

# **HHS Public Access**

J Am Geriatr Soc. Author manuscript; available in PMC 2015 December 01.

#### Published in final edited form as:

Author manuscript

J Am Geriatr Soc. 2014 December ; 62(12): 2350–2356. doi:10.1111/jgs.13135.

# Higher Perceived Stress Scale scores are associated with higher pain intensity and pain interference levels in Older Adults

# Robert S. White, M.D.,

Saul B. Korey Department of Neurology, Department of Epidemiology and Population Health and the Einstein Aging Study, from the Albert Einstein College of Medicine, Bronx, NY.; Department of Anesthesiology, Weill Cornell Medical Center, New York, NY

# Julie Jiang, B.S.,

Saul B. Korey Department of Neurology, Department of Epidemiology and Population Health and the Einstein Aging Study, from the Albert Einstein College of Medicine, Bronx, NY

# Charles B. Hall, Ph.D.,

Department of Epidemiology and Population Health, Saul B. Korey Department of Neurology, and the Einstein Aging Study, from the Albert Einstein College of Medicine, Bronx, NY

# Mindy J. Katz, M.P.H.,

Saul B. Korey Department of Neurology and the Einstein Aging Study, from the Albert Einstein College of Medicine, Bronx, NY

# Molly E. Zimmerman, Ph.D.,

#### Author Contributions:

Robert S. White: preparation of manuscript, study concept or design, analysis and interpretation of data, statistical analysis, acquisition of data, study supervision, obtaining funding.

Corresponding author: Richard B. Lipton, M.D., Albert Einstein College of Medicine, Louis and Dora Rousso Building, 1165 Morris Park Avenue, Room 332, Bronx, NY 10461, richard.lipton@einstein.yu.edu, Phone number: (718) 430-3886, Fax: (718) 430-3857. Alternative Corresponding author: Robert S. White, M.D., robert.white@med.einstein.yu.edu.

**Conflict of Interest:** Dr. White is supported by grants UL1TR000086, TL1RR000087, and KL2TR000088; reports no conflict of interest. Ms Jiang is supported by grants UL1TR000086, TL1RR000087, KL2TR000088, and T32-GM007288; reports no conflict of interest. Dr. Hall is supported by CDC grants 1U01-OH10412-01 (Project Primary Investigator), 1U01OH010411-01, 1U01OH010513-01, NIH grants P01 AG03949, R01 AG034119, R01 AG022092, UL1 RR025750, and P30 CA013330, along with CDC contracts 200-2011-39378 and 200-2011-39489; reports no conflict of interest. Ms. Katz is supported by grants NIA 2P01 AG03949, NIA R03 AG045474, and NIA R01 AG039409; has an outstanding contract with Bristol Myers Squibb, Inc. Dr. Zimmerman is a co-investigator on NIH AG003949; reports no conflict of interest. Dr. Sliwinski is supported by NIA grant number R01 AG39409; reports no conflict of interest. Dr. Sliwinski is supported by NIA grant number R01 AG39409; reports no conflict of interest. Dr. Sliwinski is supported by NIA grant number R01AG027734 (Project Leader), R01AG025119 (Investigator), R01AG02374-06A2 (Investigator), R01AG034119 (Investigator), R01AG03857 (Mentor), K23NS05140901A1 (Mentor), and K23NS47256 (Mentor), the National Headache Foundation, and the Migraine Research Fund; serves on the editorial boards of Neurology and Cephalalgia and as senior advisor to Headache, has reviewed for the NIA and NINDS, holds stock options in eNeura Therapeutics (a company without commercial products); serves as consultant, advisory board member, or has received honoraria from: Allergan, American Headache Society, Autonomic Technologies, Boston Scientific, Bristol Myers Squibb, Cognimed, Colucid, Eli Lilly, eNeura Therapeutics, GlaxoSmithKline, MAP, Merck, Nautilus Neuroscience, Novartis, NuPathe, Vedanta, Zogenix.

Julie Jiang: preparation of the manuscript, study concept or design, analysis and interpretation of data, statistical analysis. Charles B. Hall: preparation of the manuscript, study concept or design, analysis and interpretation of data, statistical analysis. Mindy J. Katz: preparation of the manuscript, study concept or design, analysis and interpretation of data, statistical analysis, acquisition of data, study supervision.

Molly E. Zimmerman: preparation of the manuscript, study concept or design, analysis and interpretation of data, statistical analysis. Martin Sliwinski: preparation of the manuscript, study concept or design, analysis and interpretation of data, statistical analysis. Richard B. Lipton: preparation of manuscript, study concept or design, analysis and interpretation of data, statistical analysis.

