Suicide is a leading cause of death, yet it remains a challenge to predict and prevent. In 2012, suicide was the tenth leading cause of death in the United States, accounting for 40,600 deaths.1 Further, 2012 marks the first time in history that U.S. deaths by suicide have exceeded 40,000 in a year.2 Suicide is a particularly large problem for older adults: although older adults only made up about 13.7% of the population in 2012, older adults represented 16.4% of suicides.3 Suicidal intent, which refers to the degree to which one desires to end his/her life by suicidal behavior,4 is more common in older adults than in the rest of the population, and attempts made by older adults are more likely to be fatal.4 To illustrate this, there are 100–200 attempts for every one youth suicide and 4 attempts for every one older adult suicide.5 Thus, older adult suicide attempts are much more likely to be fatal than younger adult suicide attempts. Research has identified many risk factors for older adult suicide including the loss of independence, social isolation, mental illness, and decline in physical health.4,6–8 However, despite having identified these risk factors, clinicians are still very poor at predicting suicide. Two factors that have emerged as strong candidates as suicide risk factors in recent years are insomnia symptoms and nightmares.

**Insomnia Symptoms and Suicide**

Many studies have found an association between insomnia symptoms and suicidal behavior in both community and clinical samples.9–17 For instance, in a sample of individuals who suffer from depression and insomnia being treated by hypnotic and selective serotonin reuptake inhibitor (SSRI) medication, insomnia is correlated with suicidal thinking, suggesting a link between insomnia and suicidal ideation in a clinical population.11 Although research has clearly demonstrated that insomnia symptoms are associated with suicidal behavior, findings regarding whether this association is independent of other psychopathology are mixed. Pigeon and colleagues found that among Veteran suicide decedents, Veterans with sleep disturbance died in closer proximity to their last visit to the Veterans Affairs (VA) than those who did not have sleep disturbance, even after statistically accounting for mental health disorders.13 Thus, sleep disturbances may relate to the level of imminence of suicidal behaviors. Similarly, insomnia symptoms were associated with suicidal ideation cross-sectionally after...
controlling for symptoms of depression, hopelessness, anxiety, Posttraumatic Stress Disorder (PTSD) diagnosis, and substance abuse. Additionally, a recent study among older adults found that poor sleep quality predicted future death by suicide after controlling for baseline depressive symptoms. However, other studies have found that the association between insomnia and suicide risk is mediated by other psychopathology. A recent research study found the relation between insomnia and suicidal ideation to be mediated by nightmares and dysfunctional attitudes about sleep. Additionally, our research team has also found relations between insomnia symptoms and suicide to be mediated by symptoms of psychopathology among young adults and older adults. However, an emerging factor that may help explain this discrepancy in the literature is insomnia duration. Our recent research has found that insomnia duration is associated with suicide risk independent of nightmare duration and current symptoms of depression, anxiety, PTSD, insomnia, and nightmares. However, research has yet to examine these relations among older adults.

Nightmares and Suicide

Nightmares are associated with suicide, and several studies among younger adults have found that nightmares are associated with suicidal ideation or suicide risk independent of symptoms of psychopathology. However, the only study we are aware of looking at nightmares and suicidal ideation among older adults found that the relation between nightmares and suicidal ideation was mediated by both insomnia symptoms and depressive symptoms. Nightmares have also been shown to be associated with an increased risk of suicide attempts. In a sample of 165 suicide attempters, individuals who reported frequent nightmares were over four times more likely to have at least one repeated suicide attempt at a 2-year follow-up after adjusting for the effects of depression, anxiety, PTSD, and substance use disorders. Similar to insomnia symptoms, recent research suggests that the sleep disorder duration, or how long one has had insomnia or nightmares, may be more important than current symptomology. As was the case with insomnia, nightmare duration was also associated with suicidal ideation independent of insomnia duration and current symptoms of insomnia, nightmares, anxiety, depression, and PTSD. Because sleep disturbances and nightmares are associated with depression and PTSD, it is important to examine relations between sleep disturbances and suicide while accounting for symptoms of depression (e.g., symptoms of anhedonia) and PTSD.

