Characterising the gut microbiome in veterans with Gulf War Illness: a protocol for a longitudinal, prospective cohort study

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ABSTRACT

Introduction Approximately 25%–35% of the 1991 Gulf War Veteran population report symptoms consistent with Gulf War Illness (GWI), a chronic, multi-symptom illness characterised by fatigue, pain, irritable bowel syndrome and problems with cognitive function. GWI is a disabling problem for Gulf War Veterans, and there remains a critical need to identify innovative, novel therapies. Gut microbiota perturbation plays a key role in the symptomatology of other chronic multi-symptom illnesses, including myalgic encephalomyelitis/chronic fatigue syndrome (ME/CFS). Given similarities between ME/CFS and GWI and the presence of gastrointestinal disorders in GWI patients, Veterans with GWI may also have gut abnormalities like those seen with ME/CFS. In this longitudinal cohort study, we are comparing the diversity (structure) and the metagenomes (function) of the gut microbiome between Gulf War Veterans with and without GWI. If we find differences in Veterans with GWI, the microbiome could be a target for therapeutic intervention to alleviate GWI symptoms.

Methods and analysis Participants answer questions about diet, exercise and lifestyle factors. Participants also complete a questionnaire (based on the Kansas case definition of GWI) regarding their medical history and symptoms; we use this questionnaire to group participants into GWI versus healthy control cohorts. We plan to enrol 52 deployed Gulf War Veterans: 26 with GWI and 26 healthy controls. Participants provide stool and saliva samples weekly for an 8-week period for microbiome analyses. Participants also provide blood samples at the beginning and end of this period, which we will use to compare measures of inflammation markers between the groups.

Ethics and dissemination The protocol was approved by the University of Wisconsin-Madison Health Sciences Institutional Review Board and the William S. Middleton Memorial Veterans Hospital Research and Development Committee. Results of this study will be submitted for publication in a peer-reviewed journal.

INTRODUCTION

Based on estimates by the Gulf War Research Advisory Committee and the 2010 Institute of Medicine report, approximately 25%–35% of the 1991 Gulf War Veteran population report symptoms consistent with Gulf War Illness (GWI).1–3 GWI is a chronic, multi-symptom illness characterised by fatigue, pain, irritable bowel syndrome and problems with cognitive function. GWI is a disabling problem for many Gulf War Veterans, and there have been only modest advances in the treatment for GWI. There remains a critical need for identification of innovative, novel therapies.4

Gut microbiota perturbation plays a key role in the symptomatology of chronic multi-symptom illnesses similar to GWI, such as myalgic encephalomyelitis/chronic fatigue syndrome (ME/CFS)5–7 Although GWI and ME/CFS are considered distinct conditions, both of them are characterised by similar complex, multi-organ chronic signs and symptoms. Moreover, Gulf War Veterans are more likely to meet the case definition for ME/CFS than non-deployed era Veterans.8 Given the
overlap and similarity of symptoms, it is possible that gut microbiota abnormalities as described in ME/CFS may also be responsible for the symptomatology in GWI.\textsuperscript{9,10}

Data suggest that individuals with GWI have an increased prevalence of gastrointestinal symptoms such as diarrhoea and abdominal pain and cramps,\textsuperscript{11} which may result in or be caused by intestinal dysbiosis (i.e., alteration in the microbial community), low grade inflammation and immune activation.\textsuperscript{12} Given the body’s largest immune reservoir resides in the gastrointestinal tract, interventions such as probiotics, dietary interventions or faecal microbiome transplants could favourably alter the gut microbiome, improve mucosal barrier function and decrease proinflammatory cytokines. Overall, interventions targeting the gut microbiome have the potential to positively influence not only gastrointestinal symptoms but also non-gastrointestinal symptoms in GWI Veterans. However, an understanding of the structure and function of the gut microbiome in Veterans with GWI is essential to the development of such interventions. Based on our review of the literature, the gut microbiome in Veterans with GWI and its relationship to the symptomatology of GWI have not yet been reported.

