Article

Differential Effects of Exercise on fMRI of the Midbrain Ascending Arousal Network Nuclei in Myalgic Encephalomyelitis / Chronic Fatigue Syndrome (ME/CFS) and Gulf War Illness (GWI) in a Model of Postexertional Malaise (PEM)

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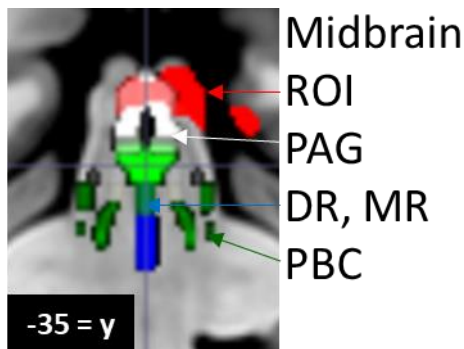   hsp9@georgetown.edu

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Abstract: Background: Myalgic Encephalomyelitis / Chronic Fatigue Syndrome (ME/CFS), Gulf War Illness (GWI) and control subjects had fMRI during difficult cognitive tests performed before and after submaximal exercise provocation (Washington 2020). Exercise caused increased activation in ME/CFS but decreased activation for GWI in the dorsal midbrain, left Rolandic operculum and right middle insula. Midbrain and isthmus nuclei participate in threat assessment, attention, cognition, mood, pain, sleep, and autonomic dysfunction. **Methods:** Activated midbrain nuclei were inferred by re-analysis of data from 31 control, 36 ME/CFS and 78 GWI subjects using a seed region approach and the Harvard Ascending Arousal Network. **Results:** Before exercise, control and GWI had greater activation during cognition than ME/CFS in left pedunculotegmental nucleus. Postexercise ME/CFS had greater activation than GWI for midline periaqueductal gray, dorsal and median raphe, and right midbrain reticular formation, parabrachial complex and locus coeruleus. The change between days (delta) was positive for ME/CFS but negative for GWI indicating reciprocal patterns of activation. Controls had no changes. **Conclusions:** Exercise caused opposite effects with increased activation in ME/CFS but decreased activation in GWI indicating different pathophysiological responses to exertion and mechanisms of disease. Midbrain and isthmus nuclei contribute to postexertional malaise in ME/CFS and GWI.

Keywords: Midbrain, postexertional malaise, PEM, arousal, exercise, fMRI, autonomic, postural tachycardia, Myalgic Encephalomyelitis / Chronic Fatigue Syndrome, ME/CFS, Gulf War Illness, GWI

Visual Abstract

Two bouts of submaximal exercise caused BOLD activity during a cognitive task to be significantly greater in the dorsal midbrain (red region of interest) of Myalgic Encephalomyelitis / Chronic Fatigue Syndrome (ME/CFS) subjects than Gulf War Illness (GWI) veterans. In this coronal section at y= -35, the region of interest (red) extended from the right inferior colliculus (top) to periaqueductal gray (PAG, white) and right lateral midbrain. DR dorsal raphe, MR, median raphe (blue), and PBC parabrachial complex (dark green) were not involved at this level of section.
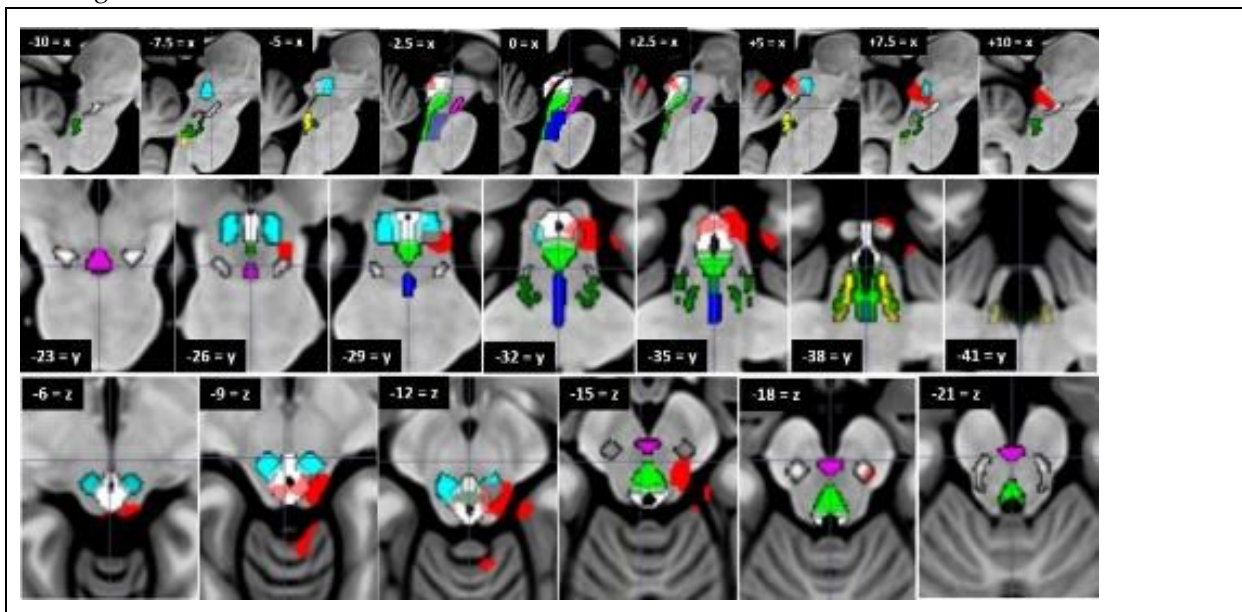
1. Introduction

Myalgic Encephalomyelitis / Chronic Fatigue Syndrome (ME/CFS) [1,2] and Gulf War Illness (GWI) [3,4] share features of post-exertional malaise (PEM, exertional exhaustion), fatigue that is not relieved by rest, unrefreshing and non-restorative sleep, total body pain and systemic hyperalgesia. ME/CFS has been considered to be a chronic consequence following flu-like epidemics [5] but in general has a sporadic heterogeneous presentation and unknown etiology. Prevalence is about 0.2 to 2% [6,7]. The 1994 Center for Disease Control criteria require moderate to severe unremitting fatigue of new onset that persists longer than 6 months and has no explanation despite appropriate medical investigations and at least four of the following eight ancillary criteria: cognitive complaints of short-term memory or concentration, sore throat, sore lymph nodes, myalgia, arthralgia, headaches including migraine, disordered sleep and post exertional malaise (PEM) [1]. Emphasis has been placed on PEM, the characteristically delayed exacerbation of the entire symptom complex following minimal physical, cognitive or emotional efforts, as a distinguishing feature of ME/CFS [3,4,7,8].

GWI affects 25 to 32% of veterans deployed to the 1990–1991 Persian Gulf War. The Centers for Disease Control criteria for Chronic Multisymptom Illness (CMI) require symptoms from at least two of three clusters: general fatigue; mood and cognitive abnormalities; and myalgia/arthralgia (pain) [3]. Mood/cognitive symptoms can range from troubles with sleep, cognitive difficulties, anxiety and depressive mood. An epidemiological comparison of symptoms between deployed and non-deployed Kansas veterans generated a more sensitive set of criteria requiring three of six domains: fatigue and sleep; pain; neurological/cognitive/mood; gastrointestinal; respiratory; and skin symptoms [4]. The etiology has been linked to exposures to neurotoxicants that were present in theatre, including organophosphates, carbamates and other pesticides, sarin/cyclosarin nerve agents and pyridostigmine bromide used as prophylactic medication against chemical warfare attacks [9,10]. Symptoms are consistent with Chronic Organophosphate Induced Neuropsychiatric Disorder (COPIND) [11–13]. Psychiatric aetiologies have been ruled out [9].

