Longitudinal Assessment of Health Symptoms in Relation to Neurotoxicant Exposures in 1991 Gulf War Veterans

The Ft. Devens Cohort

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Objective: This analysis examined the relationship between Gulf War (GW) exposures and health symptoms reported in three time periods over 20 years in Ft. Devens Cohort veterans. Methods: Repeated logistic regression models examined the association of exposures and health symptoms over time. Models included baseline age, active duty status, post-traumatic stress disorder status, sex, and time since deployment as covariates. Results: Exposure to tent heaters was associated with increased odds of crying easily and muscle twitching. Exposure to pyridostigmine bromide (PB) pills was associated with increased odds of depression and fatigue. Exposure to the Khamsiyah sarin plume was associated with increased odds of trouble concentrating and crying easily. Conclusion: This longitudinal analysis demonstrated an association between neurotoxicant exposures and increased odds of cognitive/mood, fatigue, and neurological symptoms. In addition, most symptoms increased over time since deployment regardless of exposure.

Keywords: gulf war, gulf war illness, health symptoms, longitudinal analysis, neurotoxicants exposures, veterans

Military personnel deployed during the 1990 to 1991 Gulf War (GW) were exposed to unique environmental hazards in theatre, including pesticides, petrochemicals, debris from SCUD missiles, smoke from oil well fires, depleted uranium, prophylactic medications such as pyridostigmine bromide (PB) pills, and the chemical warfare agents sarin/cyclosarin. Previous findings from cross-sectional data suggest that GW veterans self-reporting exposure to smoke from oil well fires, pesticides, debris from SCUD missiles, hearing chemical weapon alerts, and using PB anti-nerve gas pills, were more likely to report cognitive dysfunction (e.g., changes in memory), depressive symptoms, and neurological complaints (e.g., headaches). In addition, pulmonary symptoms such as asthma, chronic bronchitis, cough, and shortness of breath, were more likely to be reported among GW veterans with exposure to combustion by-products from oil well fires, propane fuel emissions from tent heaters, and chemical weapons. Several studies have now found that these symptoms may stem from chronic neuroinflammatory or neuroimmune responses following neurotoxicant exposures.

Follow-up studies among deployed GW veterans found that the number and severity of health symptoms remained stable or slightly worsened over follow-up periods of varying lengths (4–18 years), indicating neither improvement nor significant progression of symptoms. However, few longitudinal studies have examined the relationship over time between GW exposures and health symptoms in the same group. As GW veterans age, it is becoming increasingly important to evaluate these symptoms over time to study how neurotoxicants may influence patterns of aging.

The Ft. Devens cohort (FDC) is a population of former US Army Active, Reserve, and National Guard GW veterans followed prospectively through a series of surveys since their return from deployment in the Persian Gulf in 1991. These veterans were not recruited on the basis of symptom reporting but based on their military units. The findings from the FDC included early documentation of the most common health symptoms, cognitive decrements in environmentally exposed GW veterans, and differences in structural neuroimaging, including lower white matter volumes, as well as the effects of mild traumatic brain injury (mTBI) on GWI. To our knowledge, the present study is the first to utilize data from three surveys with the same veterans over a 20-year period to examine the relationship between self-reported GW deployment exposures and health symptoms over time.

Learning Objectives

- Summarize previous findings on the environmental hazards and symptoms reported by Gulf War veterans, including previous follow-up studies.
- Discuss the new 20-year follow-up findings in the Ft. Devens Cohort of Gulf War veterans, including specific symptoms associated with specific exposures.
- Discuss the study implications for ongoing care and monitoring of Gulf War veterans.
METHODS

Participants and Surveys

In the Spring of 1991 (Baseline), Active Duty, Reserve, and National Guard Army personnel who returned from deployment to the Persian Gulf through Ft. Devens, Massachusetts, were recruited to participate in a survey to assess psychological health and combat exposure. Follow-up questionnaires were designed to assess long-term health, and psychological and functional well-being, as well as Gulf-specific environmental and combat exposures. Follow-up questionnaires were taken from the baseline survey (1991). All demographic variables were taken from the baseline survey (1991). All participants gave their informed consent for inclusion at each timepoint, before they participated in the surveys. Institutional review board approvals were obtained from VA Boston Healthcare System and Boston University before initiating the surveys.