Here, we are characterising the gut microbiome of Gulf War Veterans with GWI compared with healthy Gulf War Veterans without GWI. This work is ongoing (2018–2020) and will be among the first to examine gut microbiome in Gulf War Veterans and will advance our understanding of the pathophysiology of GWI.

**STUDY AIMS AND HYPOTHESES**

In this longitudinal, prospective cohort study, we aim to compare the gut microbiomes between deployed Gulf War Veterans: 26 with GWI and 26 without GWI. Figure 1 provides an overview of the study methods. We hypothesise that the gut microbiome will be more diverse and will have a higher abundance of Bacteroidetes (rather than Firmicutes) in healthy Gulf War Veterans as compared with Veterans with GWI.

We also aim to evaluate the functional characteristics of the gut microbiome by comparing metagenomes between Gulf War Veterans with and without GWI. We hypothesise that metagenomes will differ in the enrichment of genes for predicted pathways between these groups. We will especially focus on microbiome pathways producing metabolites involved in symptoms frequently seen in GWI. For example, butyrate—a short-chain fatty acid metabolite produced by gut bacteria—has been associated with gastrointestinal symptoms.\textsuperscript{13,14} Recent work in a mouse model of GWI has also shown that butyrate priming reduces metabolic abnormalities seen in GWI, for example, by restoring levels of gut tight junction proteins and reducing activation of a Toll-like receptor (TLR4; a mediator of inflammation).\textsuperscript{15}

To test our hypotheses, we will extract, sequence and analyse genomic DNA from serial stool samples of Gulf War Veterans. We will compare the alpha and beta diversity of the microbiota between groups. We will also perform a shotgun metagenomics analysis on the collected samples.

To provide additional context to the gut microbiome results, we will also analyse the diversity of the oral microbiota through the collection and analysis of serial saliva samples. We will also collect blood samples to measure inflammation (C-reactive protein levels). We anticipate Veterans with GWI will have higher levels of C-reactive protein and inflammatory markers as compared with Gulf War Veterans without GWI.

**METHODS AND ANALYSIS**

**Recruitment and consent**

Participants are recruited primarily through invitation letters sent to local Gulf War Veterans identified from VA databases. Additional advertising strategies, including fliers, social media posts and local newspaper ads are also used. Veterans interested in participating complete a screening by telephone, and those who are eligible and interested in participating attend an in-person enrolment visit. Participants provide weekly stool and saliva samples over an 8-week period and provide blood samples at Weeks 1 and 8.

**Eligibility criteria and cohort assignment**

Inclusion and exclusion criteria for participants are listed in boxes 1 and 2. We are utilising a modified Kansas case...
definition to determine eligibility and cohort assignment for this study. Veterans are excluded from the study if they report active, serious medical/psychological diagnoses which account for GWI-like symptoms or prevent accurate GWI-like symptom reporting. Veterans with well-controlled (or in remission) symptoms and who have not been hospitalised due to these conditions up to 5 years prior to enrolment may be eligible to participate.

We also use the Kansas case definition to group participants into two cohorts: deployed Gulf War Veterans with GWI and deployed healthy Gulf War Veterans. During enrolment, participants use the Kansas case definition questionnaire to self-report the severity of symptoms they experience within the six GWI symptom domains: fatigue, neurological/cognitive/mood, pain, gastrointestinal, respiratory and skin. For each symptom, participants select whether the symptom has no, mild, moderate or severe impacts on their daily lives. To be grouped into the GWI cohort, participants must endorse one or more moderate to severe symptoms in three or more of the six domains. One of the domains with moderate/severe symptoms must be fatigue. Participants who do not meet these criteria are grouped into the healthy Gulf War Veteran control cohort.