The overlapping symptoms of GWI and ME/CFS have generated unifying hypotheses [10]. However, pathophysiological mechanisms and objective findings that can be used for disease diagnosis and prediction of potential therapies have been more difficult to identify. We approached this problem by developing a two day submaximal exercise provocation paradigm with symptom, heart rate variability, and functional magnetic resonance imaging to assess PEM and

other changes induced by the exertional challenge. In a previous study, we examined blood oxygenation level dependent (BOLD) activation during a difficult, high cognitive load continuous 2-back working memory task and compared preexercise and postexercise scans. Control, ME/CFS and GWI were equivalent prior to exercise (baseline), but after exercise ME/CFS had a significant increase in blood oxygenation level dependent (BOLD) activation while GWI had a significant decrease in the dorsal midbrain, right middle insula and left Rolandic operculum [14]. The midbrain region of interest extended from the left to right periaqueductal gray (PAG) and to the adjacent right midbrain reticular formation (MRF), inferior colliculus and lateral lemniscus, and caudally to the right lateral isthmus (Figure S1). Because these nuclei have profound influences on threat assessment, pain, negative emotion, attention, wakefulness, and instinctual neurobehaviours, it was of interest to assess the activation of relevant anatomical midbrain nuclei.

SOM Figure S1.

In this report, we re-analyze the original BOLD data [14] using a seed region approach to gain a preliminary understanding of the nuclei that were activated within the midbrain region of interest. The seed regions were selected from the ascending arousal network that was defined from histological sections and diffusion studies of brainstem white matter tracts [14]. BOLD signals in each of the target nuclei were assessed on preexercise and postexercise days. The aim was to identify which midbrain nuclei were affected by exercise in ME/CFS and GWI subjects and to judge effect sizes to guide future confirmatory studies. We propose that affected nuclei may participate in the pathology of postexertional malaise.

2. Methods

2.1 Ethics

All subjects gave written informed consent to this protocol that was approved by the Georgetown University Institutional Review Board (IRB 2009-229, 2013-0943 and 2015-0579) and Department of Defense Congressionally Directed Medical Research Program (CDMRP) Human Research Program Office (HRPO) (A-15547 and A-18479) and listed in clinicaltrials.gov (NCT01291758 and NCT00810225). All clinical investigations were conducted according to the principles expressed in the Declaration of Helsinki.

2.2 Demographics

GWI, ME/CFS and healthy control subjects were recruited to this four day long in-patient study in the Clinical Research Unit of the Georgetown – Howard Universities Center for Clinical and Translational Science. Subjects had history and physical examinations to ensure their inclusion by meeting Chronic Multisymptom Illness [3] and Kansas [4] criteria for GWI, Fukuda [1] and Canadian [2] criteria for ME/CFS, confirmation of sedentary lifestyle for control subjects (less than 40 min of aerobic activity per week) and exclusion because of serious medical or psychiatric conditions such as psychosis [4,15–17]. History of posttraumatic stress disorder (PTSD) [18] or depression [19] were not exclusions unless the subject had been hospitalized in the past 5 years. Subjects completed Chronic Fatigue Syndrome Symptom Severity [20], SF-36 quality of life [21], Chalder Fatigue [22] and McGill Pain [23] questionnaires, and had systemic hyperalgesia tested by dolorimetry [24,25].

Light sensitivity was assessed by comparing scores while supine looking up at standard hospital ward fluorescent lighting turned on or off with curtains shut and background ambient light. Sound sensitivity was assessed by dropping a standard hardboard clip board (#44292, Item #1671406, Model #ST44292-CC, Staples, Inc.) from 3 feet onto a linoleum floor. The sound intensity 6 feet away at the subject's ear was approximately 60 dB (ambient noise ~ 35 dB) (Physics Toolbox Sensor Suite). Severity was scored using the modified 0 to 20 point anchored ordinal Intensity scale of the Gracely Box score [26–28].

2.3 Exercise provocation

Two submaximal bicycle exercise tests were performed 24 h apart. Subjects cycled for 25 min at 70% predicted maximum HR (220 minus patient's age), followed by a climb to 85% maximum HR to reach anaerobic threshold [14,26,27,29,30].

2.4 Orthostatic postural tachycardia phenotypes

Subjects rested supine for 5 minutes with continuous EKG measurements and arm cuff blood pressure every minute. Average recumbent heart rate (HR) was calculated. Subjects stood up and maintained their posture for 5 minutes. EKG and blood pressure were recorded each minute. The differences between standing HR at each minute and average recumbent HR were calculated (ΔHR). The procedure was performed at least twice before the first exercise, then 1, 3, 8 and 24 hr postexercise. ΔHR was used to define Orthostatic status [26,27].

(i) Postural orthostatic tachycardia (POTS) was defined by ΔHR ≥ 30 beats per minute at least 4 time points before exercise and during each postexercise measurement period.

(ii) Stress Test Activated Reversible Tachycardia (START) was defined by normal ΔHR before exercise, but at least 2 episodes with ΔHR ≥ 30 beats per minute after exercise. The phenomenon was transient as postural tachycardia returned to normal within 36 to 48 hr.

(iii) The normal postural response was defined as Stress Test Originated Phantom Perception (STOPP) based on original findings in GWI [30,31] where ΔHR was in the normal range of 12 ± 5 beats per minute and never exceeded 30 beats per minute.

2.5 Verbal working memory task

Subjects practiced the 0-back and 2-back working memory task in a mock scanner until they felt proficient [14,30,31]. In the scanner, subjects viewed an instruction panel stating 'REST' for 0.8 s followed by 19.2 s of a blank screen. The instruction '0-BACK' was viewed for 0.8 s followed by 1.2 s of a blank screen, and then a string of nine pseudorandomized letters (A, B, C, D) seen for 0.8 s each followed by 1.2 s of a blank screen per letter. Each time they saw a letter, subjects pressed the corresponding button on an MRI compatible fibre-optic four button box that was

used with both hands. After a second 'REST' period, they saw the instruction '2-BACK' and again viewed a string of nine letters. They had to view and remember the first two letters, then press the button for the first letter when they saw the third letter (i.e. '2 back', 4 s delay). The 2-back task continued for seven responses. This cycle was repeated five times.