Examining the Theoretical Basis of the Association of Sleep and Suicide

As outlined above, a substantial amount of research in recent years has focused on trying to understand the mechanism through which sleep disorders confer suicide risk by statistically controlling for the effects of other forms of psychopathology. However, until recently, the association between sleep problems and suicide risk had never been tested within a theoretical framework. In a recent paper, we examined whether the interpersonal-psychological theory of suicide (IPTS), a well-known and empirically supported model of suicidal behavior, may explain the association between sleep disturbance and suicide risk. The theory consists of 3 constructs: perceived burdensomeness, thwarted belongingness, and acquired capability to enact lethal self-harm. The interaction of these three constructs indicated the greatest risk for suicide attempts versus suicidal ideation in a clinical sample of young adults with suicidality. In our study, we found that nightmares were independently associated with suicide risk and past suicide attempts in 2 independent data sets; however, our data were mixed regarding insomnia symptoms and suicide, with the IPTS mediating the relation in one sample but not the other. Thus, it appears that the IPTS cannot fully explain the mechanism by which nightmares confer suicide risk, but that the theory may be useful in explaining the association between insomnia symptoms and suicidal behavior.

Statement of the Problem

There are several significant gaps in the empirical literature that warrant discussion and further examination. First, there is a paucity of studies examining the theoretical basis of the association between sleep disturbance and suicidality. This is important, as understanding how sleep disturbances lead to suicidality will likely aid clinicians in detecting and treating suicide risk. A second substantial limitation is that very few studies have examined sleep and suicide in older adults despite older adults having higher rates of sleep disturbances and suicide. Additionally, older adults experience an earlier circadian sleep-wake cycle, decreased total sleep time, increased wakefulness after onset of sleep, decreased sleep efficiency, and less slow wave sleep. Because of these differences in sleep and suicidal behavior, it cannot be assumed that the relations among sleep and suicide that occur in young people will be the same in older adults. Thus, there is a need for more research examining sleep and suicide among older adults.

Present Study

Although the duration of sleep dysfunction has been examined in relation to suicidal ideation in a young adult sample, and some research has been conducted examining the relations between sleep and suicide in older adults, substantial gaps in the literature remain. Specifically, research has yet to examine the impact of sleep disturbance duration on suicide risk among older adults. Further, the association between sleep disturbances and suicide risk among older adults has yet to be examined in relation to a leading suicide theory. To address these limitations in the empirical literature, the current study will examine the specific relations among duration of insomnia and nightmares and suicide risk, including examining these relations within the IPTS framework.

In line with previous findings related to younger adults, we hypothesize that insomnia duration and nightmare duration will be positively associated with suicide risk independent of current levels of insomnia symptoms and nightmare severity. Our second hypothesis is that insomnia duration and nightmare duration will be positively associated with suicide risk independent of current symptoms of insomnia, nightmares, PTSD, and anhedonia. Our third hypothesis is that insomnia duration and nightmare duration will be associated with suicide risk independent of current symptoms of insomnia,
nightmares, and the IPTS (i.e., thwarted belongingness, burdensomeness, acquired capability, and relevant interactions).12

METHODS

Participants

The current study consists of a sub-sample of 167 individuals aged 55 or older who were obtained through combining data from 2 larger data collections (age 18+) utilizing Amazon.com’s Mechanical Turk (mTurk), an online venue on which people can participate in online opportunities for nominal payments. Sample demographics from mTurk are of equal or greater representativeness and diversity than conventional samples.31 Further, and of importance to this study, samples collected through mTurk match or exceed incidence of clinical symptoms such as depression and anxiety and exposure to trauma when compared to the levels in the general population.32 Of the 167 respondents, 124 (74.3%) were female. The sample ranged in age from 55 to 75 years (mean = 60.6, SD = 4.94). The majority (91.6%) of participants were Caucasian. Participants received nominal payment (i.e., $0.25) for completion of the study; however, some analyses regarding motivation to complete mTurk studies suggest internal motivations, such as for entertainment, for participating in online studies.31 Further, the potential to reach Mechanical Turk master status may influence mTurk participants’ completion of studies. In order to work toward this status, workers are encouraged to submit Human Intelligence Tasks, or HITs, with accuracy on a breadth of subjects and from a variety of requesters; further, this status is not something a worker can request; workers’ performance can distinguish them for master status.33 Thus, participants may have completed our study for low pay in hopes of reaching master status.