## Enrolment

During the enrolment visit, participants provide medical history and health and lifestyle factors which can impact the microbiome. Participants complete an in-house assessment addressing living situation (eg, type of home, rural/urban), health issues (eg, recent stays in long-term care facilities), oral health practices, current medication use and use of alcohol and tobacco. The participants also complete validated instruments: Holmes-Rahe Stress Inventory, Multidimensional Fatigue Inventory (MFI-20),International Physical Activity Questionnaire—short form, and National Cancer Institute’s Diet History Questionnaire III.

Participants report on branch of service during their Gulf War deployment and chemical, biological and pharmaceutical exposures from their deployment using an in-house questionnaire based on the VA list of Gulf War exposures. Participants also complete the Kansas case definition assessment as described earlier.

### Sample collection

#### Stool and saliva samples

Participants are provided with collection kits, shipping materials and instructions to self-collect and mail their stool and saliva samples. Stool sample collection kits include a pair of clean gloves, stool collection tub, wooden tongue depressor, biohazard bag and pre-labeled sterile specimen cup. Saliva sample collection kits include a CryoELITE Cryogenic Storage Vial (Wheaton) saliva collection tube and a biohazard bag. The stool and saliva collection containers do not contain preservative agents. The participants are provided with ice packs and an insulated shipping container.

Participants collect at least five tablespoons of stool and 1–3.5 mL of saliva samples at their convenience once per week for 8 weeks and mail samples to the study team in the supplied shipping containers.

Participants have a window of 5 days each week to collect samples. The participant records the date and time of collection on the label provided for each sample and refrigerates the samples on collection (no freezing). Within 24 hours of collection, the participant ships the samples overnight on completely frozen ice packs. Samples received more than 72 hours after the date noted on the sample label are rejected and are not included in microbiome analyses.

A member of the study team conducts a weekly phone check-in with each participant to ensure timely sample collection and to ask about any antibiotic/probiotic use or gastrointestinal symptoms over the week. This information is noted on study case report forms.

#### Blood samples

Two whole blood samples are drawn by venipuncture at enrolment and at the 8-week point: one sample is drawn into a 2 mL sodium heparin Vacuette (454302, Greiner Bio-one) and one sample is drawn into a 4 mL Vacuette containing potassium ethylenediaminetetraacetic acid (EDTA) (454021, Greiner Bio-one). Whole blood drawn in sodium heparin Vacuettes are processed immediately on collection for phagocytosis assay. Whole blood from
the EDTA Vacuette are centrifuged at 1690×g for 15 min at room temperature immediately on collection, after which plasma is aspirated and aliquoted into cryovials to be stored at −80°C.

Microbiome analyses

The microbiome analyses methods follow the standard protocol developed by our research group, as previously published. Both 16S rRNA-targeted amplicon sequencing of the V4 region and shotgun metagenomics will be performed on the stool and saliva samples to address the aims of this project. 16S rRNA sequencing will be done on an Illumina MiSeq. A subset of the samples will be analysed using shotgun metagenomics on an Illumina HiSeq. All sequencing will be carried out at the University of Wisconsin-Madison Biotechnology Center. DNA extraction methods for each sample type are described below. For 16S rRNA sequencing, purified DNA is normalised to 5 ng/µL, amplified using PCR with barcoded primers for the 16S V4 region and sequenced using 2×250 nt reads.