The Health Symptom Checklist

The Health Symptom Checklist is a pre-set list of health symptoms originally adapted from Bartone et al. This measure asks participants to indicate whether they had experienced any of the specific health symptoms. At Follow-up 1, a 20-item Health Symptom Checklist asked veterans to indicate the frequency of 20 symptoms over the past 4 weeks using a Likert-scale rating (none, a little, often, very often, yes to no). Follow-up 2 included a 52-item Expanded Health Checklist, which asked veterans to indicate whether they experienced each health symptom over the past 4 weeks (yes or no). If the veteran indicated yes, they were asked to indicate the frequency of the symptom as either sometimes, or a lot. Follow-up 3 included a 34-item Health Symptom Checklist, which asked veterans to endorse whether they experienced a symptom (yes or no) over the past 30 days. If veterans indicated yes, they were asked to indicate the frequency of the symptom (rarely or once or twice in all, some (about once per week), often (several times per week), very often (almost every day)). All responses were dichotomized as present or absent. If a veteran initially indicated not experiencing a symptom, but then endorsed a frequency, the response was recoded as being present.

On the basis of the epidemiological literature examining GW exposures and reported health symptoms, symptoms were chosen that could be characterized as belonging to one of the following categories: mood-cognition, fatigue, and neurological or physical symptoms. A total of 12 symptoms fit these categories and were included in all follow-up surveys: difficulty concentrating, feeling depressed, crying easily, feeling anxious, trouble sleeping, lack of energy, dizziness, headache, muscle twitching or trembling, rapid heart rate, skin rash, and shortness of breath.

Gulf War Exposure Characterization

At Follow-up 2 and Follow-up 3, FDC veterans were asked about environmental and combat exposures specific to GW deployment. Exposure to PB pills and chemical weapons were assessed at follow-up 2 and Follow-up 3. There was substantial agreement for PB pill exposure as assessed by the Kappa statistic (0.628 95% confidence interval, 0.516 to 0.740). However, agreement was low for chemical weapon exposure (0.098 95% CI = 0.005 to 0.201). This may be due to the differential wording for the assessment of chemical weapons at Follow-up 2 and Follow-up 3. Follow-up 2 assessed number of times a participant was on formal alert for a chemical attack, and Follow-up 3 assessed whether or not the participant was ever exposed to chemical/biological warfare. To minimize the length of time between deployment and recall, exposure data were taken from the Follow-up 2 survey, 6 years after deployment.

Veterans were asked to recall whether or not they had consumed PB pills and heard formal chemical weapons alerts in a dichotomous, yes or no response. They were also asked whether they had experienced exposure to smoke from tent heaters according to these categories: “not exposed,” “exposed but did not feel ill,” “exposed and felt ill.” Answers for these three GW-specific exposures were dichotomized into exposed and not exposed.

In addition, veterans were measured in regard to exposure to the Khamisiyah weapons demolition. In March of 1991, the Army Corp of engineers detonated underground munitions bunkers with thousands of Iraqi rockets, which they were unaware at the time had chemical weapons, sarin/cyclosarin, in the tips of the rockets. When the explosions occurred, it was estimated that over 100,000 GW veterans were exposed to low-level sarin/cyclosarin due to their proximity to the air plumes from the explosions. The Department of Defense (DoD) eventually notified exposed veterans by letter and a registry was established. A list of FDC veterans who received notification letters of potential exposure to sarin based on their proximity to the Khamisiyah weapons depot detonations and presumed to be exposed to low levels of sarin/cyclosarin based on wind plume modeling in 2000 was obtained by the DoD. These exposures were categorized as a dichotomous variable, yes (received a notification letter from the 2000 DOD exposure model) or no (never received notification letter from the 2000 DOD exposure model).