2.6 MRI data acquisition, preprocessing and analysis

All structural and functional MRI data were acquired on a Siemens 3 T Tim Trio scanner located within the Center for Functional and Molecular Imaging at Georgetown University Medical Center equipped using a transmit-receive body coil and a commercial 12-element head coil array as described previously [14,30,31]. Parameters for structural 3D T1-weighted magnetization-prepared rapid acquisition with gradient echo (MPRAGE) images were: TE =2.52 ms, TR =1900ms, TI = 900ms, flip angle = 9, FOV = 250mm, 176 slices, slice resolution = 1.0 mm and voxel size 1x1x1mm. Images were processed in SPM12 [32]. fMRI data consisted of T2*-weighted gradient-echo planar images (EPIs) acquired during the n-back tasks. EPI data acquisition parameters were: TR/TE = 2500/30 ms, flip angle = 90, FoV =205 mm^2, matrix size =64x64, number of slices = 47, voxel size = 3.2 mm^3 (isotropic). Raw EPI data were preprocessed through the default pipeline within the CONN toolbox [33]. Briefly, steps were: (i) slice timing correction, (ii) subject motion estimation and correction, (iii) outlier detection for 'scrubbing' based on Artifact Detection Tools, (iv) co-registration with structural data, (v) segmentation and spatial normalization into standard Montreal Neurological Institute (MNI) space [34] and (vi) spatial smoothing with a stationary Gaussian filter of 6 mm full-width at half maximum (FWHM). Voxel size was 2.0 mm^3 (isotropic) after spatial normalization and conversion to Montreal Neurological Institute (MNI) space.

All within-subject and group-level image analyses were performed using the SPM12 software package [35]. After accounting for magnetic saturation by removing the first 6 scans, a paradigm based on the timing of events in the 2-back task (Fig. 1) was applied to preprocessed EPI data to sort individual subject scans into instruction, fixation, 0-back and 2-back bins. In the original analysis, one sample t-tests contrasted the BOLD signals from the 2-back and 0-back scans of each subject and included estimates of the translation (x, y, and z) and rotation (roll, pitch, and yaw) as covariates of non-interest. The resulting 2-back>0-back contrast maps from every subject were sorted into the control, GWI, and ME/CFS groups [14].

For this seed region re-analysis, we used the Harvard Ascending Arousal Network atlas [36,37] in the Lead DBS software package [38–40] to define regions of interest (ROI) for midbrain nuclei. The mean BOLD signal for each ROI was extracted from each subject's contrast map, re-centered to a population grand mean of 0, and the normalized data analyzed in the MarsBaR 0.44 toolbox [41,42]. MarsBaR output was the BOLD activation levels for the 2-back>0-back contrast condition in each midbrain nucleus for the control, ME/CFS and GWI groups on the pre- and post-exercise days.

Significant differences were determined by analysis of variance (ANOVA) with Tukey Honest Significant Difference or 2-tailed unpaired t-test with Bonferroni correction to correct for multiple comparisons, multivariate general linear modeling (mGLM) of relevant demographic and other independent variables, and partial correlation analysis performed in R and SPSS v27. mGLM of BOLD activation in each seed region began with Disease status (control, ME/CFS, GWI), Orthostatic status (START, STOPP, POTS), PTSD and gender as fixed factors with age, BMI and dolorimetry pressure thresholds as independent variables. Results were shown in the Supplementary Online Materials.

Nuclei that were significantly altered based on ANOVA were annotated onto the DBS Harvard anatomical figure (Figure 1) [36,37]. When viewed from the anterior (ventral) and posterior (dorsal) sides, the nuclei in the ascending arousal network suggest three layers in the coronal plane of MRI space that approximate their embryological origins [43–45]. Mesomere 1 contributes the superior (SC) and inferior colliculus (IC), midbrain reticular formation

(MRF) and periaqueductal grey (PAG). The second layer is derived from rhombomeres 0 and 1 and includes the superior dorsal raphe (DR), pontis oralis (PO), and more lateral penduculotegmental nuclei (PTN, formerly pedunculopontine nuclei or PPN) [43–45]. The ventral tegmental area (VTA) is anterior when viewed at this level but extends from the isthmus to diencephalon. The most caudal layer had midline DR and median raphe (MR), and bilateral locus coeruleus (LC) and parabrachial complex (PBC) that are derived from rhombomeres 1 and 2. Embryological origins do not align with the coronal MRI projection because of the marked ventral flexion of the midbrain during development and distortion of the original neural tube structures. Nuclei with significant differences by BOLD were manually highlighted by red outlines using Procreate software.

2.7 Supplementary Materials

Supplementary materials contain partial correlation coefficient analysis and multivariate general linear modeling results. Appendix A contains demographics, questionnaire and fMRI data.

3. Results

3.1 Demographics and questionnaires

As expected, there were more females in ME/CFS because of the known female predominance [6] and more male veterans with GWI from assault divisions deployed to the Persian Gulf (Table 1). PTSD was more common in GWI. Pressure induced pain sensitivity tested by dolorimetry was not different because male and female subjects were combined [25]. ME/CFS and GWI had comparable symptom scores that were significantly worse than controls with the exception of pain, SF36 Role Emotional and Mental Health that indicated more impairment in GWI than ME/CFS.

Preexercise BOLD values were assessed by self-reported demographics variables in a multivariable general linear model. The significant covariates were Orthostatic status, low back pain, depression, heart disease, gender and marital status (Table S1). The next iteration used the significant covariates as fixed factors and removed the other variables. Orthostatic status was the only variable to be significant (Table S2).

Table 1. Demographics. Questionnaire scores were significantly different between control, ME/CFS and GWI by ANOVA followed by Tukey Honest Significant Difference or Kruskal-Wallis and Mann Whitney tests (*) corrected for multiple comparisons. Mean ± SD. (Score ranges.)

	Control	ME/CFS	GWI	Control vs ME/CFS	Control vs GWI	ME/CFS vs GWI
N	31	36	78			
Age	43.2 ± 16.5	47.3 ± 13.1	47.1 ± 7.4			
Female *	38.7%	69.4%	21.8%	0.029		0.0001
BMI	28.3 ± 4.5	26.2 ± 5.6	29.4 ± 5.3			0.008
Dolorimetry (kg)	4.6 ± 2.4	3.9 ± 1.9	4.2 ± 2.1			
PTSD *	9.7%	13.8%	43.6%		0.001	0.004
Migraine *	13.3%	41.7%	63.5%		<0.00001	
Chalder Fatigue (0 to 33)	12.3 ± 5.4	23.3 ± 6.2	25.3 ± 4.7	<0.00001	<0.00001	
McGill pain (0 to 45)	3.6 ± 6.4	13.4 ± 11.0	23.8 ± 9.0	0.00012	<0.00001	<0.00001
CFS Severity Questionnaire (0 no symptom to 4 severe)						
Fatigue	1.3 ± 1.2	3.4 ± 0.8	3.5 ± 0.7	<0.00001	<0.00001	
Postexertional Malaise	0.6 ± 1.1	3.5 ± 0.8	3.4 ± 1.0	<0.00001	<0.00001	
Sleep	1.7 ± 1.4	3.2 ± 0.9	3.5 ± 0.8	<0.00001	<0.00001	
Memory, Concentration	1.2 ± 1.2	2.9 ± 0.9	3.1 ± 0.8	<0.00001	<0.00001	
Muscle Pain	0.6 ± 0.9	2.5 ± 1.3	3.1 ± 1.0	<0.00001	<0.00001	0.011
Joint Pain	0.8 ± 1.0	1.9 ± 1.4	3.2 ± 1.0	0.00026	<0.00001	<0.00001
Headache	1.0 ± 1.2	2.0 ± 1.3	2.7 ± 1.2	0.0022	<0.00001	0.025
Sore Throat	0.3 ± 0.7	1.0 ± 1.0	1.4 ± 1.2	0.031	0.000013	
Lymph Nodes	0.1 ± 0.4	1.0 ± 1.1	1.5 ± 1.3	0.004	<0.00001	
SF-36 (100 best to 0 worst)						
Physical Function	85.2 ± 24.2	44.4 ± 26.5	46.9 ± 24.6	<0.00001	<0.00001	
Role Physical	80.0 ± 36.8	10.0 ± 25.9	9.5 ± 24.8	<0.00001	<0.00001	
Bodily Pain	82.9 ± 20.1	47.2 ± 27.7	29.5 ± 18.3	<0.00001	<0.00001	0.00024
General Health	69.8 ± 22.8	33.2 ± 22.9	26.4 ± 19.2	<0.00001	<0.00001	
Vitality	60.2 ± 20.7	18.0 ± 15.9	16.6 ± 15.3	<0.00001	<0.00001	
Social Function	80.0 ± 25.1	30.7 ± 27.0	30.8 ± 24.5	<0.00001	<0.00001	
Role Emotional	86.7 ± 31.1	70.5 ± 44.1	30.7 ± 38.4		<0.00001	<0.00001
Mental Health	73.6 ± 16.8	67.8 ± 17.4	54.8 ± 22.3		0.000084	0.0056