Saul B. Korey Department of Neurology and the Einstein Aging Study, from the Albert Einstein College of Medicine, Bronx, NY

#### Martin Sliwinski, Ph.D., and

Department of Human Development and Family Studies and the Center for Healthy Aging, from Pennsylvania State University, University Park, PA

#### Richard B. Lipton, M.D.

Saul B. Korey Department of Neurology, Department of Epidemiology and Population Health and the Einstein Aging Study, from the Albert Einstein College of Medicine and the Montefiore Medical Center, Bronx, NY

# Abstract

**Objectives**—To determine the prevalence of bodily pain measures (pain intensity and pain interference) in elderly people and their relationship with perceived stress scale (PSS) scores.

Design—Cross-sectional.

Setting—Community.

Participants—A representative community sample of 578 subjects aged 70 and older.

**Measurements**—The prevalence of pain intensity and pain interference and their relationship with perceived stress scale scores, demographic factors, past medical history, and neuropsychological testing scores were examined. Pain intensity and pain interference were measured by the SF-36 bodily pain questions.

**Results**—The study sample of 578 participants has a mean age of 78.8 years and is 63% female. Bivariate analysis for pain measures showed that higher scores on the perceived stress scale, lower neuropsychological test scores, and medical histories were associated with both pain intensity and interference. Logistic regression showed that higher scores on the perceived stress scale were significantly associated with increased odds of having moderate/severe pain intensity and moderate/severe pain interference (with and without the inclusion of for pain intensity in the models).

**Conclusion**—Higher PSS scores are associated with higher levels of pain intensity and pain interference. In this cross-sectional analysis, directionality cannot be determined. As both perceived stress and pain are potentially modifiable risk factors for cognitive decline and other poor health outcomes, future research should address temporality and the benefits of treatment.

#### **Keywords**

chronic pain; pain intensity; pain interference; elderly

# INTRODUCTION

The Institute of Medicine classifies chronic pain as a public health problem that affects 100 million adults with an annual economic cost ranging between \$560–\$635 billion<sup>1</sup>. Where applicable, research should identify potentially modifiable risk factors and stakeholders should enact approaches aimed at pain prevention, treatment, and care<sup>1</sup>.

Cognitive, psychological, and behavioral factors influence the pain experience<sup>2–4</sup>. Chronic pain is associated with psychopathology, including psychiatric and psychological disorders<sup>2</sup>. Psychological factors are important in the coping, quality of life, and disability experienced in chronic pain sufferers<sup>4</sup>. Research has identified an association between perceived stress and various pain syndromes; including recurrent, orofacial, and arthritis related pain<sup>5–7</sup>. The perceived stress scale (PSS) is a psychosocial measurement of an individual's appraisal of life events as stressful<sup>8–10</sup>. It focuses on the subjective experience and how life is unpredictable, uncontrollable, and overloading. The PSS is correlated with life event scores, but is considered a more accurate reflection of stress experienced. Objective stress scales count number of stressful events, ignoring personal and contextual factors. In comparison, the PSS measures a cognitively mediated emotional response to an objective event incorporating an individual's social support system, robustness, and locus of control<sup>8–10</sup>.

The relationship between perceived stress and bodily pain has not been studied, particularly in an elderly community based population. There is a paucity of epidemiologic research focused on temporally concurrent perceived stress and its association with pain measures. Concurrent analysis is important, because perceived stress is variable over time and changes in response to an individual's current daily hassles, major life events, and present coping ability<sup>9</sup>.

Herein, we examined the association of perceived stress with bodily pain measures in the elderly over a 4-week period using cohort data from the Einstein Aging Study (EAS). Our pain outcomes included pain intensity (severity) and pain interference (pain related disability). We hypothesized that higher levels of perceived stress would be associated with increased levels of both measures.

# METHODS

# Study Population

Participants in this study were sampled from the EAS, a methodically-recruited, populationbased longitudinal study of adults age 70 and older who reside in Bronx County, New York. Study design, enrollment procedures, and methods have been previously described<sup>11</sup>. Katz et al. has demonstrated that the EAS is representative of the Bronx English speaking elderly community by age, gender, and education<sup>11</sup>. Participants were recruited using voter registration lists and Medicare eligibility information. Exclusion criteria included non-English speaking, severe audiovisual disturbances, prevalent dementia, institutionalization, or any condition that would interfere with participation (active psychiatric symptoms). Written informed consent was obtained during clinic visits in accordance with study protocols approved by the Institutional Review Board of the Albert Einstein College of Medicine.

#### **Clinical Evaluation**

Participants were evaluated by demographic surveys, structured medical history form, and queries concerning personal events. The entirety of our study sample was dementia free as

ascertained at case conferences with neuropsychology and neurology input using standardized clinical criteria from the DSM-IV<sup>12</sup>.