FIGURE 1. Overall timeline of the Ft. Devens Cohort (FDC) study.
a notification letter) for Khamisiyah exposure status as a proxy for sarin chemical weapons exposure during their deployment.

**Data Analysis**

Descriptive statistics were used to describe the demographics and characteristics of the full FDC and the current study sample. The frequencies and percentages of health symptoms at each follow-up time point are reported. A generalized estimating equation using a logit link and an unstructured correlation was built for each health symptom, which included the GW-specific exposures to tent heaters, PB pills, self-reported exposure to chemical warfare agents and notification of proximity to sarin/cyclosarin air plumes, as well as the additional covariates of baseline age, active duty status, post-traumatic stress disorder (PTSD) status, and sex.

In addition, dates of survey completion were used to determine the amount of time in years since baseline survey completion to evaluate the effect of time on health symptom reporting. Interaction terms for years since deployment and each exposure were initially included in each model. Interaction terms that were not significant were removed from final models. Odds ratios (ORs) and 95% CIs were calculated from the logistic regression models. In interactions models, years since deployment were centered at 1.24 years, the mean time since deployment at Follow-up 1, so that ORs for exposure involved in interactions are describing associations at Follow-up 1. P values less than 0.05 were considered significant. Adjustments were not made for multiplicity. All analyses were performed using SAS 9.4 (SAS, Cary, North Carolina).

**RESULTS**

**Demographics and Baseline Characteristics**

A total of 2949 FDC veterans completed the initial survey upon return from the GW in 1991. A total of 295 FDC veterans responded at all three time points, of whom 292 veterans had health symptom data at all three time points. In this subsample, the average age immediately after deployment was approximately 32.3 years, the mean time since deployment at Follow-up 1, so that ORs for exposure involved in interactions are describing associations at Follow-up 1. Of the 2949 veterans, 91.6% reported exposure to tent heaters. Approximately, 41.4% of veterans received notification of exposure to sarin gas from the Khamisiyah munitions demolition. On average, the number of years since deployment was 1.24 ± 0.28 years at Follow-up 1, 5.95 ± 0.18 years at Follow-up 2, and 23.91 ± 0.92 years at Follow-up 3. Frequencies and percentages of reported health symptoms at each Follow-up period are displayed in Table 2. Frequency of symptoms ranged from as low as 13.0% (rapid heart rate) to as high as 66.4% (trouble sleeping).

**Longitudinal Associations Between Gulf War Exposure and Health**

The relation between GW-specific exposures and health symptoms are displayed in Table 3. Overall, years since deployment was significantly associated with an increase in reporting of 9 of the 12 symptoms and a decrease in one symptom (headache) over time (Table 3). The increase odds of reporting these symptoms ranged from 2% to 6% per year since deployment. Self-reported exposure to chemical weapons was not significantly associated with health symptoms. However, self-reported exposure to tent heaters was associated with increased odds of crying easily (OR = 1.77, 95% CI = 1.03 to 3.04, P = 0.038) and muscle twitch or trembling (OR = 1.50, 95% CI = 1.00 to 2.23, P = 0.048) compared with veterans not reporting exposure to tent heaters. Significant interactions between tent heaters and years since deployment were noted for lack of energy (P = 0.036) and shortness of breath (P = 0.009), with differing patterns. Those reporting exposure to tent heaters initially reported lower frequency of shortness of breath (at Follow-up 1 OR = 0.59, 95% CI = 0.33 to 1.04, P = 0.068), but eventually reported a higher frequency (at Follow-up 3 OR = 1.33, 95% CI = 0.75 to 2.37, P = 0.331), than those unexposed. Those reporting exposure to tent heaters initially reported higher frequency of trouble sleeping (at Follow-up 1 OR = 1.60, 85% CI = 0.92 to 2.76, P = 0.09), but eventually reported lower frequencies (at Follow-up 3 OR = 0.84, 95% CI = 0.50 to 1.43, P = 0.52), than those unexposed. Self-reported exposure to PB pills was associated with increased odds of feeling depressed (OR = 1.65, 95% CI = 1.06 to 2.55, P = 0.026), crying easily (OR = 1.86, 95% CI = 1.01 to 3.43, P = 0.046), and lack of energy (OR = 1.74, 95% CI = 1.14 to 2.64, P = 0.010) compared with veterans not reporting PB pill usage. In addition, a significant interaction was noted between PB pill exposure and years since deployment for trouble sleeping (P = 0.008), where those reporting exposure initially reported higher frequency of trouble sleeping (at Follow-up 1 OR = 2.09,