3.2 Partial correlations

Partial correlations compared BOLD signal intensities for each node on Day 1, Day 2 and the delta with subjective questionnaires about CFS symptoms, SF36 domains, psychological and depression complaints with disease status, orthostatic status, gender, age and BMI as covariates. BOLD data were internally correlated within Day 1, Day 2 and delta, positively correlated between Day 2 and delta, and negatively correlated for Day 1 vs delta (Tables S3-S6). There were no significant correlations between subjective questionnaire data and objective preexercise or postexercise BOLD outcomes. The magnitudes of the significant correlations ($p < 0.05$ corrected) were low ($R < 0.4$ and $R > -0.4$ for the inversely scored SF36 domains,).

3.3 ANOVA

BOLD data were compared between groups defined by disease status (control, ME/CFS, GWI) and orthostatic status (START, STOPP, POTS) on the preexercise and postexercise study days. In the original study, dorsal midbrain activation was not different between groups before exercise [14].

Visual inspection suggested a trend for differences in BOLD between groups. Data from all seed region datapoints and subjects were contrasted between groups. Prior to exercise, ME/CFS had numerically lower BOLD (0.108 ± 0.032, mean $\pm$ 95%CI for n=504 datapoints = all regions of interest in all ME/CFS subjects) compared to control (0.297 ± 0.037, mean $\pm$ 95%CI, n=434 datapoints) and GWI (0.235 ± 0.024, mean $\pm$ 95%CI, n=1092 datapoints). This suggested reduced blood flow during the 2-back>0-back condition for ME/CFS at baseline.

In the ascending arousal network, control was significantly more activated than ME/CFS in bilateral PTN, L_PBC, and VTA, while GWI was higher than ME/CFS in L_PTN and L_PO (Figure 1, Table 2).

The only difference based on Orthostatic status on Day 1 was greater L_PBC activation in POTS than STOPP (Table S7).

Males (0.249 ± 0.023, mean $\pm$ 95%CI) showed a trend towards higher BOLD activation than females (0.167 ± 0.028, $p = 0.000012$ by unpaired 2-tailed t-test) when all nodes and subjects were assessed. The difference was significant for L_PO ($p = 0.046$) (Table S8).

Figure 1. Ascending arousal network before exercise. A. Anterior coronal view of the ventral midbrain with the plane of the posterior thalamus (Thal), pineal (P), superior (SC) and inferior colliculus (IC) as the backdrop and Basis Pontis as the "floor". Control had greater activation than ME/CFS for ventral tegmental area (VTA), bilateral pedunculotegmental nuclei (L and R PTN, formerly penduculopontine nuclei) and left parabrachial complex (L_PBC). GWI had greater activation than ME/CFS in the L_PTN and pontis oralis (L_PO). B. The view from the posterior (dorsal) side was oriented to show the coronal plane with centromedian/parafascicular nucleus (CEM/Pf), reticular nucleus (Ret) and central lateral nucleus (CL) of the thalamus and midbrain red nuclei (RN) for orientation. The red highlighted regions had significantly lower BOLD activation in ME/CFS than control and GWI groups (ANOVA and Tukey Honest Significant Difference $p < 0.05$) on the preexercise day (Table 2). Control had elevated BOLD compared to ME/CFS in VTA, bilateral PTN and L PBC, while GWI was higher than ME/CFS for left PTN and PO.
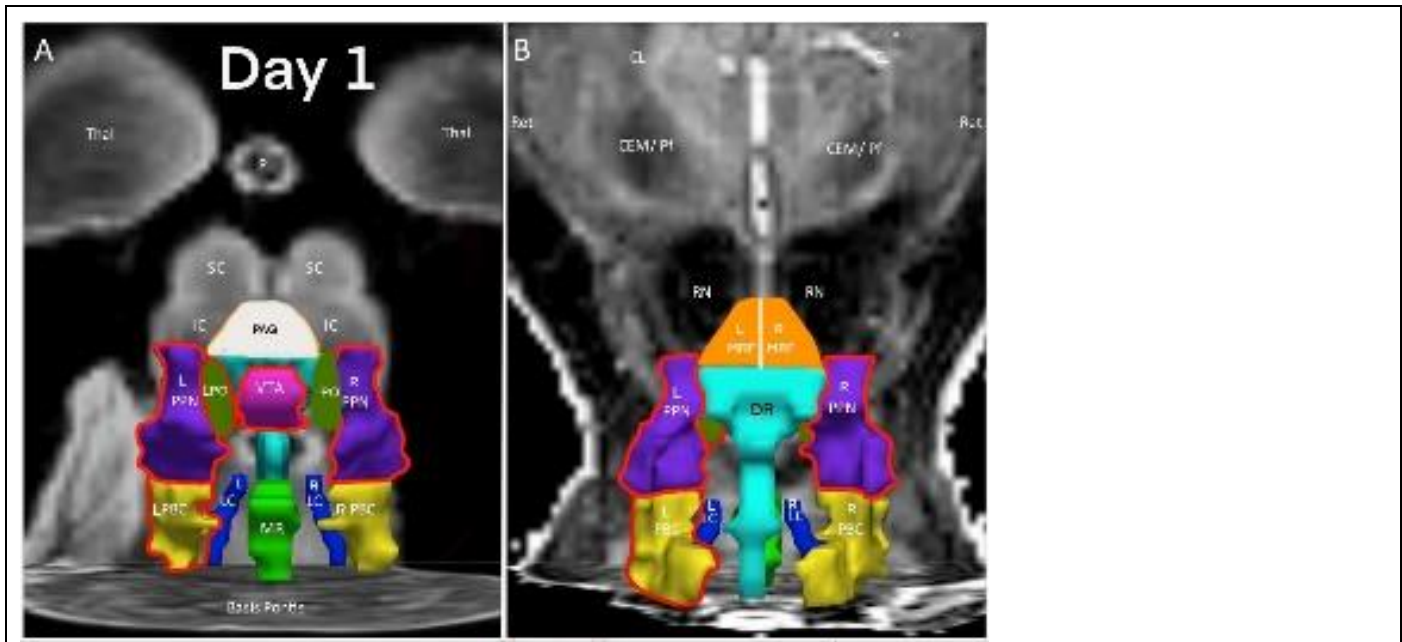
Table 2. ANOVA for Preexercise differences based on Disease status. BOLD signals for SC and GWI were numerically and statistically higher than ME/CFS (mean ± 95%CI, Tukey Honest Significant Difference, HSD). L and R indicate left and right, respectively. Hedges' g.