We focused on medical conditions with the highest prevalence in our study population; including diabetes, congestive heart failure, hypertension, angina, myocardial infarction, stroke, chronic obstructive lung disease, and osteoarthritis. Lifestyle variables, alcohol consumption and smoking history were included because of their impact on pain. Alcohol consumption over the past year (in grams) was categorized into tertiles; the lowest tertile serving as reference. Smoking history was categorized as never, former, and current smoker.

### Neuropsychological Evaluation

Prior literature has shown an inverse relationship between neuropsychological function and pain measures, necessitating inclusion to control for possible confounding. The Reading Subtest grade equivalency score from the Wide Range Achievement Test (WRAT3)<sup>13</sup> has a continuous range from 1–13. The Wechsler Adult Intelligence Scale - Third Edition<sup>14</sup> Verbal IQ (VIQ) score reflects performance on language-based tests of comprehension and problem solving. Free Recall (0–48) is a measure of episodic memory from the Free and Cued Selective Reminding Test<sup>15</sup>. Phonemic fluency<sup>16</sup> ("FAS"; number of correctly named words) and a test of set shifting and concept formation (Trail Making Test Part B<sup>17</sup>; seconds to test completion) measured executive functioning. Depressive symptoms were evaluated using the 15- item Geriatric Depression Scale (GDS); a score of >5 was defined as positive<sup>18</sup>.

#### Stress Evaluation

The PSS is a global assessment of an individual's perception of psychological stress during the past month<sup>9</sup>; the original scale consisted of 14-items. PSS was included in the EAS testing battery in 2006. Prior factor analysis performed in this cohort identified a two factor solution<sup>19</sup>, consistent with published literature; item 12 did not load strongly on either factor, and was not included in the analysis. Our modified measure PSS-13 had reliable factor structure and predictive validity in our population sample<sup>19</sup>. Our scale contained six negatively worded items (questions 1, 2, 3, 8, 11, and 14), and seven positively worded items (questions 4, 5, 6, 7, 9, 10, and 12). Each item was rated on a five point scale. Scores were calculated after reverse keying positive items and summation of scores. Possible total scores range from 0–52 (higher score indicates greater stress; however, this effect is non-linear). The PSS is not a diagnostic instrument and no predetermined cut-points qualify different levels of perceived stress. For our analysis, PSS was modeled two ways: 1) divided into equally weighted quartiles with the first quartile (lowest levels of stress) serving as the reference group and 2) as a continuous variable.

#### **Pain Evaluation**

Bodily pain measures were drawn from the Short Form  $36^{20}$ . Pain intensity was questioned as, "How much bodily pain have you had during the past 4 weeks?" Responses range from 1 to 6 with 1 = "None" and 6 = "Very Severe." Participants were dichotomized into two groups: no/mild pain intensity vs. moderate/severe pain intensity. Pain interference was questioned as, "During the past 4 weeks, how much did pain interfere with your normal

work (including both work outside the home and housework)?" Responses range from 1 to 5 with 1 = "Not at all" and 5 = "extremely." Participants were dichotomized into two groups: no/mild pain interference vs. moderate/severe pain interference. These methods of surveying and categorizing bodily pain outcomes have previously been used in population-based studies<sup>21, 22</sup>.

# **Statistical Analysis**

Analyses were performed using STATA software, version 12.1 (College Station, Texas). Characteristics were compared separately for the dichotomous outcome variables pain intensity and pain interference and the independent variables describing the participant's demographic and medical history. Continuous variables were compared using two-sample ttests, or, when the variables had a distribution far from normal, by Mann-Whitney Wilcoxon tests. Categorical variables were compared using the Pearson chi-square test or Fisher's exact test.

To examine the effect of PSS on pain status we fit logistic regression models to our data. We developed separate models for pain intensity (no/mild pain intensity vs. moderate/severe pain intensity) and for pain interference (no/mild pain interference vs. moderate/severe pain interference) with and without inclusion of pain intensity in the models. In an effort to take into account potential confounders we included in our models variables with bivariate baseline testing results of p < 0.25; or variables, such as age, race, and gender, that were selected a priori. This was repeated with PSS as a categorized and continuous variable. We assigned statistical significance at an alpha level of 0.05. Elevated odds ratios indicate the increased likelihood of pain intensity or pain interference. The odds ratio for continuous variables such as age represents the change in odds for each additional unit change. Model assumptions of normality and linearity were assessed both graphically and statistically; goodness of fit testing was performed.

# RESULTS

Between February 2006 and February 2012 a total of 806 individuals were evaluated for inclusion in the EAS with 19 individuals excluded for prevalent dementia at baseline. 787 dementia free individuals were considered for the current analysis. 578 individuals had available pain and stress data from the same 4 week period and were included in the present analysis; 403 individuals had complete covariate data, allowing for inclusion in logistic regression models.

