the pyruvate to lactate catalyzing enzyme LDH assessed after exertion with HGS parameters (Table 4). Both CK and LDH are marker of muscle damage, too. In females, higher CK correlated with higher Fatigue Ratio2 and lower Recovery Ratio. Higher Lactate dehydrogenase (LDH) concentrations correlated in men with lower Recovery Ratio and higher Fatigue Ratio1 and in the female patients with higher Fatigue Ratio2.

Sensitivity and specificity of HGS assessment

In order to analyze the suitability of HGS parameters as diagnostic test we conducted operator characteristics (ROC) analyses for the HGS parameters in males and females age-grouped in 20–39 years and 40 – 59 years. Cutoff values and area under the curve (AUC) for age 20–39 years are listed in Table 5. The highest values for sensitivity and specificity showed the Fmean2 with an AUC of 0.94 at a cutoff of <19.95 kg for females and an AUC of 0.91 at a cutoff of <28.76 kg for males (Fig. 3, Table 5 and Additional file 3: Figure S3). Cutoff values and AUC for females and males aged 40–59 years are shown in Additional file 5: Table S2.

Quadriceps strength measurement

We performed five consecutive quadriceps strength (QS) measurements in eight female ME/CFS patients and eight female HC. Concordant to HGS, ME/CFS patients showed a significantly lower Fmax5 compared to Fmax1 compared to the HC resulting in a significantly lower Fmean (Table 6 and Fig. 4).

Discussion

In our study we show that patients with ME/CFS have reduced Fmax and Fmean HGS compared to HC. Our finding of impaired Fmax is in line with previous studies [10–13]. In addition, the greater decrease of HGS during ten repeated measurements compared to HC resulted in a higher Fatigue Ratio as was shown already in the study by Neu et al. [12]. Further, upon repeat measurement after an hour HGS was significantly lower in ME/CFS while it was fully recovered in HC indicating an impaired recovery rate in patients with ME/CFS as a parameter for fatigability. This finding is in accordance with the study by Meeus et al. describing an impaired recovery 45 min after 18 repeat maximal contractions [10].

Our study provides further evidence that repeat measurement of muscle strength via a hand dynamometer is a simple and useful method to objectively detect muscular fatigue and fatigability. ROC analyses revealed a high diagnostic sensitivity and specificity to distinguish between HC and ME/CFS with Fmean2 yielding best discrimination in women and men in both age groups. For assessing HGS as diagnostic marker in ME/CFS, we think it is important to use the Canadian Consensus Criteria requiring PEM. In contrast Ickmans et al. using the “Fukuda criteria” for diagnosis, which do not require PEM, observed significant differences in strength and recovery only for ME/CFS patients with comorbid fibromyalgia [27, 28].

Our findings are, however, not specific for ME/CFS as CRF patients showed significantly lower HGS parameters compared to HC, too. We found fatigability and impaired recovery also in female CRF patients. For HGS in patients with multiple sclerosis (MS) two studies delivered inconsistent findings. Nacul et al. showed diminished Fmax in MS patients in contrast to normal
a) Females and Males: Graphs showing strength variation with repetition for individuals with ME/CFS initial and after 1 hour compared to controls. Bars indicate confidence intervals.

b) Females and Males: Box plots comparing maximum force (Fmax) among different groups: HC, HC after 1h, ME/CFS, ME/CFS after 1h, CFS after 1h. Significant differences are marked with asterisks (*, **).

C) Females and Males: Box plots comparing mean force (Fmean) among the same groups as in b). Significant differences are marked similarly.
Fmax and recovery of HGS in the study by Meeus et al. [10, 11].

The quadriceps force measurement provides evidence that diminished muscular strength is found in leg muscles, too, showing in a similar pattern as the HGS that ME/CFS patients have a stronger drop in force upon repeated exertion. Thus, our study confirms the findings by Paul et al. [14] and is in contrast to the report by Gibson et al. of normal quadriceps muscle function in ME/CFS [15].

The correlations between HGS and clinical parameters underline the clinical relevance of HGS assessment in ME/CFS. Lower Bell scores (higher severity of disability) correlated with lower Fmax1 in women and lower Fmax and Fmean and higher Fatigue Ratio1 in men indicating a causal relation of muscular strength to severity of the disease. Remarkably, PEM correlated in women with lower Fmean2 and in women and men with higher Fatigue Ratio2, providing evidence that HGS is an objective marker to assess the severity of PEM. In line with this,
### Table 4  Clinical characteristics of ME/CFS and correlations with handgrip strength