All participants were from the United States, with 43 states represented. Most commonly, participants were from Florida (10.2%), California (7.2%), and Tennessee (6.6%). Further, participants from all regions of the US were represented: 13.2% from the Northeast, 25.1% from the Midwest, 19.2% from the West, and 41.3% from the South; these proportions are similar to those of the United States population in 2014 (17.6% in the Northeast, 21.2% in the Midwest, 23.6% in the West, and 37.6% in the South).34 Though previous analyses have found a representative split of sex on mTurk (55% female sample),31 our study had an unrepresentative majority of females who completed the study. Our study may have self-selected a majority women because of the subject matters of suicide, insomnia, and depression.

Measures

The insomnia severity index (ISI)35 assesses severity of insomnia symptoms over the past 2 weeks. This questionnaire has 7 items with answer options that range from 0 to 4 points, for a possible maximum of 28 points.36 Scores > 15 indicate a moderate to severe level of insomnia.35 Over 3 months, the ISI has demonstrated sufficient test-retest reliability; further, the ISI has shown concurrent validity with polysomnography and patients’ sleep diaries.35,36 In the current study the ISI demonstrated acceptable internal consistency (α = 0.90). Means and standard deviations for all measures, including the ISI, can be found in Table 1.

The Disturbing Dreams and Nightmares Severity Index (DDNSI) was used to assess frequency and severity of nightmares over the past year.35,36 The DDNSI assesses frequency based on the number of nights per week with nightmares and the number of nightmares participants endorse having each week (maximum of 14 nightmares).37 Likert-type scale questions further assessed nightmare severity by measuring how often individuals are awakened by nightmares and nightmare severity and intensity. Higher scores on the DDNSI reflect higher levels of current nightmare difficulties, with scores ≥ 10 indicating the possible presence of a nightmare disorder.38 In the current study the DDNSI demonstrated acceptable internal consistency (α = 0.90).

Insomnia duration was measured by asking the participants “If you have an insomnia problem, how long have you had it for (please specify months and years)?” After converting participant answers, a duration variable of total months of reported insomnia was created.14 Nightmare duration was determined by one item from the DDNSI that does not factor into the total DDNSI score. On this item, participants were asked to provide an estimation of the number of months and/or years they have been experiencing nightmares or disturbing dreams.39 Again, data were converted to reflect the total number of months of reported nightmares or disturbing dreams. These measures of duration of insomnia symptoms and nightmares have been used in a previous study evaluating sleep disturbance duration.14

The Posttraumatic Stress Disorder Checklist-Civilian Version (PCL-CL)40 was used to measure how much trouble associated with PTSD symptoms participants have experienced related to their most significant life stressor. For each of the 17 items, participants could choose from 1 (not at all) up to

<table>
<thead>
<tr>
<th>Measure</th>
<th>Mean</th>
<th>SD</th>
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<tr>
<td>Burdensomeness</td>
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<td>Belongingness</td>
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<td>14.18</td>
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<tr>
<td>Acquired capability</td>
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<td>6.47</td>
</tr>
<tr>
<td>Suicide risk</td>
<td>6.86</td>
<td>2.87</td>
</tr>
</tbody>
</table>

n of 166 for insomnia symptoms, burdensomeness, and belongingness; n of 167 for remaining measures; insomnia symptoms are scores from the ISI; insomnia symptom duration are self-reported lengths of time having experienced insomnia symptoms; nightmares are scores from the DDNSI; nightmare duration are self-reported lengths of time having experienced nightmares; PTSD symptoms are scores of posttraumatic stress disorder symptoms from the PCL-CL; anhedonia are scores from SLIPS; burdensomeness are scores of perceived burdensomeness from INQ; belongingness are scores of thwarted belongingness from INQ; acquired capability are scores from ACSS; suicide risk are scores from SBQ. SD, standard deviation.
ACSS-FAD uses 7 items to assess the level of habituation to fear of death participants exhibit. The ACSS-FAD has demonstrated appropriate convergent and discriminant validity. In the current study the ACSS-FAD demonstrated acceptable internal consistency ($\alpha = 0.81$).