	Control	ME/CFS	GWI	Control > ME/CFS	GWI > ME/CFS
L_MRF	0.266 ± 0.164	0.123 ± 0.124	0.197 ± 0.103		
R_MRF	0.331 ± 0.181	0.216 ± 0.119	0.172 ± 0.103		
PAG	0.234 ± 0.145	0.061 ± 0.101	0.222 ± 0.089		
VTA	0.279 ± 0.127	0.103 ± 0.089	0.193 ± 0.061	p=0.034, g=0.58	
DR	0.275 ± 0.118	0.086 ± 0.103	0.233 ± 0.081		
MR	0.295 ± 0.124	0.131 ± 0.123	0.249 ± 0.087		
L_PO	0.263 ± 0.134	0.061 ± 0.126	0.257 ± 0.078		p=0.019, g=0.56
R_PO	0.290 ± 0.134	0.103 ± 0.136	0.252 ± 0.087		
L_PTN	0.316 ± 0.133	0.049 ± 0.101	0.264 ± 0.086	p=0.008, g=0.81	p=0.009, g=0.60
R_PTN	0.356 ± 0.126	0.125 ± 0.086	0.259 ± 0.080	p=0.014, g=0.78	
L_LC	0.309 ± 0.147	0.142 ± 0.158	0.226 ± 0.103		
R_LC	0.316 ± 0.181	0.143 ± 0.175	0.274 ± 0.126		
L_PBC	0.331 ± 0.139	0.078 ± 0.135	0.231 ± 0.092	p=0.029, g=0.65	
R_PBC	0.299 ± 0.148	0.088 ± 0.136	0.263 ± 0.101		

On the postexercise day, the relationship between groups became inverted as control (0.254 ± 0.035, mean ± 95%CI) and ME/CFS (0.260 ± 0.034) had significantly greater BOLD than GWI by ANOVA when all subjects and nodes were evaluated ($p < 10^{-9}$ by 2-tailed unpaired t-test with Bonferroni correction). After exertion, control was higher than GWI for bilateral MRF, VTA and R_PTN, while ME/CFS was greater than GWI for midline PAG, DR and MR and right MRF, LC and PBC (Table 3) (Figure 2).

There were no significant differences based on Orthostatic status following exercise (Table S9).

Gender was not a significant covariate for any region postexercise (Table S10). However, there was a general trend for females (0.123 ± 0.028, mean ± 95%CI) to have lower BOLD than males (0.197 ± 0.021, p = 0.000032 by 2-tailed unpaired t-test) when all nodes and subjects were compared.

Figure 2. Postexercise. Anterior (A) and posterior (B) views indicate that GWI had significantly lower BOLD signals than control in the L_MRF, R_MRF, VTA and R_PTN (outlined in red, Tukey Honest Significant Difference p < 0.05) (Table 3). GWI was significantly lower than ME/CFS in the R_MRF, PAG, DR, MR, R_PBC and R_ LC. There were no significant differences between the control and ME/CFS groups following exercise. Data were annotated as in Figure 1.

Table 3. ANOVA for postexercise by Disease status. Control and ME/CFS had significantly higher BOLD than GWI for several nodes. There was a general trend for GWI to have the lowest values in all regions. Mean ± 95%CI. Tukey Honest Significant Difference. Hedges' g.

	Control	ME/CFS	GWI	Control > GWI	ME/CFS > GWI
N	31	36	78		
L_MRF	0.341 ± 0.146	0.206 ± 0.147	0.073 ± 0.094	p=0.008, g=0.65	
R_MRF	0.364 ± 0.149	0.301 ± 0.138	0.077 ± 0.084	p=0.002, g=0.75	p=0.014, g=0.58
PAG	0.182 ± 0.141	0.271 ± 0.142	0.060 ± 0.078		p=0.016, g=0.57
VTA	0.252 ± 0.104	0.195 ± 0.099	0.078 ± 0.068	p=0.018, g=0.58	
DR	0.229 ± 0.113	0.265 ± 0.126	0.068 ± 0.070		p=0.009, g=0.60
MR	0.231 ± 0.115	0.285 ± 0.139	0.092 ± 0.075		p=0.018, g=0.54
L_PO	0.178 ± 0.125	0.241 ± 0.141	0.129 ± 0.082		
R_PO	0.248 ± 0.131	0.255 ± 0.135	0.101 ± 0.077		
L_PTN	0.233 ± 0.124	0.208 ± 0.127	0.093 ± 0.084		
R_PTN	0.308 ± 0.158	0.289 ± 0.104	0.126 ± 0.074	p=0.04, g=0.50	
L_LC	0.223 ± 0.144	0.274 ± 0.146	0.097 ± 0.089		
R_LC	0.276 ± 0.167	0.313 ± 0.150	0.077 ± 0.095		p=0.022, g=0.55

L_PBC	0.250 ± 0.133	0.250 ± 0.135	0.106 ± 0.091	
R_PBC	0.240 ± 0.135	0.289 ± 0.139	0.110 ± 0.076		p=0.041, g=0.50

ΔBOLD was positive in ME/CFS and negative in GWI indicating the significant dynamic effects caused by exercise in these two diseases (Figure 3). ME/CFS had higher increments than GWI for all seed regions except MRF and L_LC (Table 4). There were no differences between groups defined by Orthostatic status or gender.

Figure 3. Exercise effects. Incremental changes (ΔBOLD) were significantly larger (more positive) for ME/CFS than GWI (negative, diminished BOLD) in the midline PAG, VTA, DR and MR, bilateral PBC, PO and PTN, and R_LC (red highlighted nuclei) in the anterior (A) and posterior (B) views. Controls had no net changes. Data were from ANOVA analysis (Table 4).
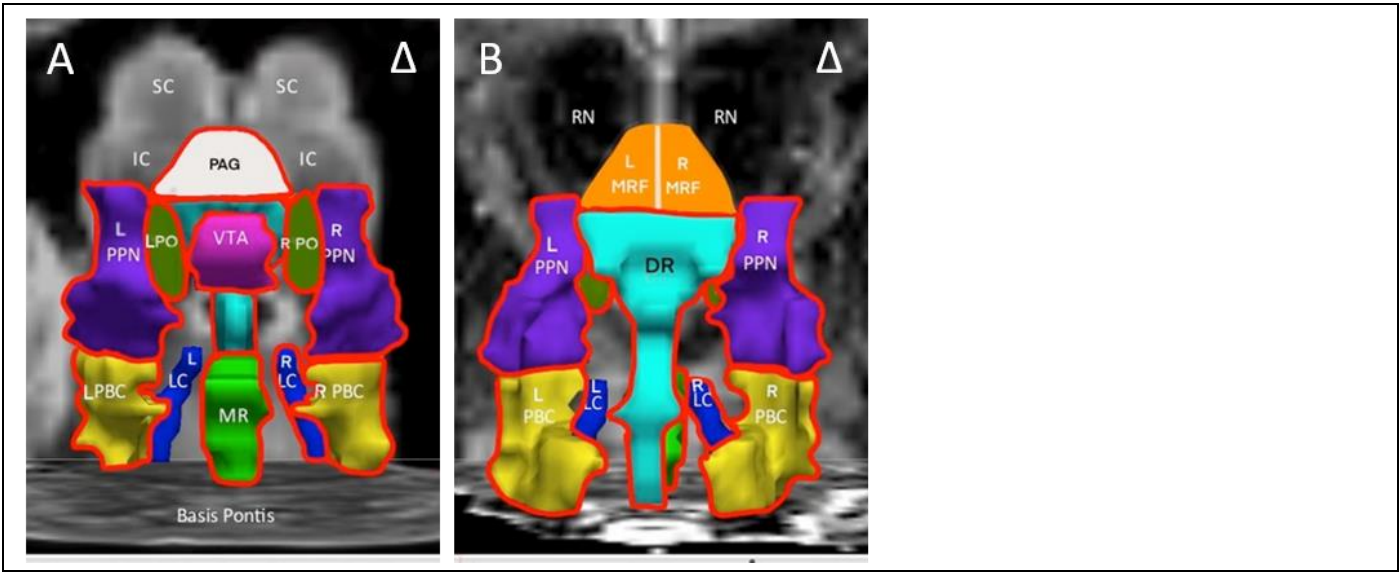