|                | Female |                      | Male |                      |
|----------------|--------|-----------------------|------|-----------------------|
|                | Median | Range | n | Median | Range | n |
| **Bell**       |        |        |   |        |        |   |
| Correlations   | r      | p      | type | r      | p      | type |
| Fmax1          | 0.2670 | 0.0409  | lin  | 0.115  | 0.4682  | lin |
| Fmean1         | 0.1226 | 0.3551  | non-lin | 0.3255 | 0.0354  | non-lin |
| Fmean2         | 0.1444 | 0.2710  | non-lin | 0.3175 | 0.0405  | non-lin |
| Fatigue Ratio1 | 0.07946 | 0.5497  | non-lin | −0.3995 | 0.0088  | non-lin |
| Fatigue Ratio2 | −0.211 | 0.0643  | non-lin | −0.3624 | 0.0183  | lin  |
| **PEM**        |        |        |   |        |        |   |
| Correlations   | r      | p      | type | r      | p      | Type |
| Fmean2         | −0.334 | 0.0499  | non-lin | −0.1987 | 0.2526  | non-lin |
| Fatigue Ratio2 | 0.4040 | 0.0161  | lin  | 0.34  | 0.0457  | lin  |
| **Muscle Pain**|        |        |   |        |        |   |
| Correlations   | r      | p      | type | r      | p      | Type |
| Fmean2         | −0.3095 | 0.0287  | lin  | −0.1643 | 0.3175  | lin  |
| Recovery Ratio | −0.282 | 0.0472  | non-lin | −0.1542 | 0.3488  | non-lin |
| **COMPASS vasomotor** |        |        |   |        |        |   |
| Correlations   | r      | p      | Type | r      | p      | Type |
| Fmax1          | −0.2664 | 0.0643  | non-lin | −0.3537 | 0.0251  | non-lin |
| Fmax2          | −0.3635 | 0.0102  | non-lin | −0.4327 | 0.0026  | non-lin |
| Fmean1         | −0.3013 | 0.0354  | non-lin | −0.3784 | 0.0161  | non-lin |
| Fmean2         | −0.3663 | 0.0096  | non-lin | −0.5767 | <0.0001 | non-lin |
| Fatigue Ratio2 | 0.03817 | 0.7946  | lin  | 0.4869 | 0.0014  | lin  |
| Recovery Ratio | −0.1938 | 0.1821  | non-lin | −0.628 | <0.0001 | non-lin |
| **SF-36 phys. funct** |        |        |   |        |        |   |
| Correlations   | r      | p      | Type | r      | p      | Type |
| Fmax2          | 0.1724 | 0.2217  | non-lin | 0.3474 | 0.0351  | non-lin |
| Fmean2         | 0.2326 | 0.0970  | lin  | 0.3605 | 0.0284  | lin  |
| **CK**         |        |        |   |        |        |   |
| Correlations   | r      | p      | Type | r      | p      | Type |
| Fatigue Ratio2 | 0.3054 | 0.0348  | lin  | −0.1856 | 0.3092  | lin  |
| Recovery Ratio | −0.2935 | 0.0429  | lin  | −0.1045 | 0.5691  | lin  |
| **LDH**        |        |        |   |        |        |   |
| Correlations   | r      | p      | Type | r      | p      | Type |
| Fatigue Ratio1 | −0.06696 | 0.6547  | non-lin | 0.3522 | 0.0411  | non-lin |
| Fatigue Ratio2 | 0.3630 | 0.0322  | non-lin | 0.2683 | 0.125  | non-lin |
| Recovery Ratio | −0.0336 | 0.8226  | non-lin | −0.378 | 0.0275  | non-lin |

Clinical parameters with significant correlations with HGS and HGS ratios. Pearson (lin) or Spearman (non-lin) correlation coefficients were calculated. Bold: significant correlations with p < 0.05
severity of muscle pain correlated with lower $F_{\text{mean2}}$ and impaired recovery in the female cohort. A potential explanation is that more severe PEM is associated with more muscular fatigability and muscle pain. This might explain why pacing as a strategy to avoid PEM results in improvement of physical fatigue [29]. In the studies by Nacul et al. and Neu et al. fatigue assessed by a fatigue questionnaire correlated with $F_{\text{max}}$ HGS. In the male cohort we found correlations between the SF-36 physical functioning, and $F_{\text{max2}}$ and $F_{\text{mean2}}$. In our study we could not find an association of HGS parameters with the Chalder Fatigue Scale, which is, however, a questionnaire focusing mostly on mental fatigue (Additional file 6: Table S3).