The Specific Loss of Interest and Pleasure Scale (SLIPS) assesses recent changes in symptoms of anhedonia. This measure includes 23 items with ordinal scales that allow individuals to indicate the level of change in interest or pleasure from certain activities from 0 (no change), 1 (some decrease), 2 (marked decline), and 3 (interest/pleasure has never been present). Because items originally scored as 3 inflate scores and relate to trait anhedonia, those items were re-coded as a score of 0, such that the SLIPS score in the current study would capture recent changes in levels of anhedonia. The SLIPS has demonstrated high levels of internal consistency ($\alpha = 0.94$) and strong incremental validity when compared with other measures of anhedonia. In the current study the SLIPS demonstrated acceptable internal consistency ($\alpha = 0.94$).

The Interpersonal Needs Questionnaire (INQ) has 15 items used to assess participant’s level of perceived burdensomeness with 6 items and perceived levels of thwarted belongingness with the remaining 9 items. Based on the IPTS, the INQ examines levels of suicidal desire where higher scores are associated with higher levels of perceived burdensomeness or thwarted belongingness. In the current study both burdensomeness ($\alpha = 0.95$) and belongingness ($\alpha = 0.94$) demonstrated acceptable internal consistency.

The Acquired Capability for Suicide Scale – Fearlessness about Death (ACSS-FAD) is a self-report measure adapted from the original ACSS to differentiate fearlessness about death from the concept of physiological pain tolerance – both of which make up acquired capability within the IPTS. The ACSS-FAD uses 7 items to assess the level of habitation to fear of death participants exhibit. The ACSS-FAD has demonstrated appropriate convergent and discriminant validity. In the current study the ACSS-FAD demonstrated acceptable internal consistency ($\alpha = 0.81$).

The Suicidal Behaviors Questionnaire – Revised (SBQ-R) is a 4-item self-report, derived from the SBQ, that is used to measure suicidal risk levels. A score ranging from 3 to 18 reflects past suicidal thoughts, ideation, threats, and attempts as well as future suicidal behaviors. The SBQ-R has demonstrated acceptable internal consistency ($\alpha = 0.87$, clinical sample; $\alpha = 0.88$, nonclinical sample). In the current study the SBQ-R demonstrated acceptable internal consistency ($\alpha = 0.74$).

**Analyses**

Data were initially collected from 177 individuals aged 55 and older. Steps were taken in order to check and help ensure data quality, as the data were collected via an online survey. As a first step, any participant who did not make it to the demographics questions of the study was removed. Participants who endorsed the same answer for each item (i.e., all 1s, all 2s, all 3s) on the INQ or the ACSS-FAD, both of which have items that are reverse-scored, were removed. After removing participants for these reasons, the current sample included data from 167 individuals.

The SBQ was assessed for distribution normality. No deviation from normality was noted (skew = 0.89, kurtosis = 0.01), so no transformations were necessary. Descriptive statistics and correlations for the current sample can be found in **Table 1** and **Table 2**, respectively. An analysis of variance (ANOVA) was utilized to test for effects of gender. Gender, $F_{1,165} = 0.31$, $p = 0.58$, was not significantly related to suicide risk. Thus, gender was not included as a covariate in our analyses.

**RESULTS**

A multiple linear regression assessed the first hypothesis which stated that insomnia duration and nightmare duration...
are associated with suicide risk independent of current levels of insomnia symptoms and current nightmare severity (see Table 3). The overall regression was significant, n = 165, \( F_{4,161} = 9.56, p < 0.01, R^2 = 0.19 \). Level of current insomnia symptoms, \( \beta = 0.16, p < 0.01 \), and nightmare duration, \( \beta = 0.03, p = 0.03 \), were significantly associated with suicide risk; however, duration of insomnia symptoms and current nightmares were not statistically significant in the model. Thus, our first hypothesis was only partially supported, as duration of insomnia symptoms was not associated with suicide risk.