Table 4. ΔBOLD. Exercise-induced changes in the 2-back>0-back condition were analyzed. Incremental changes (ΔBOLD = postexercise minus preexercise) were assessed by ANOVA then corrected for multiple comparisons with Tukey Honest Significant Difference. ME/CFS had increased BOLD while GWI had diminished BOLD after exercise. Control had no net change and was not statistically different from either of the other groups. Mean ± 95%CI. Hedges' g.

	SC	ME/CFS	GWI	ME/CFS > GWI
N	31	36	78	
L_MRFΔ	0.075 ± 0.201	0.083 ± 0.163	-0.124 ± 0.129	
R_MRFΔ	0.033 ± 0.216	0.085 ± 0.171	-0.095 ± 0.122	
PAGΔ	-0.052 ± 0.167	0.210 ± 0.175	-0.162 ± 0.110	p=0.001, g=0.75
VTAΔ	-0.027 ± 0.138	0.092 ± 0.130	-0.114 ± 0.080	p=0.016, g=0.57
DRΔ	-0.046 ± 0.139	0.179 ± 0.159	-0.165 ± 0.103	p=0.001, g=0.75
MRΔ	-0.065 ± 0.153	0.155 ± 0.173	-0.157 ± 0.113	p=0.005, g=0.62
L_POΔ	-0.085 ± 0.167	0.180 ± 0.181	-0.128 ± 0.108	p=0.006, g=0.62
R_POΔ	-0.041 ± 0.178	0.152 ± 0.175	-0.151 ± 0.123	p=0.014, g=0.56

L_PTNΔ	-0.083 ± 0.154	0.158 ± 0.166	-0.171 ± 0.107	p=0.002, g=0.69
R_PTNΔ	-0.048 ± 0.168	0.164 ± 0.155	-0.132 ± 0.100	p=0.004, g=0.66
L_LCΔ	-0.086 ± 0.172	0.133 ± 0.201	-0.129 ± 0.130	
R_LCΔ	-0.039 ± 0.218	0.169 ± 0.220	-0.197 ± 0.165	p=0.024, g=0.52
L_PBCΔ	-0.081 ± 0.181	0.171 ± 0.198	-0.125 ± 0.128	p=0.025, g=0.52
R_PBCΔ	-0.059 ± 0.167	0.202 ± 0.195	-0.153 ± 0.129	p=0.005, g=0.62

3.4 Multivariate general linear models (mGLM)

mGLM of preexercise data used Disease, Orthostatic status, PTSD and gender as fixed factors with age, BMI and dolorimetry pressure thresholds as independent variables. Disease status was significant (Table S11). L_PTN activation was significantly lower in ME/CFS (0.018 ± 0.143, mean ± 95%CI) than control (0.326 ± 0.198, p = 0.047 univariate significance) and GWI (0.286 ± 0.127, p = 0.018) (Table S12). This was comparable to the ANOVA outcomes (Table 2). Orthostatic status was significant with L_PBC being significantly lower in STOPP (0.062 ± 0.121, mean ± 95%CI) than POTS (0.394 ± 0.219, p = 0.006 Tukey Honest Significant Difference) and START (0.397 ± 0.164, p = 0.034) (Table S13). Age, gender, PTSD, BMI and dolorimetry pressure thresholds were not significant covariates prior to exercise.

Postexercise mGLM evaluated the same fixed factors and independent variables. Disease status was significant after exercise. ME/CFS and Control had significantly higher BOLD activation than GWI in VTA, L_MRF and R_PTN (Table S14). Overall, ME/CFS was greater than GWI for all regions except L_PO, L_LC and L_PBC. More nodes were significant by mGLM than ANOVA (Table 3). Gender was significant for R_LC and R_PBC as males had greater BOLD activation than females after adjustment for the other variables (Table S15). Other significant interactions between disease, orthostatic and PTSD status, age and dolorimetry thresholds (kg) were detailed in the Supplementary Online Material (Tables S16-S18), but must be interpreted with caution because of concerns about the number of variables, sample sizes and potential overfitting of the data.

Incremental changes in BOLD between days (ΔBOLD) reinforced the differences between diseases found by ANOVA (Table 4). Estimated marginal means for Disease status bracketed zero for controls, were positive for ME/CFS and negative for GWI (Table S19).

Orthostatic status had a significant impact with higher activity for STOPP than START in R_LCΔ and bilateral PBCΔ (Table 5). The 95% confidence intervals for POTS and STOPP bracketed zero indicating no change after exercise. However, ΔBOLD was negative for the START group indicating a dynamic exercise-induced effect on brainstem activation in this phenotype. Effect sizes were small indicating that it may be difficult to reproduce the finding.

