Measuring anxiety in the elderly: psychometric properties of the state trait inventory of cognitive and somatic anxiety (STICSA) in an elderly Italian sample

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ABSTRACT

Background: Despite its increasing personal and societal impact, assessment of late-life anxiety has received relatively little attention in psychiatric research. Differential symptom presentation and physical comorbidities among the elderly, relative to younger cohorts creates a need for anxiety measures that are psychometrically validated in the elderly.

Methods: The present study examined the factor structure and discriminant validity of the state-trait inventory for cognitive and somatic anxiety (STICSA) in a sample of Italian middle-aged and older adults. Participants were 396 community-dwelling middle-aged (50–64 years) and older (≥65 years) adults. In addition to the STICSA, participants completed two depression measures and a general well-being survey with physical and mental health subscales.

Results: Factor analysis supported the validity of both state–trait and cognitive–somatic distinctions underlying the STICSA, all dimensions exhibited excellent internal consistency (Cronbach’s α coefficients ≥ 0.86), and correlations with depression measures provided limited evidence for differentiation of anxious and depressive symptoms. The STICSA also showed evidence of discriminating anxious symptoms from physical health symptoms, a particularly relevant feature of a valid anxiety measure in elderly samples.

Conclusions: The STICSA appears to be a valid measure of cognitive and somatic anxiety in the elderly.

Key words: anxiety, factor analysis, geriatrics, physical health, depression

Introduction

Late-life anxiety is an increasingly relevant psychiatric condition. With increasing numbers of older adults, anxiety will become an increasing cause of health care utilization contributing to elevated personal and societal costs (Wolitzky-Taylor et al., 2010). Unfortunately, the detection of anxiety disorders in late life is complicated by cognitive decline, newly emergent changes in life circumstances, high medical comorbidity, and a symptom presentation that is markedly different from younger age groups (Magni and DeLeo, 1984; Kogan et al., 2000; Cully et al., 2006; Seignourel et al., 2008; Wolitzky-Taylor et al., 2010; Therrien and Hunsley, 2012). For many of the above reasons, anxiety disorders in late life are more likely to go unnoticed and untreated relative to anxiety in younger populations. Assessing the presence and severity of clinical anxiety in the elderly is an important challenge for researchers and clinicians alike. However, relatively little is known about the assessment of anxiety in older adults (Ayers et al., 2007).

Self-report measures are by far the most common method of assessment of anxiety (Alwahhabi, 2003; Dennis et al., 2007). Self-report inventories are easily administered and limit patient/participant burden. Approximately 12 different anxiety measures have been identified as commonly used for the assessment of anxiety in older adults (Therrien and Hunsley, 2012). Most of these measures were originally developed and validated in college samples and therefore lack norms and sufficient psychometric evidence for use with older
adults. Additionally, many of these measures fail to capture the unique phenomenology of anxiety in the elderly (Edelstein et al., 2008). For example, an overabundance of somatic items can obfuscate differences between medical and psychological causes of anxiety in this population (Therrien and Hunsley, 2012). Moreover, instruments developed in younger populations should be normed with samples of older adults, providing age-appropriate norms for this population. It is well known that the clinical expression and severity of anxiety symptoms may change with age (Wolitzky-Taylor et al., 2010). Another important issue for the assessment of late-life anxiety is the frequent comorbidity with other mental disorders (Kogan et al., 2000). Particularly, the high comorbidity of anxiety and depression among younger adults (Mineka et al., 1998) becomes even more impressive in older adults, with approximately half of depressed older adults also meeting criteria for an anxiety disorder (Beekman et al., 2000). Thus, anxiety instruments able to disentangle symptoms of anxiety and depression are especially needed among elderly samples.

To address many of these limitations, some researchers have investigated the psychometric properties of existing anxiety questionnaires in older adults (such as the abbreviated Penn State Worry Questionnaire (Hopko et al., 2003; Crittendon and Hopko, 2006; Nuevo et al., 2007; Wuthrich et al., 2014) and the Geriatric Anxiety Inventory-short form (Byrne and Pachana, 2011)). Additionally, new measures have been created specifically for use with older adults (e.g. Geriatric Anxiety Inventory; Pachana et al., 2007, Adult Manifest Anxiety Scale-Elderly Version; Reynolds et al., 2003, Geriatric Anxiety Scale; Segal et al., 2010).

Despite the strengths of modified and newly developed measures for use in older adults (e.g. adequate internal consistency, strong convergent validity), these measures present several limitations which may restrict their clinical utility in specific groups of older adults. For example, some of these scales exhibit poor discrimination between anxiety and depression or health-related variables. Some of the aforementioned measures use dichotomous response formats, which can provide less information (when compared to dimensional rating) regarding the severity of symptoms. Finally, many anxiety measures exhibit poor reliability for items assessing physical or sleep disturbances in the elderly, potentially due to difficulty in differentiating anxiety from medical disorders and elevated prevalence of sleep difficulties in later life (Foley et al., 2004; Crittendon and Hopko, 2006; Andrew and Dulin, 2007; Byrne et al., 2008; Diefenbach et al., 2009; Segal et al., 2010; Yochim et al., 2011; Gould et al., 2014; Johnco et al., 2014; Mueller, 2014). Moreover, geriatric anxiety measures cannot be used for investigating anxiety in younger age groups, a particular challenge for studies investigating anxiety across the lifespan. While age-specific measures have their benefits, the availability of general anxiety questionnaires with age-based norms can be preferable to age-specific instruments. Consequently, examination of the psychometric properties in currently available measures of anxiety when used with older adults (in lieu of revising these instruments for age-specific use) is helpful for many of the aforementioned reasons.