Using a multiple linear regression, the second hypothesis, which stated that insomnia duration and nightmare duration are associated with suicide risk independent of current insomnia symptoms, nightmares, PTSD symptoms, and symptoms of anhedonia, was tested (see Table 4). Current insomnia symptoms, nightmares, PTSD symptoms, and anhedonic symptoms were added in the first step. Insomnia duration and nightmare duration were added in step two. The overall regression was significant in step one, n = 165, \( F_{4,161} = 18.36, p < 0.01, R^2 = 0.31 \). Initially, only symptoms of anhedonia, \( \beta = 0.12, p < 0.01 \), were associated with suicide risk, consistent with the previous finding that anhedonic symptoms of depression are related to suicidal ideation independent of cognitive and affective symptoms of depression. After adding insomnia duration and nightmare duration to the regression in step two (\( R^2 \) change = 0.03, \( F = 3.44, p = 0.04 \)), symptoms of anhedonia, \( \beta = 0.12, p < 0.01 \), and nightmare duration, \( \beta = 0.03, p = 0.01 \), were the only variables associated with suicide risk. Again, our hypothesis was partially supported. Although nightmare duration was associated with suicide risk independent of pathology and current nightmares, insomnia duration again did not have a significant relation with suicide risk within this model.

To test the third hypothesis, which predicted that duration of insomnia and nightmares would be associated with risk of suicide independent from current insomnia symptoms, current nightmares, and the IPTS, a multiple linear regression was utilized (see Table 5). In the first step, the constructs and interactions of the IPTS were added; in the second step, current insomnia symptoms and current nightmares were added in. In the third and final step, nightmare duration and insomnia duration were added to the model. The overall regression was significant in step one, n = 164, \( F_{1,157} = 12.31, p < 0.01, R^2 = 0.35 \).

<table>
<thead>
<tr>
<th>Table 3—Insomnia duration and nightmare duration predicting suicide risk, controlling for current insomnia and nightmare symptoms.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Predictors</td>
</tr>
<tr>
<td>Model</td>
</tr>
<tr>
<td>Insomnia Symptoms</td>
</tr>
<tr>
<td>Nightmares</td>
</tr>
<tr>
<td>Insomnia Duration</td>
</tr>
<tr>
<td>Nightmare Duration</td>
</tr>
</tbody>
</table>

Insomnia symptoms are scores from the ISI; insomnia symptom duration are self-reported lengths of time having experienced insomnia symptoms; Nightmares are scores from the DDNSI; nightmare duration are self-reported lengths of time having experienced nightmares.

Thwarted belongingness, \( \beta = 0.09, p < 0.01 \), was positively associated with suicide risk; however, no other part of the IPTS was significantly associated with suicide risk. After adding insomnia severity and current nightmares to the model in step two, thwarted belongingness, \( \beta = 0.08, p < 0.01 \), and current nightmares, \( \beta = 0.07, p = 0.04 \), were positively associated with suicide risk, but no other aspect of the IPTS or current insomnia symptoms were associated with suicide risk. In the final step of the regression (\( R^2 \) change = 0.03, \( F = 4.21, p = 0.02 \)), thwarted belongingness, \( \beta = 0.08, p < 0.01 \), and duration of nightmares, \( \beta = 0.03, p < 0.01 \), were positively associated with suicide risk. Thus, our hypothesis was partially supported in that nightmare duration is related to suicide risk independent of the IPTS, current insomnia symptoms, and current nightmares. However, the duration of insomnia symptoms was not significantly associated with suicide risk independent of IPTS, current insomnia, and nightmares and nightmare duration.

**DISCUSSION**

The current study is significant as it is the first study to examine whether insomnia and nightmare duration are related to suicide risk among older adults, as well as the first study to examine sleep disturbance duration, suicide risk, and the IPTS in any age group. Although prior findings suggest that duration of sleep disturbance is important in level of suicide risk in young people, it was important to examine the role of insomnia and nightmare duration in older adults because sleep disturbances and suicide risk in older adults has presented differently than that for younger or mixed-age samples.

We found that insomnia symptoms (but not duration of insomnia symptoms) and nightmare duration (but not the current severity of nightmares) were significantly associated with...
suicide risk. This finding is partially in line with previous research that found insomnia symptoms, but not current nightmares, to be related to suicide risk in older adults. However, once symptoms of anhedonia and PTSD (in one regression) were accounted for, current insomnia symptoms no longer showed a significant relation with suicide risk. Though previous research suggested that current nightmares in older adults may not be relevant in suicide risk assessment, which was replicated in the present study, our findings suggest that the duration of nightmares explains a significant proportion of variance in suicide risk level in older adults. In each of our analyses, nightmare duration showed a relation with suicide risk independent of other features of our models; however, insomnia symptom duration did not hold a significant relation with suicide risk for any of our models. Though duration of insomnia symptoms was associated with suicide risk in a college-aged sample, that relation was not seen in the current sample of older adults.