Results of bivariate analysis show that both moderate/severe pain intensity (Table 1) and moderate/severe pain interference (Table 2) were associated with higher levels of perceived stress. Logistic regression showed that higher levels of perceived stress was associated with increased odds of having both bodily pain measures, moderate/severe pain intensity (OR=1.05 per unit increase in PSS score, Table 3) and moderate/severe pain interference (OR=1.07 per unit increase in PSS score, Web-Table 2). As compared to the 1<sup>st</sup> quartile (lowest levels of stress), the 2<sup>nd</sup> (OR=2.12), 3<sup>rd</sup> (OR=2.47), and 4<sup>th</sup> (OR=3.47) quartiles of PSS scores had increased odds of moderate/severe pain intensity, respectively (Table 3). As compared to the 1<sup>st</sup> quartile, the 2<sup>nd</sup> (OR=5.13), 3<sup>rd</sup> (OR=5.87), and 4<sup>th</sup> (OR=7.32) quartiles

of PSS scores had increased odds of moderate/severe pain interference, respectively (showing results for models with pain intensity included as a predictor, Web-Table 2. Similar results obtained in models without pain intensity included as a predictor, Web-Table 1).

Logistic regression analysis also revealed the following significant associations for the bodily pain measures. Increased odds of having moderate/severe pain intensity were associated with congestive heart failure and osteoarthritis; increased age was associated with decreased odds of having moderate/severe pain intensity (Table 3). Increased odds of having moderate/severe pain interference was associated with moderate/severe pain intensity, being non-Caucasian/non-African American race, having depressive symptoms, and osteoarthritis; increased age, higher free recall scores, and higher consumption of alcohol were associated with decreased odds of having moderate/severe pain interference (Web-Table 1 and Web-Table 2).

# DISCUSSION

We have shown that higher levels of perceived stress were associated with both higher levels of pain intensity and pain interference in an elderly population of dementia-free individuals. This association remained significant when age, gender, race, medical history, depressive symptoms and neuropsychological test performance were included as covariates in the models; and when modeled as continuous or categorical variables, pointing to the robustness of our findings. Our research focuses on a community based sample of older adults; prior research was focused on pediatric and middle aged populations and examined specific pain syndromes<sup>5–7</sup>. This is the first study to our knowledge to examine the association between perceived stress and bodily pain measures.

The relationship between increased psychological distress and chronic pain<sup>2–4</sup> is often referred to as the diathesis-stress model of pain<sup>23, 24</sup>. Chronic stress leads to dysregulation in the body's "supersystem" (comprised of the nervous, endocrine, and immune systems) that regulate an individual's experience of pain<sup>23</sup>. The excess psychosocial distress increases the allostatic load placed on the body resulting in chronic disorders, including pain<sup>23, 24</sup>. Increased perceived stress has been associated with a broad array of adverse health outcomes<sup>25, 26</sup>. Based on our results, the diathesis or vulnerability would be higher levels of perceived stress experienced by the individual. This present study, however, did not examine other psychological constructs that are related to pain, limiting our ability to identify mediators or confounders in the perceived stress and pain pathways. A limitation of this study is that it is cross-sectional. Therefore, temporal sequences describing pain and stress cannot be established. Future research should explore longitudinal analysis to better explore these relationships.

Our model results show that higher scores on the perceived stress scale were associated with an increase in both pain intensity and pain interference (and remained significant when pain intensity level was included as a model predictor). We suggest that perceived stress may influence pain interference directly and through a pathway mediated by pain intensity.

Most likely a bidirectional relationship exists where pain exacerbates stress and stress exacerbates pain. Methods that reduce both may improve the adverse health outcomes linked to these common symptoms in late life. Research has linked the presence of high stress and chronic pain together as a risk factor for shortened telomere length and advanced cellular aging<sup>27</sup>; which is associated with negative health outcomes including cardiovascular disease, cognitive function, and immune function, among others. Presence of both frequent musculoskeletal pain and perceived stress are risk factors for reduced work performance and ability<sup>28</sup>. Therefore, methods to reduce stress should be employed and further explored. These include meditation, physical exercises, and coping strategies<sup>29</sup>, 30.

This study has several limitations. We utilized data from a sample of elderly participants, age 70 years and above, who reside in Bronx, New York; questions about external validity exist and whether our results can be applied to other samples. Studies have shown that the prevalence of pain interference rises sharply with increasing age<sup>22</sup>; the prevalence of chronic pain increases until around age 65, followed by a plateau, and a decrease in reported pain for ages 75 and older<sup>21</sup>. Prevalence of medical histories in the EAS was previously found to be similar to rates for persons over age 65 in the US<sup>11</sup>.