**TABLE 1. Demographics and Baseline Characteristics of Full Ft. Devens Cohort and Current Ft. Devens Cohort Sample**

| Demographic/Characteristic | Full Devens Cohort (N = 2,949) | Study Sample (N = 292) |
|-----------------------------|---------------------------------|------------------------|
| Age, years’                 | 30.2 ± 8.4                      | 32.3 ± 8.5             |
| Male, n (%)                 | 2,702 (91.6%)                   | 260 (89.0%)            |
| Caucasian, n (%)            | 2,443 (82.8%)                   | 272 (92.3%)            |
| Active Duty, n (%) (vs Reserve, National Guard) | 823 (27.9%) | 46 (16.4%) |
| Mississippi PTSD scale-score | 61.9 ± 13.4                     | 61.9 ± 14.2            |
| Clinical cutoff on Mississippi scale-score, n (%) | 116 (3.9%) | 16 (5.5%) |

*P < 0.05

**TABLE 2. Frequencies and Percentages of Reported Health Symptoms at Each Follow-Up**

| Symptom                          | Follow-up 1 (n = 2,894) | Follow-up 2 (n = 2,702) | Follow-up 3 (n = 2,649) |
|----------------------------------|-------------------------|-------------------------|-------------------------|
| Age, years’                      | 30.2 ± 8.4              | 32.3 ± 8.5              | 34.4 ± 9.2              |
| Male, n (%)                      | 2,702 (91.6%)           | 260 (89.0%)             | 246 (92.8%)             |
| Caucasian, n (%)                 | 2,443 (82.8%)           | 272 (92.3%)             | 281 (93.3%)             |
| Active Duty, n (%) (vs Reserve, National Guard) | 823 (27.9%) | 46 (16.4%) |
| Mississippi PTSD scale-score     | 61.9 ± 13.4             | 61.9 ± 14.2             | 62.0 ± 14.3             |
| Clinical cutoff on Mississippi scale-score, n (%) | 116 (3.9%) | 16 (5.5%) |

*P < 0.05

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Fatigue

Trouble Sleeping 0.91 (0.62–1.35) 0.65 2.09 (1.25–3.48) 0.005 0.92 (0.53–1.59) 0.76 0.96 (0.66–1.39) 0.83 1.06 (1.04–1.09) <0.001

Lack of Energy 1.30 (0.84–2.01) 0.24 1.74 (1.14–2.64) 0.010 0.80 (0.45–1.40) 0.43 1.03 (0.72–1.48) 0.88 1.03 (1.01–1.05) 0.005

Neurological or Physical

Dizziness 0.88 (0.58–1.33) 0.54 1.42 (0.84–2.38) 0.19 0.70 (0.36–1.38) 0.31 1.35 (0.83–2.21) 0.23 1.02 (1.00–1.04) 0.08

Headache 0.67 (0.45–1.00) 0.05 1.03 (0.64–1.62) 0.23 0.48 (0.16–1.52) 0.19 0.63 (0.42–0.93) 0.02 0.90 (0.62–1.31) 0.66 0.91 (0.63–1.31) 0.66

Muscle Twitch or Trembling 1.50 (1.00–2.23) 0.04 1.18 (0.75–1.86) 0.53 0.63 (0.34–1.15) 0.13 1.09 (0.72–1.66) 0.69 1.03 (1.01–1.04) <0.001