Table 5. Multivariate general linear model mGLM for ΔBOLD and Orthostatic status. Estimated marginal means for incremental changes (ΔBOLD) were evaluated with Disease and Orthostatic status, PTSD and gender as fixed factors and age, BMI and dolorimetry as independent variables. The model was significant for Disease status (Table S19) and Orthostatic status (Wilks' Lambda = 0.651, p = 0.019, Partial Eta Squared = 0.193) but not age, gender, PTSD, BMI or dolorimetry. 95% confidence intervals for POTS and STOPP bracketed zero indicating no net changes with exercise. The START phenotype was defined by exercise induced postural tachycardia and had reduced BOLD activation in the R_LC and bilateral PBC following exercise. The STOPP phenotype was the normal condition with no change in postural tachycardia. Mean ± 95%CI. Univariate significance. Hedges' g.
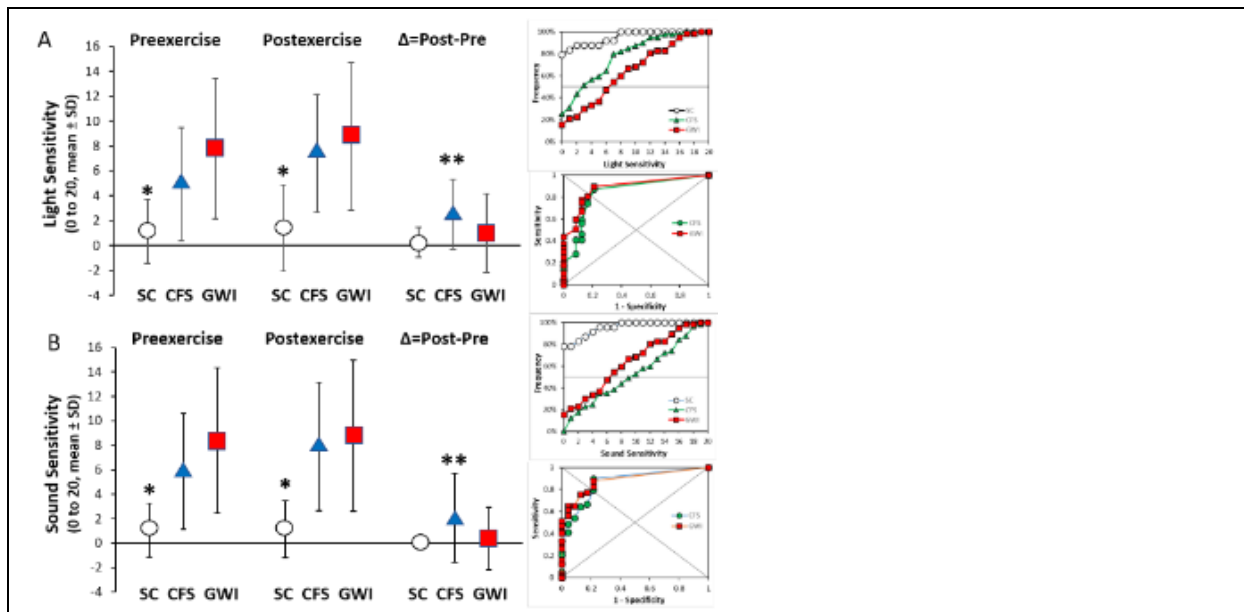
	POTS	START	STOPP	STOPP > START

L_MRFΔ	-0.009 ± 0.315	-0.093 ± 0.235	-0.063 ± 0.174	
R_MRFΔ	0.003 ± 0.319	-0.096 ± 0.239	-0.051 ± 0.177	
PAGΔ	0.004 ± 0.271	-0.140 ± 0.202	-0.026 ± 0.150	
VTAΔ	-0.062 ± 0.211	-0.146 ± 0.157	0.034 ± 0.116	
DRΔ	-0.090 ± 0.251	-0.153 ± 0.188	0.028 ± 0.139	
MRΔ	-0.164 ± 0.271	-0.166 ± 0.203	0.050 ± 0.150	
L_POΔ	-0.085 ± 0.274	-0.170 ± 0.204	0.061 ± 0.151	
R_POΔ	-0.048 ± 0.298	-0.259 ± 0.222	0.072 ± 0.164	
L_PTNΔ	-0.247 ± 0.256	-0.098 ± 0.191	-0.028 ± 0.142	
R_PTNΔ	-0.053 ± 0.260	-0.098 ± 0.194	-0.008 ± 0.144	
L_LCΔ	-0.219 ± 0.314	-0.232 ± 0.235	0.102 ± 0.174	
R_LCΔ	-0.070 ± 0.390	-0.329 ± 0.291	0.156 ± 0.216	p=0.035, g=0.42
L_PBCΔ	-0.304 ± 0.302	-0.268 ± 0.226	0.117 ± 0.167	p=0.029, g=0.29
R_PBCΔ	0.020 ± 0.316	-0.298 ± 0.236	0.110 ± 0.174	p=0.026, g=0.39

3.5 Light and sound sensitivity

Provocations with light and sound showed that sensory sensitivities were significantly worse in ME/CFS and GWI than control subjects before and after exercise (Figure 4). Frequency analysis of light sensitivity prior to exercise found scores of 0 or 1 out of 20 (no discomfort) in 83.3% of SC, 30.8% of ME/CFS and 21.1% of GWI. Exercise worsened light sensitivity in paired analysis for CFS (p = 2.7×10^{-6}) and GWI (p = 0.022), and sound sensitivity in ME/CFS (p = 0.037) by 2-tailed paired t-tests. The incremental changes (Δ) were significantly larger in ME/CFS than GWI and control (p < 0.044 by 2-tailed unpaired t-tests after Bonferroni corrections). Thresholds for significant sensitivities were ≥ 2 out of 20 by receiver operating characteristics indicating that visual and auditory hypersensitivity was common in ME/ CFS and GWI. The sound sensitivity and exaggerated startle responses were consistent with dysfunctional activity in the inferior colliculus [14].

Figure 4. Light sound sensitivity. Sensitivity was scored using modified Gracely box scores (range 0 to 20 points, mean ± SD). Light (A) and sound (B) were significantly worse for ME/CFS (blue triangles) and GWI (red squares) than SC (white circles) before and after exercise (* p<0.0001 by 2-tailed unpaired t-test after Bonferroni correction). The incremental changes (Δ) were larger in ME/CFS than SC and GWI (** p<0.044 by 2-tailed unpaired t-tests after Bonferroni correction). Paired changes were significant for light and sound in the ME/CFS group, and light in GWI. Receiver operating characteristics defined light sensitivity as ≥ 2 with 83.3% specificity and sensitivities of 74.4% for CFS and 80.7% for GWI (AUC = 0.884). Sound sensitivity of ≥ 2 had 78.3% specificity and sensitivities of 79.5% for ME/CFS and 82.5% for GWI (AUC=0.884).

4. Discussion

The BOLD data are the 2-back>0-back condition that contrasts the difficult high cognitive load continuous 2-back working memory task against the simple low cognitive load 0-back stimulus matching attention task [14]. The postexercise and incremental data reflect dynamic effects of exertion on cognition. Exercise caused changes in the 2-back>0-back condition that measures relative brain activation during the more difficult task. Specific effects on 2-back alone, 0-back alone, and 0-back>2-back condition were not assessed here.

The importance of the BOLD data is that there were differences in relative levels of regional blood flow into midbrain nuclei. ME/CFS and GWI had significant incremental changes following exercise whereas controls had no net changes. The changes were opposite in ME/CFS compared to GWI indicating distinct dynamic exercise-induced pathological consequences.

In the original study [14], dorsal midbrain activation was not different between groups prior to exercise. Reanalysis of the same BOLD data using the ascending arousal network nuclei and seed region approach found baseline differences with lower BOLD in ME/CFS than control and GWI (Figure 1, Table 2). Control had greater activation compared to ME/CFS in VTA, bilateral PTN and L PBC, while GWI was higher than ME/CFS for left PTN and PO.

The pedunculotegmental nuclei (PTN) had reduced activation in ME/CFS. This cholinergic nucleus has extensive efferents that release acetylcholine throughout the cerebrum to maintain wakefulness and induce attention. PTN assists in updating rapidly changing environmental information as required for our continuous 2-back task. Other functions and potential roles in disease dysfunction are discussed in the accompanying paper in this issue.

Exercise caused a significant dynamic switch in midbrain activation. Exercise caused an increase in BOLD in ME/CFS but decrease in GWI. Therefore, exercise had differential effects in ME/CFS compared to GWI. Following exercise, the control was greater than GWI in VTA, bilateral MRF and right PTN, while ME/CFS was higher than GWI for midline PAG, DR and MR, and right lateral MRF, PBC and LC (Figure 2, Table 3). The seed region approach was consistent with the activation found in the original 141 voxel region of interest that also included inferior colliculus (Figure S1) [14]. All 141 voxels in the region of interest analysis were contiguous, but the seed region approach assayed the net activation within the smaller volumes defined by the seeds.