The EAS does not have sufficient data on participant's anxiety, pain self-efficacy, pain coping, and pain acceptance. Future research should reexamine our findings in light of these other psychological constructs. Some pain associated disease states such as fibromyalgia were excluded from our analysis because of insufficient data. We relied on self-reported data. Recall bias is a possibility, although this would tend to be non-differential and would attenuate our results towards the null hypothesis. The bodily pain measures reflect a four week prevalence of pain and may not be representative of experienced pain for longer time periods. These pain measures were used in prior studies<sup>21, 22</sup> and we believe they are indeed representative of the pain experience. Lack thereof appropriate representation would most likely be non-differential and would serve to attenuate our findings toward the null hypothesis.

In light of the economic recession (2008–2009) and increase in perceived stress due to nationwide financial and occupational uncertainties, one potential limitation of our study was the use of PSS data ranging from 2006 to 2012. Cohen at al. examined this question and reported that the associated increase in perceived stress did not affect individuals aged 65 and older<sup>8</sup>, representative of our study population.

Our study has a number of strengths. Our sample was drawn from the EAS, a large ethnically diverse population-based study. The EAS uses well established procedures to ascertain demographic information, medical history, and neuropsychological scores. Our measures for pain<sup>21, 22</sup> and perceived stress<sup>4, 9, 10</sup> have been previously studied and shown to be psychometrically accurate and reliable. These measures were temporally balanced during the same 4-week time period allowing for concurrent associations between perceived stress and pain to be calculated. The models used in our study included all covariates with bivariate testing <0.25 and we believe that our findings for perceived stress are conservative estimates.

The present study has shown that persons with higher levels of perceived stress have increased reporting of pain intensity and pain interference. Caution must be used in interpreting our results since they show only associations and do not show causality. Future research should focus on interventional trials to determine if stress reducing techniques lead to a reduction in pain measures.

# **Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

# Acknowledgments

We thank the participants, investigators, and staff of the EAS. This work was supported by the Lenard and Sylvia Marx-Better Foundation, the Czap Foundation, and the Hollander Family Foundation.

**FUNDING:** The project described was supported by the National Center for Advancing Translational Sciences (NCATS), a component of the National Institutes of Health (NIH), through CTSA grant numbers UL1TR000086, TL1RR000087, and KL2TR000088. Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the NIH.

**Sponsor's Role:** The sponsor, the National Institutes of Health, had no role in the design, running, analysis, or interpretation of the results of this study.

### References

- 1. Relieving Pain in America: A Blueprint for Transforming Prevention, Care, Education, and Research. The National Academies Press; 2011.
- Dersh J, Polatin PB, Gatchel RJ. Chronic pain and psychopathology: Research findings and theoretical considerations. Psychosom Med. 2002; 64:773–786. [PubMed: 12271108]
- 3. Gatchel RJ. Comorbidity of chronic pain and mental health disorders: The biopsychosocial perspective. Am Psychol. 2004; 59:795–805. [PubMed: 15554853]
- Turk DC, Okifuji A. Psychological factors in chronic pain: Evolution and revolution. J Consult Clin Psychol. 2002; 70:678–690. [PubMed: 12090376]
- Alfven G, Östberg V, Hjern A. Stressor, perceived stress and recurrent pain in Swedish schoolchildren. J Psychosom Res. 2008; 65:381–387. [PubMed: 18805248]
- Harris M, Loxton D, Sibbritt D, et al. The influence of perceived stress on the onset of arthritis in women: Findings from the Australian Longitudinal Study on Women's Health. Ann Behav Med. 2013; 46:9–18. [PubMed: 23436274]
- Sanders AE, Slade GD. Gender modifies effect of perceived stress on orofacial pain symptoms: National Survey of Adult Oral Health. J Orofac Pain. 2011; 25:317–326. [PubMed: 22247927]
- Cohen S, Janicki-Deverts D. Who's Stressed? Distributions of psychological stress in the United States in probability samples from 1983, 2006, and 20091. J Appl Soc Psychol. 2012; 42:1320– 1334.
- Cohen S, Kamarck T, Mermelstein R. A global measure of perceived stress. J Health Soc Behav. 1983; 24(4):385–396. [PubMed: 6668417]
- Cohen, S.; Kessler, RC.; Underwood Gordon, L. Measuring stress: A guide for health and social scientists. New York: Oxford; 1995.
- Katz MJ, Lipton RB, Hall CB, et al. Age-specific and sex-specific prevalence and incidence of mild cognitive impairment, dementia, and Alzheimer dementia in blacks and whites: A report from the Einstein Aging Study. Alzheimer Dis Assoc Disord. 2012; 26:335–343. [PubMed: 22156756]
- American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders, DSM-IV. Washington, DC: American Psychiatric Association; 1994.
- Wilkinson, GS. Wide Range Achievement Test 3-Administration manual. Wilmington, DE: Jastak Associates, Inc; 1993.