The current paper is the first to suggest that duration of nightmares is important in understanding suicide risk among older adults. This finding has clinical relevance in the detection of suicide risk, and we believe that the assessment of suicide risk among older adults who report a long history of nightmares is warranted.

Though the current study did not ultimately find relations among current insomnia symptoms, duration of insomnia, and suicide risk independent of the IPTS, the current study builds off the previous finding that insomnia and nightmares are associated with suicide risk independent of the IPTS in younger adults. In older adults, current nightmares and not current insomnia symptoms relate to suicide risk independent of the IPTS; however, nightmare duration mediates the relation between nightmares and suicide risk, suggesting that how long an individual has been experiencing nightmares is more predictive of suicide risk than the current presence of nightmares. These findings suggest that the IPTS cannot fully explain why nightmare duration increases suicide risk among older adults and that further research in this area is warranted.

Lastly, our current findings along with our previous findings among younger adults suggest that failing to treat nightmares may increase one’s risk for suicide. In assessing patients with suicide risk, it is pertinent to investigate not only current presence of nightmares, but also their history of nightmares. Further, suicide risk should be assessed in patients who report a history of nightmares, as the duration of nightmares is predictive of suicide over and above the typically assessed factors. Given that duration of nightmares is related to suicide risk, treating nightmares early on may prevent suicide risk from increasing. However, to date, no research has found that treating nightmares reduces suicide risk. Based upon our findings, we believe that this research would greatly benefit the literature and is clearly warranted.

**Limitations**

The current study has a few limitations that must be considered. The first limitation is that the data comprising our sample was combined from two separate studies conducted on mTurk, so it was not a continuous data collection. However, our analyses failed to find any participants who took the study more than once, so we believe the responses meet the statistical assumption of independence. Second, our sample included individuals ages 55 and older, whereas many older adult studies include only those over age 60 or 65. However, we do not believe this to detract from the current findings, as other studies examining insomnia and older adults have used individuals as young as age 50. Additionally, a limitation of our current sample is that there was an uneven gender split; however, given that there were no differences in our outcome variable based on gender, the current findings are generalizable and valid. The current sample is not a clinical sample; thus, the findings may

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**Table 5—Insomnia duration and nightmare duration predicting suicide risk, controlling for symptoms of IPTS and insomnia and nightmare symptoms.**

<table>
<thead>
<tr>
<th>Predictors</th>
<th>$R^2$</th>
<th>$\beta$</th>
<th>t</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Step 1</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Burden × belong</td>
<td>0.04</td>
<td>0.82</td>
<td>0.41</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Burden × capability</td>
<td>&lt; 0.01</td>
<td>1.05</td>
<td>0.30</td>
<td></td>
</tr>
<tr>
<td>Belong × capability</td>
<td>&lt; 0.01</td>
<td>-0.05</td>
<td>0.96</td>
<td></td>
</tr>
<tr>
<td>Belong × belong × capability</td>
<td>&lt; 0.01</td>
<td>-0.76</td>
<td>0.45</td>
<td></td>
</tr>
<tr>
<td>Nightmares</td>
<td>0.07</td>
<td>2.05</td>
<td>0.04</td>
<td></td>
</tr>
<tr>
<td><strong>Step 2</strong></td>
<td>0.39</td>
<td>0.01</td>
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<td></td>
</tr>
<tr>
<td>Burden × belong</td>
<td>0.03</td>
<td>0.62</td>
<td>0.54</td>
<td></td>
</tr>
<tr>
<td>Burden × capability</td>
<td>0.08</td>
<td>4.03</td>
<td>&lt; 0.01</td>
<td></td>
</tr>
<tr>
<td>Belong × belong</td>
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<td>-0.60</td>
<td>0.55</td>
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<tr>
<td>Belong × capability</td>
<td>&lt; 0.01</td>
<td>1.09</td>
<td>0.28</td>
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<tr>
<td>Belong × belong × capability</td>
<td>&lt; 0.01</td>
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<tr>
<td>Insomnia symptoms</td>
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<td>1.49</td>
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<tr>
<td>Nightmares</td>
<td>0.07</td>
<td>2.05</td>
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<tr>
<td><strong>Step 3</strong></td>
<td>0.42</td>
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<tr>
<td>Burden × belong</td>
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<tr>
<td>Burden × capability</td>
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<tr>
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</table>