Serum LDH and CK are both marker of muscle metabolism and damage and known to rise after exertion. Interestingly LDH and CK, which we determined after repeat HGS measurements also correlated with HGS parameters. Higher LDH concentrations correlated with higher Fatigue Ratio1/2 ($F_{\text{max}}/F_{\text{mean}}$) and thus fatigability in both sexes. LDH is an enzyme catalyzing the conversion of pyruvate to lactate and may indicate a preferential energy production via glycolysis rather than the more efficient oxidative phosphorylation. Diminished pyruvate dehydrogenase function is described in ME/

**Table 5** Cutoff values for hand grip strength (Age 20–39, ME/CFS vs HC)

| Sex       | Cutoff Value | AUC   | Sensitivity | Specificity |
|-----------|--------------|-------|-------------|-------------|
| $F_{\text{max1}}$ Females | < 23.55 kg | 0.8656 | 70 | 93.75 |
| Males     | < 33.35 kg  | 0.8762 | 68.42 | 94.12 |
| $F_{\text{max2}}$ Females | < 24.40 kg | 0.9156 | 90 | 93.75 |
| Males     | < 36.55 kg  | 0.9025 | 78.95 | 88.24 |
| $F_{\text{mean1}}$ Females | < 19.74 kg | 0.9188 | 80 | 100 |
| Males     | < 29.36 kg  | 0.8947 | 73.68 | 94.12 |
| $F_{\text{mean2}}$ Females | < 19.95 kg | 0.9438 | 85 | 100 |
| Males     | < 28.76 kg  | 0.9071 | 68.42 | 100 |
| Fatigue Ratio1 Females | > 1.161 | 0.8500 | 75 | 87.5 |
| Males     | > 1.197     | 0.5325 | 36.84 | 100 |
| Fatigue Ratio2 Females | > 1.200 | 0.8563 | 65 | 93.75 |
| Males     | > 1.189     | 0.5728 | 36.84 | 94.12 |
| Recovery Ratio Females | < 0.914 | 0.7906 | 70 | 93.75 |
| Males     | < 1.125     | 0.5820 | 100 | 23.53 |

Results of ROC Analysis for males and females aged 20–39 years. ME/CFS patients (females $n = 20$, males $n = 19$) and HC (females $n = 16$, males $n = 17$). Bold: most distinct parameter

**Table 6** Results of quadriceps strength measurement

| Max. quadriceps strength [Newton]          | HC, $n = 8$ | ME/CFS, $n = 8$ |
|-------------------------------------------|-------------|-----------------|
| QS initial ($F_{\text{max1}}$)            | 333 (82)    | 264 (108)       |
| 5 min ($F_{\text{max2}}$)                 | 324 (46)    | 237 (88)        |
| 10 min ($F_{\text{max3}}$)                | 333 (50)    | 246 (88)        |
| 15 min ($F_{\text{max4}}$)                | 334 (39)    | 212 (83)        |
| 20 min ($F_{\text{max5}}$)                | 342 (40)    | 212 (90)        |
| $F_{\text{max5}}$ in % of $F_{\text{max1}}$ | 106.4 (21.8)| 76.1 (13.0) p = 0.0045 |
| $F_{\text{mean}}$                         | 330 (46)    | 223.5 (88)      |
| Fatigue Ratio                             | 1.12 (0.058)| 1.23 (0.110) p = 0.1171 |

Mean value of quadriceps strength with standard deviation in brackets; p-value refers to comparison with healthy controls (t-Test)
CFS resulting in increased conversion of pyruvate to lactate [30]. Thus enhanced LDH may indicate diminished capacity of oxidative phosphorylation, which may well explain lower muscle strength during repeat exercise. Further we found that higher CK levels are associated with poorer Recovery Ratio and higher Fatigue Ratio in women. CK plays a central role in the energy supply of the muscle and muscular strain can increase the serum concentration of CK, although there are inter-individual differences [31]. A previous study found that CK concentrations are lower in patients with severe ME/CFS [32, 33]. This is not in contrast to our study as CK was analyzed after exertion in our study.

The pathomechanism of muscular fatigue and fatigability in ME/CFS is not fully elucidated yet. Histologic alterations with an increase in the more fatigue-prone, energetically expensive fast fibre type was described in muscle biopsies from ME/CFS patients, while contractile properties of muscle fibres were preserved [34]. Further metabolic alterations with altered expression of genes for mitochondrial and energy function were described [35]. Another study reported impaired glucose uptake and adenosine monophosphate activated protein kinase (AMPK) activity in cultured muscle cells from ME/CFS patients [36]. Fluge et al. showed that myoblasts grown in presence of serum from patients with severe ME/CFS showed increased mitochondrial respiration and excessive lactate secretion indicating impaired pyruvate dehydrogenase function [30]. Impaired oxygen supply to muscles upon exertion in ME/CFS was described, too. ME/CFS patients showed evidence of reduced hyperemic flow and reduced oxygen delivery [37]. In line with this hypoperfusion in cerebral vessels was recently shown [38]. A MRI study revealed that patients with ME/CFS have abnormalities in recovery of intramuscular pH following exercise which is related to autonomic dysfunction [39]. Paradox vasoconstriction due to β2 adrenergic dysfunction resulting in muscle hypoperfusion is considered as an important pathomechanism in ME/CFS [4]. In an own study we observed disease severity to correlate with endothelial dysfunction in ME/CFS [40]. In line with this concept, we observed here a strong correlation between vasomotor dysregulation assessed by COMPASS questionnaire with Fmax and Fmean of repeat testing.