- Grober E, Lipton RB, Hall C, et al. Memory impairment on free and cued selective reminding predicts dementia. Neurology. 2000; 54:827–832. [PubMed: 10690971]
- Spreen, OSE. A compendium of neuropsychological tests: Administration, norms and commentary.
  Oxford University Press; New York: 1998.
- Bowie CR, Harvey PD. Administration and interpretation of the Trail Making Test. Nat Protoc. 2006; 1:2277–2281. [PubMed: 17406468]
- Marc LG, Raue PJ, Bruce ML. Screening Performance of the 15-Item Geriatric Depression Scale in a Diverse Elderly Home Care Population. Am J Geriatr. 2008; 16:914–921.
- 19. Ezzati A, Jiang J, Katz MJ, et al. Validation of the Perceived Stress Scale in a community sample of older adults. Int J Geriatr Psychiatry. 2014; 29:645–652. [PubMed: 24302253]
- 20. Ware JE Jr, Sherbourne CD. The MOS 36-item short-form health survey (SF-36). I. conceptual framework and item selection. Med Care. 1992; 30:473–483. [PubMed: 1593914]
- Blyth FM, March LM, Brnabic AJM, et al. Chronic pain in Australia: A prevalence study. Pain. 2001; 89:127–134. [PubMed: 11166468]
- 22. Thomas E, Peat G, Harris L, et al. The prevalence of pain and pain interference in a general population of older adults: Cross-sectional findings from the North Staffordshire Osteoarthritis Project (NorStOP). Pain. 2004; 110:361–368. [PubMed: 15275787]
- 23. Chapman CR, Tuckett RP, Song CW, et al. Pain and stress in a systems perspective: reciprocal neural, endocrine, and immune interactions. J. 2008; 9:122–145.
- 24. Turk D. A diathesis-stress model of chronic pain and disability following traumatic injury. Pain Res Manage. 2002; 7:9–19.
- Arnold SV, Smolderen KG, Buchanan DM, et al. Perceived stress in myocardial infarction: Longterm mortality and health status outcomes. J Am Coll Cardiol. 2012; 60:1756–1763. [PubMed: 23040574]
- Cohen S, Doyle WJ, Skoner DP. Psychological stress, cytokine production, and severity of upper respiratory illness. Psychosom Med. 1999; 61:175–180. [PubMed: 10204970]
- 27. Sibille K, Langaee T, Burkley B, et al. Chronic pain, perceived stress, and cellular aging: an exploratory study. Mol Pain. 2012; 8:12. [PubMed: 22325162]
- Lindegård A, Larsman P, Hadzibajramovic E, et al. The influence of perceived stress and musculoskeletal pain on work performance and work ability in Swedish health care workers. Int Arch Occup Environ Health. 2014; 87:373–379. [PubMed: 23609321]
- Deckro GR, Ballinger KM, Hoyt M, et al. The evaluation of a mind/body intervention to reduce psychological distress and perceived stress in college students. J Am Coll Health. 2002; 50:281– 287. [PubMed: 12701653]
- Hartfiel N, Burton C, Rycroft-Malone J, et al. Yoga for reducing perceived stress and back pain at work. Occup Med (Lond). 2012; 62:606–612. [PubMed: 23012344]