Step 1: burdensomeness are scores of perceived burdensomeness from INQ; belongingness are scores of thwarted belongingness from INQ; capability are scores from the ACSS; burden × belong is the interaction between burdensomeness and belongingness; burden × capability is the interaction between burdensomeness and capability; belong × capability is the interaction between belongingness and capability; burden × belong × capability is the interaction between burdensomeness, belongingness, and capability. Step 2: insomnia symptoms are scores from the ISI; nightmares are scores from the DDNSI. Step 3: insomnia duration are self-reported lengths of time having experienced insomnia symptoms; Nightmare Duration are self-reported lengths of time having experienced nightmares.
not generalize to those at severe risk of suicide. Previous findings, however, have indicated that respondents to mTurk studies exhibit similar or higher levels of anxiety, depression, and exposure to trauma as compared to individuals in the general population.42 Fourth, the current sample relies on retrospective reporting as a measure of duration of insomnia symptoms and nightmares; thus, the data may be less reliable than sleep diaries, actigraphy, or polysomnography for insomnia or nightmare diaries for nightmares. However, the ISI has demonstrated concurrent validity with polysomnography and sleep diaries, suggesting that participants’ reports are representative of their current sleep.35,36 A further limitation is that other sleep disorders, such as obstructive sleep apnea, are prevalent among older adults and could not be controlled for in the current study. Although obstructive sleep apnea cannot be ruled out as a cause of sleep disturbance in the current sample, our results indicate that the resulting subjective distress associated with sleep problems can be associated with risk for suicide, though this relation is no longer significant when symptoms of anhedonia and PTSD are statistically removed. Despite this mediation, the literature would benefit from future studies that include sleep studies to examine insomnia and obstructive sleep apnea separately. Additionally, although the DDNSI is a widely used, validated measure of nightmares, it does not include a definition of nightmares and thus relies on respondents’ interpretation of whether what they are experiencing is a nightmare. Therefore, it is possible that some respondents affirmatively based on night terrors instead of nightmares. However, given that the prevalence rate of night terrors in adults 45 and older ranges from 1% to 2.3%,31 the current findings are unlikely to be impacted by individuals who may have answered the questions based on night terrors.

**Future Directions**

The current findings indicate that insomnia symptoms and nightmare duration are clinically relevant and should be evaluated in older adults. Further, if endorsed, further screening of mental disorders and suicide risk is warranted. Future longitudinal studies may help further elucidate the timing and mechanism of insomnia and nightmares in relation to suicide risk. Finally, assessing whether nightmare treatment for older adults with nightmares reduces the risk for suicide in those individuals is pertinent. With increased knowledge of risk factors for suicide in older adults, it is important to determine what may eliminate or decrease those risk factors effectively.

**ABBREVIATIONS**

ACSS-FAD, Acquired Capability for Suicide Scale – Fearlessness about Death
ANOVA, analysis of variance
DDNSI, Disturbing Dreams and Nightmares Severity Index
HIT, Human Intelligence Tasks
INQ, Interpersonal Needs Questionnaire
IPTS, Interpersonal-Psychological Theory of Suicide
ISI, insomnia severity index
mTurk, Amazon.com’s Mechanical Turk
PCL-CL, Posttraumatic Stress Disorder Checklist – Civilian Version
PTSD, posttraumatic stress disorder
SBQ-R, Suicidal Behaviors Questionnaire – Revised
SLIPS, Specific Loss of Interest and Pleasure Scale
SSRI, Selective Serotonin Reuptake Inhibitor
VA, Veterans Affairs

**REFERENCES**

2. McIntosh JL. Personal communication. 2014 December 23.