The STICSA (Ree et al., 2008) was recently developed to overcome several limitations issues associated with Spielberger’s State-Trait Anxiety Inventory (STAI; Spielberger et al., 1983; Kennedy et al., 2001; Therrien and Hunsley, 2012). To date, the STICSA has been shown to have excellent psychometric properties in both nonclinical and clinical samples of adults (Gros et al., 2007; Ree et al., 2008; Gros et al., 2010; Van Dam et al., 2013) and to differentiate anxiety from depression better than other commonly used anxiety measures (e.g. the STAI; Gros et al., 2007). The fact that the STICSA assesses cognitive and somatic dimensions of anxiety, and particularly that it can differentiate anxiety and depression, makes it an attractive tool for assessing anxiety in the elderly.

To our knowledge, no previous studies have investigated the psychometric properties of the STICSA in older adults. Accordingly, our aims were to assess the factor structure and discriminant validity of the Italian version of the STICSA in a sample of middle-aged and older adults. We predicted confirmation of the original four-factor structure (e.g. state vs. trait and cognitive vs. somatic dimensions). Concurrent and discriminant validity were investigated by administering two self-report measures of depression and a measure of quality of life, the latter assessing mental and physical health functioning. Our hypotheses were that: (1) cognitive and somatic subscales of the STICSA are correlated reciprocally more strongly than they are correlated with measures of depression; (2) the STICSA correlates more strongly with mental health functioning than with physical health functioning, indicating the ability of STICSA scores to discriminate between anxiety and physical health issues.

Additionally, we assessed whether age (older adults – 65 years and older – vs. middle-aged adults – 50–64 years old) and sex had an effect on STICSA scores. While there are no clear-cut boundaries between adulthood and late adulthood, 65 years is a common cutoff used for “old age” (Federal Interagency Forum on Aging-Related Statistics, 2012; World Health Organisation, 2014),

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Table 1. Descriptive statistics of the sample (N = 396)

<table>
<thead>
<tr>
<th></th>
<th>Count</th>
<th>Percentage</th>
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<tbody>
<tr>
<td>Women</td>
<td>200</td>
<td>50.5%</td>
</tr>
<tr>
<td>Age – M±SD (range) years</td>
<td>69.04 ± 8.41 (51/94)</td>
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<tr>
<td>Marital status</td>
<td></td>
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<tr>
<td>Married</td>
<td>288</td>
<td>72.7%</td>
</tr>
<tr>
<td>Divorced</td>
<td>23</td>
<td>5.8%</td>
</tr>
<tr>
<td>Widowed</td>
<td>62</td>
<td>15.7%</td>
</tr>
<tr>
<td>Single</td>
<td>23</td>
<td>5.8%</td>
</tr>
<tr>
<td>Educational attainment – M±SD years</td>
<td>8.12 ± 4.43</td>
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</tr>
<tr>
<td>Education ≤ 8 years</td>
<td>265</td>
<td>66.9%</td>
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<tr>
<td>Retired</td>
<td>230</td>
<td>58.1%</td>
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<tr>
<td>GDS – M±SD</td>
<td>8.57 ± 5.55</td>
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<tr>
<td>TDI – M±SD</td>
<td>21.68 ± 11.88</td>
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<tr>
<td>SF-12 PCS – M±SD</td>
<td>42.10 ± 9.72</td>
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<tr>
<td>SF-12 MCS – M±SD</td>
<td>51.13 ± 8.62</td>
<td></td>
</tr>
</tbody>
</table>

Note. TDI = teate depression inventory; GDS = geriatric depression scale; PCS = SF-12 physical composite score; MCS = SF-12 mental composite score.

Methods

Participants
Participants were 396 community-dwelling middle-aged and elderly adults (196 men). Mean age of the sample was 69.04 years (SD = 8.41; 69.08 ± 8.21 and 69.01 ± 8.62, respectively, for men and women; t(394) = 0.08; p = 0.94). Just under 65% (64.8%) of the sample was 65 years or older. Sociodemographic characteristics of the sample are listed in Table 1. No clinical diagnoses were provided. However, according to the previously reported cut-off scores of the STICSA (Van Dam et al., 2013), 30.8% (n = 122) of the participants met criteria for clinical anxiety (STICSA Trait scale total score > 43), while 5.3% (n = 21) of the participants met criteria for the possible presence of anxiety disorders (STICSA Trait scale total score > 40).