#### Table 1

#### Demographic and medical characteristics of subjects by presence of pain intensity

Variables	Total (n=578)	No/mild Pain Intensity (n=384) 66.4%	Moderate/Severe Pain Intensity (n=194) 33.6%	p-value
Demographics				
Age Mean (± SD)	78.8 (5.4)	78.9 (5.6)	78.5 (5.1)	0.44
Female, N (%)	367 (63.5)	217 (56.5)	150 (77.3)	< 0.001
Race - African American, N (%)	168 (29.1)	102 (26.6)	66 (34.0)	0.11
Race - Other, N (%)	48 (8.3)	30 (7.8)	18 (9.3)	
Lifestyle, N (%)				
Non-smoker	244 (44.4)	162 (44.0)	82 (45.3)	
Former smoker	279 (50.8)	190 (51.6)	89 (49.2)	0.76
Current Smoker	26 (4.7)	16 (4.4)	10 (5.5)	
Alcohol – 1 <sup>st</sup> tertile	158 (28.7)	96 (26.0)	62 (34.3)	0.008
Alcohol – 2 <sup>nd</sup> tertile	181 (32.9)	115 (31.2)	66 (36.5)	
Alcohol – 3 <sup>rd</sup> tertile	211 (38.4)	158 (42.8)	53 (29.3)	
Perceived Stress				
PSS Mean (S.D.)	16.4 (7.7)	15.6 (7.8)	17.9 (7.2)	< 0.001
1 <sup>st</sup> Quartile (PSS scores 0–11), N (%)	163 (28.2)	124 (32.3)	39 (20.1)	
2nd Quartile (PSS scores 12-16), N (%)	145 (25.1)	97 (25.3)	48 (24.7)	0.003
3 <sup>rd</sup> Quartile (PSS scores 17–22), N (%)	153 (26.5)	64 (16.7)	53 (27.3)	
4th Quartile (PSS scores 23–50), N (%)	117 (20.2)	64 (16.7)	53 (27.3)	
Psychological Testing				
Depressive symptoms	64 (11.1)	33 (8.6)	31 (16.0)	0.008
WRAT3 Mean (± SD)	12.04 (2.1)	12.10 (2.0)	11.93 (2.2)	0.34
Free Recall Mean (± SD)	30.5 (6.7)	30.5 (6.9)	30.6 (6.2)	0.87
VIQ Mean (± SD)	109.2 (16.4)	110.6 (16.5)	106.4 (15.8)	0.012
FAS Mean (± SD)	35.6 (12.9)	35.9 (13.0)	34.9 (12.6)	0.38
Trails B Mean (± SD)	142.3 (73.1)	141.4 (73.7)	144.1 (72.2)	0.67
History of Medical Illnesses, N (%)				
Hypertension	374 (64.7)	238 (62.0)	136 (70.1)	0.054
Myocardial infarction	39 (6.8)	23 (6.0)	16 (8.3)	0.31
Stroke	44 (7.6)	28 (7.3)	16 (8.3)	0.68
Diabetes	115 (19.9)	68 (17.7)	47 (24.2)	0.064
Angina	32 (5.5)	18 (4.6)	14 (7.2)	0.21
Osteoarthritis	377 (65.2)	220 (57.3)	157 (80.9)	< 0.001
COPD	44 (7.6)	19 (5.0)	25 (12.9)	0.001
Congestive Heart Failure	22 (3.8)	11 (2.9)	11 (5.7)	0.096

Note: All data presented in mean (standard deviation), unless otherwise specified. Continuous variables analyzed by ANOVA; categorical variables analyzed by Pearson chi-square test or Fisher's exact test. P-values refer to comparisons between pain intensity levels. PSS: perceived stress scale. Depressive symptoms were evaluated using the 15- item Geriatric Depression Scale. VIQ: verbal IQ. COPD: Chronic Obstructive Pulmonary Disease. WRAT3 is measured by the Wide Range Achievement Test. Free Recall is measured by the Free and Cued Selective Reminding Test. Verbal IQ score is a summary measure drawn from the Wechsler Adult Intelligence Scale - Third Edition.

#### Table 2

#### Demographic and medical characteristics of subjects by presence of pain interference

Variables	Total (n=578)	No/mild Pain Interference (n=481) 83.2%	Moderate/Severe Pain Interference (n=97) 16.8%	p-value
Demographics				
Age Mean (± SD)	78.8 (5.4)	78.9 (5.5)	78.4 (5.4)	0.48
Female, N (%)	367 (63.5)	301 (62.6)	66 (68.0)	0.31
Race - African American, N (%)	168 (29.1)	137 (28.5)	31 (32.0)	0.17
Race - Other, N (%)	48 (8.3)	36 (7.5)	12 (12.4)	
Lifestyle, N (%)				
Non-smoker	244 (44.4)	205 (44.9)	39 (42.4)	
Former smoker	279 (50.8)	230 (50.3)	49 (53.3)	0.87
Current Smoker	26 (4.7)	22 (4.8)	4 (4.4)	
Alcohol – 1 <sup>st</sup> tertile	158 (28.7)	125 (27.2)	33 (36.3)	0.032
Alcohol – 2nd tertile	181 (32.9)	147 (32.0)	34 (37.4)	
Alcohol – 3 <sup>rd</sup> tertile	211 (38.4)	187 (40.7)	24 (26.4)	
Perceived Stress				
PSS Mean (± SD)	16.4 (7.7)	15.7 (7.6)	19.4 (7.3)	< 0.001
1st Quartile (PSS scores 0–11), N (%)	163 (28.2)	152 (31.6)	11 (11.3)	
2 <sup>nd</sup> Quartile (PSS scores 12–16), N (%)	145 (25.1)	120 (25.0)	25 (25.8)	< 0.001
3rd Quartile (PSS scores 17-22), N (%)	153 (26.5)	125 (26.0)	28 (28.9)	
4th Quartile (PSS scores 23-50), N (%)	117 (20.2)	84 (17.5)	33 (34.0)	
Psychological Testing				
Depressive Symptoms	64 (11.1)	40 (8.3)	24 (24.7)	< 0.001
WRAT3 Mean (± SD)	12.04 (2.1)	12.13 (2.0)	11.62 (2.4)	0.027
Free Recall Mean (± SD)	30.5 (6.7)	30.7 (6.7)	29.5 (6.3)	0.11
VIQ Mean (± SD)	109.2 (16.4)	110.0 (16.3)	105.5 (16.5)	0.032
FAS Mean (± SD)	35.6 (12.9)	35.7 (12.5)	34.8 (1.5)	0.57
Trails B Mean (± SD)	142.3 (73.1)	138.84 (71.0)	159.0 (81.3)	0.01
History of Medical Illnesses, N (%)				
Hypertension	374 (64.7)	307 (63.8)	67 (69.1)	0.32
Myocardial infarction	39 (6.8)	29 (6.0)	10 (10.3)	0.13
Stroke	44 (7.6)	34 (7.1)	10 (10.3)	0.27
Diabetes	115 (19.9)	88 (18.3)	27 (27.8)	0.032
Angina	32 (5.5)	25 (5.2)	7 (7.2)	0.43
Osteoarthritis	377 (65.2)	297 (61.8)	80 (84.5)	< 0.001
COPD	44 (7.6)	27 (5.6)	17 (17.5)	< 0.001
Congestive Heart Failure	22 (3.8)	16 (3.3)	6 (6.2)	0.18