Skin Rash 0.90 (0.56–1.43) 0.65 1.49 (0.85–2.60) 0.17 0.61 (0.29–1.27) 0.19 1.21 (0.77–1.89) 0.40 1.03 (1.01–1.04) <0.001

Shortness of Breath 0.59 (0.33–1.04) 0.07 1.51 (0.89–2.57) 0.13 0.84 (0.42–1.67) 0.61 1.60 (0.92–2.76) 0.09 1.02 (1.00–1.05) 0.09

Models included baseline age, sex, active military status, and PTSD status as covariates.

Interactions between years since deployment and exposures were assessed in each model. Significant interactions (denoted with *) were found between years since deployment and tent heaters for lack of energy (P = 0.036) and shortness of breath (P = 0.009), between years since deployment and PB pills for trouble sleeping (P = 0.008), and between years since deployment and Khamisiyah Notification for crying easily (P = 0.021), dizziness (P = 0.028), skin rash (P = 0.003), and shortness of breath (P = 0.037). Odds ratios for exposure with interactions correspond to the effect at Follow-up 1, at 1.24 years post deployment.

CI, confidence interval; OR, odds ratio.

95% CI = 1.25 to 3.48, P = 0.005), but eventually endorsed trouble sleeping at the same frequency as those unexposed (at Follow-up 3 OR = 0.83, 95% CI = 0.47 to 1.46, P = 0.053).

Exposure to the sarin plume from the Khamisiyah demolition operation was associated with increased odds of trouble concentrating (OR = 1.56, 95% CI = 1.06 to 2.28, P = 0.023) and decreased odds of headaches (OR = 0.63, 95% CI = 0.42 to 0.93, P = 0.021) compared with FDC veterans who did not receive Khamisiyah notification letters. A significant interaction was noted between exposure to sarin and years since deployment for crying easily (P = 0.021), where those exposed initially reported higher frequency of crying easily (at Follow-up 1 OR = 1.94, 95% CI = 1.03 to 3.65, P = 0.040). Additional significant interactions between notification of exposure to the sarin plume from Khamisiyah and years since deployment were seen for dizziness (P = 0.028), skin rash (P = 0.003), and shortness of breath (P = 0.037). For each of these four symptoms, the exposed group reported initially reported higher frequency of symptoms than the unexposed group, but eventually reported either similar or lower rates than the unexposed group.

**DISCUSSION**

In a longitudinal analysis controlling for baseline age, active duty status, PTSD status, and sex, exposure to tent heaters was associated with higher endorsement of crying easily and muscle twitching/trembling. Our findings are similar to other studies that have shown tent heater exposure to be related to increased reporting of cardiac, neurological, and pulmonary symptoms, as well as chronic multisymptom illness (CMI), involving fatigue, cognitive mood, and musculoskeletal symptoms.

Self-reported exposure to PB pills was significantly associated with higher endorsement of feeling depressed, crying easily, trouble sleeping, and lack of energy. These results are consistent with previous reports of nearly 250,000 GW veterans, which suggested consumption of PB pills is a causal factor of GWI (GWI), a disorder that affects approximately 30% of GW veterans and is characterized by a combination of symptoms that include fatigue, cognitive and mood dysfunction, musculoskeletal pain, gastrointestinal complaints, respiratory symptoms, and skin rashes. Similarly, other studies demonstrated that veterans who consumed PB pills had significantly higher odds of CMI, which is a commonly used definition for GWI. In addition, self-reported PB exposure has recently been associated with medical conditions, including myocardial infarction, diabetes, and chronic fatigue syndrome.