Stool total genomic DNA extraction

0.8–0.1 g of stool is added to a bead-beat tube containing approximately 300 mg of 1.0 mm diameter zirconia/silica beads and vortexed to homogenise the stool. The sample is then centrifuged for 15 min at 4°C at 500×g. A total of 800 µL of 2× Sodium Chloride-Tris-EDTA (STE) buffer is added to the supernatant and up to 1000 µL is transferred to a new bead-beat tube containing 0.1 mm diameter zirconia/silica beads and one 4 mm stainless steel bead. For chemical lysis, 115 µL of an enzymatic cocktail containing 50 µL lysozyme (10 mg/mL), 10 µL mutanolysin (1 mg/mL), 5 µL lysostaphin (5 mg/mL) and 50 µL 20% sodium dodecyl sulfate is added to each tube. Additionally, 700 µL phenol:chloroform:isoamyl alcohol is added to the sample. Bead-beat tubes are then vortexed and incubated at 56°C for 30 min. For mechanical lysis, bead-beat tubes are vortexed, placed in a Mini-BeadBeater-24 (Cat 112011, Biospec Products, Bartlesville, Oklahoma, USA) and beat for 3 min. Tubes are centrifuged at 16 000×g for 10 min at 4°C. The top aqueous layer is transferred to a clean 2 mL tube, washed with 500 µL of an enzymatic cocktail containing 10 µL lysozyme (10 mg/mL), 50 µL mutanolysin (1 mg/mL), 50 µL lysostaphin (5 mg/mL) and 50 µL 20% sodium dodecyl sulfate is added to each tube. Additionally, 700 µL phenol:chloroform:isoamyl alcohol is added to the sample. Bead-beat tubes are then vortexed and incubated at 56°C for 30 min. For mechanical lysis, bead-beat tubes are vortexed, placed in a Mini-BeadBeater-24 (Cat 112011, Biospec Products, Bartlesville, Oklahoma, USA) and beat for 3 min. Tubes are centrifuged at 16 000×g for 10 min at 4°C. The top aqueous layer is transferred to a clean 2 mL tube, washed with an additional 500 µL phenol:chloroform:isoamyl alcohol and vortexed. The sample is then centrifuged at 16 000×g for 10 min at 4°C. The phenol:chloroform:isoamyl alcohol wash is repeated between 2 and 10 times to remove impurities from the sample until the aqueous layer is clean. The top aqueous layer is then transferred to a clean 2 mL microcentrifuge tube containing 70 µL of 3 M sodium acetate and 700 µL isopropanol. The samples are inverted several times and subsequently incubated at −20°C for 30 min to 1 hour. Each sample is centrifuged at 16 000×g (4°C) for 20 min to collect the DNA pellet, which is then washed with 500 µL cold 70% ethanol. The ethanol wash is repeated, and sample DNA pellets are dried for 5 min using a Savant SpeedVac (DNA120-230, Thermo Scientific, Waltham, Massachusetts, USA). Finally, dried DNA pellets are resuspended in 100 µL TE buffer and stored overnight at 4°C or at 37°C for 1 hour to dissolve the DNA pellet. Samples are then purified using NucleoSpin Gel and PCR cleanup kit according to manufacturer’s directions (Macherey-Nagel, Germany) and eluted in 40 µL TE buffer. DNA is quantified using PicoGreen in a microplate reader (BioTek Instruments) and stored long term at −80°C.

Saliva total genomic DNA extraction

1× PBS is added to the saliva tube to reach a volume of 1 mL which was then transferred to a bead-beat tube containing 0.1 mm diameter zirconia/silica beads and one 4 mm stainless steel bead. For chemical lysis, 25 µL of an enzymatic cocktail containing 5 µL lysozyme (10 mg/mL), 15 µL mutanolysin (1 mg/mL), 5 µL lysostaphin (5 mg/mL) is added to each sample, vortexed and incubated at 37°C for 30 min. Following incubation, 60 µL of an enzymatic cocktail containing 10 µL Proteinase K (20 µg/mL) and 50 µL 10% sodium dodecyl sulfate is added to each tube for a second chemical lysis step. The samples are then vortexed and incubated at 55°C for 45 min. Mechanical lysis is carried out by bead-beating the samples on high on a Mini-BeadBeater-24 (Cat 112011, Biospec Products) for 3 min. Samples are centrifuged at 16 000×g for 3 min at 4°C. The top aqueous layer is transferred to a sterile 2 mL tube and 0.1 vol of 3 M sodium acetate was added along with 0.6 vol isopropanol and mixed by inversion. The samples are incubated at −20°C for 30 min to 1 hour. Each sample is centrifuged at 16 000×g (4°C) for 20 min to collect the DNA pellet, which is then washed with 500 µL cold 70% ethanol. The ethanol wash is repeated, and sample DNA pellets are dried for 5 min using a Savant SpeedVac (DNA120-230, Thermo Scientific). Finally, dried DNA pellets are resuspended in 100 µL TE buffer and stored overnight at 4°C or at 37°C for 1 hour to dissolve the DNA pellet. Samples are then purified using NucleoSpin Gel and PCR cleanup kit according to manufacturer’s directions (Macherey-Nagel) and eluted in 40 µL TE buffer. DNA is quantified using PicoGreen in a microplate reader (BioTek Instruments) and stored long term at −80°C.