After exercise, the midline nuclei and MRF had significantly lower BOLD activation in GWI than the other two groups.

PAG is integral to threat assessment and instantaneous responses. Detection of a threat activates the PAG and midbrain reticular formation and causes a transition from relaxed wakefulness to high general attention [46]. Active responses range from freeze with conscious tonic immobility if motion would lead to detection by a nearby predator; defensive approach to assess an ambiguous threat (modeled as rumination) [47,48]; flight via an escape route or to shelter; and defensive attack if the predator is within a dangerous distance and escape is not possible [49,50].

VTA is a dopaminergic nucleus that stimulates the locus coeruleus to promote wakefulness [51,52]. It has a role in reward situations and positive emotion. Dysfunction is associated with anhedonia.

DR and MR are serotoninergic nuclei that project to the limbic system during active stress, conflicts and anxiety [53,54]. They may initiate fight or flight decisions. MR participates in tolerance and coping strategies with aversive stimuli as well as arousal, wakefulness and long term memory.

In humans, the MRF is activated during the transition from a relaxed awake state to an attention-demanding state during reaction-time tasks [46] and during focused interrogation of threats while interpreting the proximity of danger (e.g. freezing in place) [55]. The caudal portion of the MRF extends into the cuneiform region that is correlated with cardiovascular dysfunction in ME/CFS [56,57].

The right locus coeruleus (R_LC) had significantly lower activation in GWI than ME/CFS after exercise. [58,59]. It is the predominant source of noradrenergic innervation in the brain [58,59]. Integrated PAG, amygdala and sensory information activate the LC to generate diffuse efferent outputs to the cerebrum and brainstem [60]. They act in an instantaneous fashion like a tripwire for immediate instinctual responses such as freeze-fight-flight, focused cerebral attention and sympathetic activation for immediate action. Inappropriate or dysfunctional activation contributes to anxiety and PTSD [61]. Atrophy of the right locus coeruleus was found at autopsy in veterans with PTSD [62].

Exercise caused no incremental changes in controls (ΔBOLD). However, exertion led to significant positive increments in ME/CFS and, by contrast, decreases in GWI. The dynamic changes were significantly different between ME/CFS and GWI for midline PAG, DR, MR and VTA, right LC, and bilateral PTN, PBC, and PO (Figure 3, Table 4). The opposite directions of change indicated that distinctly different pathological mechanisms that regulate midbrain blood flow and neurovascular coupling were modulated by exercise in the two diseases indicating that ME/CFS and GWI were excellent "illness controls" for each other.

Both ME/CFS and GWI had significant light and sound sensitivity (Figure 4). These sensations are monitored in the superior and inferior colliculus, respectively. The inferior colliculi are innervated by the ascending hindbrain auditory pathway (the lateral lemniscus), somatosensory pathways from the medulla, pons, and arousal nuclei [63]. The inferior colliculus participates in multimodal sensory perceptions, vestibulo-ocular reflex, predator aversion and escape, prey localization, social communication, analgesia and fear related behaviours. Sharp acoustic stimulation initiate the startle response with acutely accentuated attention and surveillance leading to visual and truncal orientation towards the sound, generalized hyperarousal and aversive behaviors [64–67]. Inferior colliculus is relevant to the light and sound sensitivity and heightened startle response in ME/CFS, GWI and veterans with PTSD [68].

The inferior colliculus is highly metabolically active and vulnerable to toxic injury [63]. ME/CFS have reduced cerebral blood flow during heads up tilt [69] and exercise [70] that may reduce oxygen supply to the susceptible inferior colliculus and lead to dorsal midbrain dysfunction. Exercise – induced lability of cerebral blood flow and neurovascular coupling may contribute to the dysfunctional BOLD patterns in the midbrain, insula and cerebellum vermis in GWI and ME/CFS [14] but with different mechanisms and statistically reciprocal outcomes in the two diseases.

The general linear models assessed the relative influences and interactions between Disease, Orthostatic and gender status on each day and adjusted for age, PTSD, BMI and dolorimetry thresholds. Outcomes for disease status were generally comparable to the ANOVA results.

Orthostatic status was significant before exercise as START and POTS had higher BOLD than STOPP in the left PBC (Table S13). POTS had postural tachycardia before and after exercise, while START were defined by exercise in-duced postural tachycardia [26,27,31]. After exercise, START had dynamic incremental depression of BOLD activation in the R_LC and bilateral PBC (Table 5) compared to the STOPP subjects who had no changes in BOLD or postural heart rate. Involvement of the locus coeruleus in START implicated exercise-induced autonomic dysfunction. The PBC interrogates pain and interoceptive visceral sensations then forwards the information to the PAG, thalamus, hypothal-amus, and amygdala for further processing [71]. PBC is recruited in states of malaise as an adaptive component of the sickness response [72], and so may participate in the experience of postexertional malaise.

Hedges' g ranged from 0.50 to 0.81 which suggests moderate to high effect sizes for replication of the significant results using the same protocols.

It is important to appreciate the limitations of these disease and exercise effects. The findings were inferred based on the seed regions extracted from the ascending arousal network. Coordinates of the seed regions may im-prove as newer standards are created [40]. The actual metric being compared is the 2-back>0-back differential activity during the difficult cognitive working memory task. The results may not be applicable to the resting state or other cognitive tasks. The neural and vascular responses combined to generate these data without providing insights into functional connectivity or molecular mechanisms. Brainstem motion may blur the borders of nuclei in this seed region approach. Therefore, we consider the results to be a general predictor of changes in BOLD for nuclei in the ascending arousal network that are congruent with the dorsal midbrain region of interest found in our previous study [14]. The results do suggest that significant differences will be found in future studies that specifically target these nuclei in ME/CFS and GWI when suitable sample sizes are compared and advanced motion correction algorithms are applied [73–76]. The most significant differences were induced by exercise with elevated BOLD in ME/CFS but reductions in GWI, and were most clearly exposed by comparison of ME/CFS versus GWI rather than differences from control sub-jects (Tables 5 and S19).

5. Conclusion

The seed region approach based on the ascending arousal network extended our previous finding of exercise induced changes in BOLD during a high cognitive load 2-back working memory task. The salient findings were sig-nificantly lower BOLD in the midbrain at baseline in ME/CFS compared to GWI and control, and significant dynamic changes after exercise with elevation of BOLD in ME/CFS but reduction in GWI. Review of the functions of midbrain nuclei provides a fresh perspective on potential neural pathologies affecting inferior colliculus (startle), oculomotor and visual systems, PAG, MRF and other nuclei for threat assessment, anxiety, negative emotion, pain and tenderness and other aspects of the ME/CFS and GWI clinical experiences. The data provide an initial framework to power future studies of postexertional malaise and midbrain dysfunction.

Conflict of Interest

Supplementary Materials The following are available online at www.mdpi.com/xxx/s1, Figure S1: Midbrain Region of Interest and Tables S1-S19.

Institutional Review Board Statement: The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the Institutional Review Board (or Ethics Committee) of GEORGETOWN UNIVERSITY 2013-0943, 2015-0579, 2009-229, 29th June, 2021, and Department of Defense Human Research Protection Office Log Number A-18749, 16th September, 2019.

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: Primary data are attached in Appendix A.

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