PB reversibly binds to the same cholinergic receptors as nerve agents such as sarin and soman causing the receptors to be unavailable for these nerve agents. Thus, a Food and Drug Administration waiver was obtained by the DoD for PB pills to be administered as a prophylactic medication during the GW. Safety was assumed, as PB pills were commonly used to treat a chronic autoimmune neurumocutaneous disorder, myasthenia gravis. However, PB is an acetylcholinesterase inhibitor (AChEi), which reversibly inactivates the AChE enzyme preventing the breakdown of acetylcholine (ACh). Subsequent accumulation of ACh in the synapse stimulates muscarinic and nicotinic cholinergic receptors and has been shown to be associated with later cognitive, muscle, and sleep dysfunction. Further, oligodendrocytes have now been shown to have functional cholinergic receptors that are affected by GW-relevant AChE inhibitors, suggesting brain myelination may also be affected.

Exposure to AChEis, such as PB pills and sarin nerve gas, may also trigger brain-immune responses due to activation of microglia and astrocytes, the brain’s immune sentences. These sentinel enhance cytokine and chemokine signaling leading to chronic neuroinflammation. In addition, neuroendocrine and autonomic nervous system disruptions can occur, as well as alteration of innate immune function as cholinergic receptors also exist outside the central nervous system (CNS). Exposure to AChEis, such as PB pills and sarin nerve gas, may also trigger brain-immune responses due to activation of microglia and astrocytes, the brain’s immune sentences. These sentinel enhance cytokine and chemokine signaling leading to chronic neuroinflammation. In addition, neuroendocrine and autonomic nervous system disruptions can occur, as well as alteration of innate immune function as cholinergic receptors also exist outside the central nervous system (CNS). Exposure to AChEis, such as PB pills and sarin nerve gas, may also trigger brain-immune responses due to activation of microglia and astrocytes, the brain’s immune sentences. These sentinel enhance cytokine and chemokine signaling leading to chronic neuroinflammation. In addition, neuroendocrine and autonomic nervous system disruptions can occur, as well as alteration of innate immune function as cholinergic receptors also exist outside the central nervous system (CNS). Exposure to AChEis, such as PB pills and sarin nerve gas, may also trigger brain-immune responses due to activation of microglia and astrocytes, the brain’s immune sentences. These sentinel enhance cytokine and chemokine signaling leading to chronic neuroinflammation. In addition, neuroendocrine and autonomic nervous system disruptions can occur, as well as alteration of innate immune function as cholinergic receptors also exist outside the central nervous system (CNS). Exposure to AChEis, such as PB pills and sarin nerve gas, may also trigger brain-immune responses due to activation of microglia and astrocytes, the brain’s immune sentences. These sentinel enhance cytokine and chemokine signaling leading to chronic neuroinflammation. In addition, neuroendocrine and autonomic nervous system disruptions can occur, as well as alteration of innate immune function as cholinergic receptors also exist outside the central nervous system (CNS).
exposure was associated with increased odds of trouble concentrating and crying easily and decreased odds of headaches. These symptoms of altered mood and cognitive functioning after nerve agent exposure are consistent with those reported in the literature for other known exposed groups, including those poisoned from the Tokyo subway terrorist attacks in 1995. The DoD notified approximately 100,000 veterans who may have been exposed to sarin following the demoli-

tion plume, based on where their unit was located. Although we recognize that the Khamisiyah notified groups may not be the only GW veterans exposed to chemical weapons during the war, the Khamisiyah incident remains at least one verified and sensitive measure of sarin exposure, particularly for those who were at close proximity to the demolition plume. Our group was the first to report a dose–response association of brain imaging outcomes with the Khamisiyah plume estimated exposure levels obtained by DOD and this finding has since been partially replicated. Therefore, these results suggest that self-reported exposure to chemical weapons may not be as sensitive as the Khamisiyah notification letter for longitudinal analyses.