Blood analyses

C-reactive protein analyses

Human C-reactive protein is detected by analysis with a commercially available ELISA kit (Human C-Reactive Protein ELISA Kit, KHA0031, Invitrogen). In short, a whole blood sample collected in a 4 mL potassium EDTA vacutainer is centrifuged at 1690×g for 15 min. Plasma from the sample is aliquoted into three cryogenic vials and stored at −80°C. Samples are batch processed for C-reactive protein using a commercially available ELISA kit (Human C-Reactive Protein ELISA Kit, KHA0031, Invitrogen) and plates are read with a BioTek Synergy HT microplate reader using BioTek Gen5 V.2.09 software.
Flow cytometry analyses

Results from flow cytometry acquisition are analysed using FlowJo V.10 software. Monocytes and granulocytes are initially gated via Side Scatter (SSC) versus Forward Scatter (FSC). Those populations are further analysed via SSC versus fluorescence (BLA-1) where the fluorescent populations were further gated. The gated fluorescent populations indicated activity resulting from the phagocytosis assay (pHrodo Green E. coli BioParticles Phagocytosis Kit, P3538I, Invitrogen) where increased fluorescence is indicative of increased cell death. ThermoFishcer Attune Next Flow Cytometers used to collect phagocytosis assay data are located at the University of Wisconsin Carbone Cancer Center Flow Lab. Gating of this population using SSC versus BLA-1 allows for the generation of a curve measuring counts against emission wavelength. Curves for each condition are averaged together; SDs are calculated and p values comparison is computed.

Sample size determination and statistical analysis

Using data from other studies of the gut microbiome, including studies on ME/CFS, we performed simulation-based estimates of power using the micropower package in R. A proposed sample size of 26 healthy Gulf War Veterans and 26 Veterans with GWI will provide 80% power to detect an effect size (omega-squared) of 0.042 for the weighted UNIFRAC using a PERMANOVA test. The detectable effect size is comparable with, and more conservative than, the effect sizes observed in few studies that exist of the gut microbiome in ME/CFS, including one by our research group in which we compared the gut microbiota of 10 patients with ME/CFS to 10 healthy controls.

For 16S rRNA sequencing, raw sequences will be processed using VSEARCH and mothur. Contigs will be compiled and low-quality reads will be removed. Chimeras and short reads will be removed using VSEARCH. The GreenGenes database will be used to assign operational taxonomic units to the genus level (97% similarity) where possible. Alpha diversity will be assessed using the Shannon and Inverse Simpson’s diversity indices and richness will be calculated using the Abundance-based Coverage Estimator (ACE) and Chaoc1 indices. Beta diversity will be assessed to provide insights into community structure.

Shotgun metagenomics analysis will be conducted on a subset of samples. Phylogenetic binning will be conducted to determine if there is enrichment for specific taxa between samples. Baseline annotations will then be determined using a variety of databases to reconstruct metabolic and physiological pathways. Enrichment analyses will be conducted between samples to determine if specific sets of genes are over-represented or under-represented. These data will then be correlated to the other contextual data collected from each individual. We will focus our analyses on known pathways predicted to be implicated in modulating differences between Veterans with and without GWI, such as increases in short-chain fatty acids like butyrate and acetate.