Note: All data presented in mean (standard deviation), unless otherwise specified. Continuous variables analyzed by ANOVA; categorical variables analyzed by Pearson chi-square test or Fisher's exact test. *P*-values refer to comparisons between pain intensity levels. PSS: perceived stress scale. Depressive symptoms were evaluated using the 15- item Geriatric Depression Scale. VIQ: verbal IQ. COPD: Chronic Obstructive Pulmonary Disease. WRAT3 is measured by the Wide Range Achievement Test. Free Recall is measured by the Free and Cued Selective Reminding Test. Verbal IQ score is a summary measure drawn from the Wechsler Adult Intelligence Scale - Third Edition.

#### Table 3

Logistic regression models for pain intensity (no pain/mild pain intensity vs. moderate/severe pain intensity)

	PSS categori	zed	PSS continuous	
Variables	OR (95% CI)	p-value	OR (95% CI)	p-value
PSS 1st Quartile (scores 0-11)	1 (N/A)	N/A	-	-
PSS 2 <sup>nd</sup> Quartile (scores 12–16)	2.12 (1.06-4.21)	0.033	-	-
PSS 3 <sup>rd</sup> Quartile (scores 17–22)	2.47 (1.24-4.92)	0.010	-	-
PSS 4 <sup>th</sup> Quartile (scores 23–50)	3.24 (1.61–6.52)	0.001	-	-
PSS continuous (per unit)	-	-	1.05 (1.02–1.08)	0.003
Age years	0.96 (0.92–1.00)	0.045	0.96 (0.92–1.00)	0.047
Sex	1.42 (0.85–2.37)	0.18	1.37 (0.82–2.27)	0.23
Race - African American	1.19 (0.67–2.12)	0.56	1.22 (0.69–2.16)	0.50
Race - Other	0.82 (0.29–2.28)	0.71	0.83 (0.30-2.28)	0.72
Alcohol - 2nd tertile	0.92 (0.53-1.61)	0.78	0.96 (0.55–1.67)	0.89
Alcohol – 3 <sup>rd</sup> quartile	0.55 (0.31-1.01)	0.053	0.57 (0.31-1.03)	0.061
Depressive symptoms	1.01 (0.83–1.23)	0.09	1.02 (0.84–1.25)	0.82
VIQ	1.00 (0.98–1.01)	0.70	1.00 (0.98–1.01)	0.64
Hypertension	1.28 (0.78–2.11)	0.33	1.26 (0.77–2.07)	0.37
Diabetes	1.10 (0.63–1.95)	0.73	1.12 (0.64–1.97)	0.70
Osteoarthritis	2.57 (1.53-4.30)	< 0.001	2.61 (1.56-4.35)	< 0.001
Angina	0.64 (0.23–1.82)	0.41	0.62 (0.22–1.74)	0.37
COPD	1.81 (0.78-4.21)	0.17	1.86 (0.81–4.27)	0.15
Congestive Heart Failure	3.87 (1.20–12.46)	0.023	4.22 (1.32–13.55)	0.015

PSS: perceived stress scale; COPD: Chronic Obstructive Pulmonary Disease

OR of participant characteristics predicting pain intensity in our logistic regression models, (on the left using PSS categorized or on the right PSS continuous) are presented with confidence intervals and p-values. Models include demographic factors and medical history variables with bivariate testing results with pain intensity 0.25. Caucasian race and first tertile of alcohol consumption were used for reference groups, respectively.