Previous research indicates that symptom reporting remains relatively stable or has increased slightly in GW veterans over time. In this sample, significant associations between time since deployment and symptom reporting were observed in 10 of 12 symptoms. Generally, these symptoms significantly increased over time, with the exception of headaches, which decreased. Although speculative, this may be due to veterans seeking effective treatments early after return from deployment for this symptom. The changes in symptom trajectories could indicate emerging or worsening symptoms with age or could reflect normative aging. Porter et al found that rates of CMI significantly increased over time in all groups assessed, with GW veterans having the highest prevalence rate compared with era and non-era personnel at each time point. It is particularly striking that the symptoms involved in the criteria for CMI are increasing over time in veteran comparison groups (era and non-era personnel) and thus may be related to normal aging symptomatology. However, it was recently demonstrated that GW deployment is associated with accelerated aging such that the average 50-year-old GW deployed veteran exhibits rates of medical conditions that are similar to those seen in 70-year-old non-veterans. This suggests that symptom trajectories may be altered in this aging population and GW veterans should be carefully monitored for age-related chronic illnesses and treated to prevent further health complications.

**Limitations and Strengths**

The current analysis was limited to subjects that had data at all three time points, which may limit the generalizability of our results to the greater Ft. Devens cohort and the population of GW veterans as a whole. Indeed, compared with the greater Ft. Devens cohort, the current sample appeared to be more likely to meet the clinical cut off for PTSD and more likely to be Caucasian. Selection bias is also a concern for a variety of reasons. First, selection bias is inherent in surveys, as it is possible that veterans who respond may be different than veterans that do not respond. Concern for selection bias is higher in longitudinal studies, as individuals with more health problems may be more likely to remain in the study than healthy individuals. Alternatively, longitudinal studies also exclude the sickest veterans, who are either too ill to participate, or may have died during the course of the study. However, it is important to note that the FDC is unlike other veteran samples, as they were not recruited from patients at the Department of Veteran Affairs hospitals but rather were recruited and interviewed as a cohort study days after their return from the war.

Another important challenge to studying the effects of GW exposures on veteran health is a lack of records or measurements quantifying chemical exposures during deployment. It may also be the case that other exposures that were not assessed in this cohort may be contributing to chronic health symptoms in this cohort including depleted uranium or other heavy metals that may be better suited to biomarker studies for assessment. However, although three of the four exposures analyzed were self-reported, recall bias was minimized by using exposure data from Follow-up 2, which was less than 5 years after return from the GW and before many exposure-outcome reports had been made public. It is also important to note that the first round of DOD Khamisiyah notification letters were sent out in 1997. The first two surveys included in this study were conducted before the Khamisiyah letters were sent. Therefore, the subjective reporting of symptoms may be less biased, as veterans may not have known that they were exposed to sarin and thus would not attribute their symptoms as such.

The current study was limited to using 12 health symptom variables that were reported at all three time points. Some of the health symptoms assessed differed slightly in wording across the three surveys, and thus may have contributed to the decrease in symptoms observed at follow-up 2. It would be of interest to explore other symptoms commonly reported among GW veterans, including gastrointestinal symptoms such as nausea and diarrhea, respiratory symptoms such as coughing and wheezing, and additional cognitive symptoms including difficulty remembering and trouble finding words. In addition, the analyses presented in this paper reflect group data. Therefore, individual variation of symptoms over time may have been masked. Future studies should investigate individual trajectories of symptoms and their associations with exposure. Similarly, results are from a limited sample of GW veterans, and may lack generalizability to a larger population. Finally, analyses were not adjusted for multiple comparisons which increases the chance of type 1 error.

**CONCLUSION**

The current longitudinal analysis demonstrated higher odds of symptom reporting associated with exposure to tent heaters, consumption of PB pills, and exposure to sarin/cyclosarin from the Khamisiyah demolitions plume, among a group of GW veterans while controlling for baseline age, active duty status, PTSD status, and sex. In addition, overall, time since deployment was significantly related to symptom reporting indicating that symptoms are increasing over time in this cohort of GW veterans. This increase in symptoms may be the result of normative aging or may be specific to neurotoxicant-induced accelerated aging patterns in certain instances that should be further studied. These results raise concerns about GW veterans’ health more than 25 years after the war. GW veterans should continue to be followed over time by their care providers to properly manage long-term health issues and treat new health issues that may arise.

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