A limitation of our study is that HGS may be influenced by inactivity although much less than leg muscle strength [41]. Further HGS is associated with age. Thus, a control group of physically inactive HC closely matched by age would be best for defining normal values of HGS parameters.

Conclusion
HGS measurement is a simple diagnostic tool to assess the severity of muscle fatigue in ME/CFS. Repeat HGS assessment further allows to objectively assess fatigability and impaired recovery. Advantages of HGS measurement are easy handling, low cost and the low risk of causing PEM. Thus, it can be implemented easily in both primary care and research as an objective outcome parameter in clinical studies and drug development.

Abbreviations
AMPK: Adenosine monophosphate activated protein kinase; ANOVA: Analysis of variance; AUC: Area under curve; BMI: Body mass index; CCC: Canadian consensus criteria; CDC: Center of disease control; CFS: Chronic fatigue syndrome; CK: Creatine kinase; CPET: Cardiopulmonary exercise testing; CRF: Cancer related fatigue; HC: Healthy controls; HGS: Handgrip strength; LDH: Liver dehydrogenase; ME/CFS: Myalgic encephalomyelitis/chronic fatigue syndrome; OCR: Oxidative phosphorylation rate; PCR: Polymerase chain reaction; PEM: Post-exertional malaise; RER: Respiratory exchange ratio; RQ: Respiratory quotient; RPE: Rating of perceived exertion; VO2: Oxygen consumption; VO2peak: Peak oxygen consumption; V̇O2: Oxygen uptake; WBT: Warm bath test; X: Abbreviation.
Lactic dehydrogenase; ME: Myalgic encephalomyelitis; MS: Multiple sclerosis; NIH: National Institutes of Health; PEM: Post-exertional malaise; QS: Quadriiceps strength; ROC: Receiver operating characteristic; SD: Standard deviation.

Supplementary Information

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Additional file 1. Figure S1 Correlations in male ME/CFS patients. All relevant correlations of HGS measurements with other clinical parameters with p<0.005 (Pearson and Spearman correlation coefficients, respectively). Line shows linear regression in Pearson correlations.

Additional file 2. Figure S2 Correlations in female ME/CFS patients. All relevant correlations of HGS measurements with other clinical parameters with p<0.005 (Pearson and Spearman correlation coefficients, respectively). Line shows linear regression in Pearson correlations.

Additional file 3. Figure S3 Fmean2 and cutoff value. Distinguishing between 20–39-year-old ME/CFS patients and HC using cut off values (dotted line) of Fmean2 determined by ROC analysis. Left: females (circles: ME/CFS, n=20, squares: HC, n=16), right: males (circles: ME/CFS, n=20, squares: HC, n=17).

Additional file 4. Table S1 Correlations of clinical parameters with HGS and HGS ratios. Pearson (lin) or Spearman (non-lin) correlation coefficients were calculated. Bold: significant correlations with p < 0.05.

Additional file 5. Table S2 Results of ROC-Analysis for females and males aged 40-59 years. ME/CFS patients (females n=35, males n=21) and HC (females n=18, males n=10). Bold: most distinct parameter.

Additional file 6. Table S3 Raw Data.

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Authors’ contributions

BJ conducted the HGS measurements, collected and analyzed and interpreted the data and was a major contributor in visualization and writing the manuscript. CK, PG and KW relevantly contributed to the investigation process by conducting the consultation, diagnosing, and inclusion of patients and data assessment. ST was responsible for project administration and data collection. ST performed HGS measurements. NS conducted the QS measurements, analyzed and interpreted the data and contributed to write the manuscript. NS and WD conceptualized and administered QS experiments. CS and HF analyzed and interpreted the data and was a major contributor in visualization and writing the manuscript. Additionally, CS executed project administration in writing the manuscript. Additionally, CS executed project administration in writing the manuscript. CK, PG and KW relevantly contributed to the investigation process by conducting the consultation, diagnosing, and inclusion of patients and data assessment. ST was responsible for project administration and data collection. ST performed HGS measurements. NS conducted the QS measurements, analyzed and interpreted the data and contributed to write the manuscript. NS and WD conceptualized and administered QS experiments. CS and HF analyzed and interpreted the data and was a major contributor in visualization and writing the manuscript.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article and its Additional files.

Declarations

Ethics approval and consent to participate

All participants were informed in advance and gave their written consent. The study was approved by the ethics committee of the Charité (EA2/038/14 from 12th Aug 2015) and the data protection regulations were followed.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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