To account for covariates, mixed effects linear regression will be used to estimate mean species diversity as a function of disease status (GWI vs control) as well as contextual factors including age, dietary fibre intake, tobacco and alcohol use and living situation. Confounders such as branch of Gulf War service and self-reported Gulf War chemical and biological exposures will also be controlled for using mixed linear regression models. C-reactive protein levels will be compared between Gulf War Veterans with GWI and healthy Veterans using non-parametric tests. We expect to find that levels of inflammatory markers will be higher in Veterans with GWI than in the control group.

Patient and public involvement

We have worked with Veteran representatives and stakeholders, including local Veterans Service Officers, to share the study to Veterans who may be interested in participating. We also solicited feedback from these stakeholders into our recruitment materials and strategies to ensure communications were appropriate for this Veteran population. We plan to disseminate results back to the Veteran community through venues such as Veteran conferences/meetings, working with local Veteran Service Organizations and working with the Madison VA Patient Education Resource Center.

DISCUSSION

GWI is a disabling problem for many Gulf War Veterans. To date, there has been little information on the pathology of the disease. There are no scientifically validated, effective treatments for GWI, though there have been modest advances in GWI treatments. Thus, the need for innovative treatment strategies remains critical. The gut microbiome presents a novel target for investigation. The high prevalence of gastrointestinal dysfunction in Gulf War Veterans in the context of literature demonstrating intestinal dysbiosis in patients with ME/CFS, another complex condition, suggests that alterations in the gut bacteria could play a role in the pathophysiology of GWI. This study is among the first to examine the gut microbiome structure and function in Gulf War Veterans.

The brain–gut axis—biochemical signalling between the gastrointestinal tract and the central nervous system—is increasingly recognised as important to health. The gut microbiome is known to impact the production, expression and turnover of neurotransmitters such as serotonin. These neurotransmitters affect cognitive and functional nervous system processes such as behaviour, mood and fatigue. The microbiome also affects the intestinal barrier and tight junction integrity, and bacterial metabolites influence mucosal immune regulation. Previous studies have shown chemical exposures common to the Gulf War (eg, the anti-nerve agent pyridostigmine bromide) can be used to create a mouse...
model of GWI. Through analyses of inflammatory markers as well as the structure and function of the microbiome in Veterans with GWI, our study aims to improve our understanding of the brain–gut axis and the role of the microbiome in GWI in humans.

Preliminary data from studies in mice and humans with ME/CFS have found a potential benefit in the use of probiotics to ameliorate fatigue symptoms. Initial studies have also found improvement in ME/CFS symptoms through the use of faecal microbiota transplant; additional studies are ongoing into the use of faecal microbiota transplant to treat ME/CFS symptoms.

If we find that the gut microbiome is altered in Veterans with GWI, then probiotics, dietary intervention or faecal microbiota transplant may represent low-cost, low-risk treatment options for GWI symptoms. At the conclusion of this study, we expect to be positioned to pursue a clinical trial to investigate interventions targeting the gut microbiome as a way to alleviate symptoms in Veterans with GWI.

ETHICS AND DISSEMINATION

The study protocol was approved on 12 January 2018 and this manuscript reports on the most updated version of the protocol approved on 29 August 2018. All participants are informed prior to enrolment that participation is completely voluntary, that they can withdraw from participation at any time and that their decision to participate or not will not affect their healthcare in any way.

On completion of the study, we will present the results at major scientific conferences and will publish the results in a peer-reviewed journal.

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Contributors
NS, DBC, DR, GS and RG conceived of the study concept and design. JAK drafted the overall protocol, with critical input from CS and KB for recruitment, screening, enrolment and data collection. AEK drafted protocol sections for DNA extraction, microbiome analyses and statistical analyses; NP-B, LW and ND for blood analyses and KW for flow cytometry. All authors provided critical feedback and approved the final version of the manuscript.

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Disclaimer
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None declared.

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