The sample was recruited through advertisements (flyers, newspapers, and online ads) posted for established community groups (senior centers, retirement communities, and church groups, which focused on older adults). Study participants contributed voluntarily and anonymously; no honorarium was given for completing the assessments. All participants provided written, informed consent. The study was approved by the local institutional review board.

Measures
All participants were administered Italian versions of the STICSA, the Teate Depression Inventory (TDI; Balsamo and Saggino, 2013; 2014; Balsamo et al., 2014), the Geriatric Depression Scale (GDS; Yesavage et al., 1982), and the short-form-12 (SF-12) Health Survey (Ware et al., 1996). All the respondents completed paper-and-pencil versions of the questionnaires in a fixed order (a sociodemographic checklist, the GDS, the STICSA, the SF-12, and the TDI) on site at within established community groups. The protocol was administered by licensed psychologists who had a four-hour training wherein the objectives of the research, characteristics of the instruments administered, and information about common issues in psychological assessment among older adults (e.g. the effects of visual impairment or cognitive deficits on psychological testing) was explained. Since participants anonymously completed the assessments, those who scored highly on the depression and anxiety scales did not receive advice regarding psychological treatment. The STICSA is a 21-item measure designed to assess cognitive (e.g. “I feel agonized over my problems,” “I think that others won’t approve of me”) and somatic (e.g. “My heart beats fast,” “My muscles are tense”) symptoms. It can be administered in both trait and state variations. In the trait-anxiety subscale, the individual rates how often a statement is true in general, whereas in the state-anxiety subscale she/he rates how she/he feels at the moment of assessment. In total, the overall...
scale is made up of four subscales: State–Somatic (SS), Trait–Somatic (TS), State–Cognitive (SC), and Trait–Cognitive (TC). Each statement is rated on a four-point Likert-type scale from 1 (not at all) to 4 (very much so). Items of the STICSA were translated into Italian according to standard procedures of forward- and back-translation (van de Vijver and Tanzer, 1997). The translation was performed first from English into Italian and then from Italian into English by two independent bilingual professionals. The translators were two independent bilingual psychologists with previous experience in the cross-cultural validation of mental health measures. The back-translated version of the STICSA was submitted and approved by the author of the original scale (M. Ree, personal communication, February 04, 2013), who did not find any discrepancy in comparison to the original English version.

The GDS is a 30-item self-report questionnaire designed to measure depression in a geriatric population. Items represent symptoms of depression commonly seen in the elderly across affective (e.g. sadness, apathy, crying) and cognitive domains (e.g. thoughts of hopelessness, helplessness, guilt, worthlessness). The GDS does not assess somatic symptoms of depression (e.g. disturbances in energy level, appetite, sleep, sexual interest), which may confer less discriminability between depression and health conditions (Smarr and Keefer, 2011). Respondents rate each item on a dichotomous (yes/no) scale and the total score ranges between 0 and 30, with higher scores indicating more severe depression. In the present sample, Cronbach’s $\alpha$ was 0.84.

The TDI is a new 21-item self-report instrument designed to assess major depressive disorder as specified by the latest editions of the DSM (DSM-IV-TR and DSM-V; American Psychiatric Association, 2000; 2013) (Balsamo and Saggino, 2013). It was developed via Rasch logistic analysis of responses, within the framework of item response theory (Rasch, 1980; Andrich, 1995), in order to overcome inherent psychometric weaknesses of existing measures of depression, including the BDI-II (Balsamo and Saggino, 2007). Each item is rated on a five-point Likert-type scale, ranging from 0 (always) to 4 (never). A small but growing literature suggests that the TDI has strong psychometric properties in both clinical and nonclinical samples, including an excellent Person Separation Index, no evidence of bias due to item-trait interaction, good discriminant and convergent validity, and control of major response sets (Balsamo and Saggino, 2013, 2014; Balsamo et al., 2013, 2014; Innamorati et al., 2013). In a recent study, three cut-off scores were recommended in terms of sensitivity, specificity, and classification accuracy for screening for varying levels (minimal, mild, moderate, and severe) of depression severity in a group of patients diagnosed with major depressive disorder (Balsamo and Saggino, 2014). In the present sample, Cronbach’s $\alpha$ was 0.88.

The SF-12 is a 12-item SF of the SF-36 (Ware and Sherbourne, 1992). The SF-12 was developed for the Medical Outcomes Study (Stewart and Ware, 1992), a multi-year study of patients with chronic conditions. The instrument was designed to reduce respondent burden while achieving minimum standards of precision for purposes of group comparisons involving multiple health dimensions. The SF-12 assesses health-related quality of life in the last four weeks and covers the same health domains as the original SF-36 with one or two questions per domain, and provides two composite scores of physical (PCS) and mental health (MCS). Internal consistency coefficients were 0.80 and 0.77 for PCS and MCS, respectively.

**Statistical analysis**

We conducted confirmatory factor analysis (CFA) using the statistical package LISREL 8.7 (Jöreskog and Sörbom, 2006). All analyses were conducted using asymptotic covariance matrices and robust maximum-likelihood estimation methods because the distributions of some model variables deviated from normality.

We tested four possible factor models underlying items of the STICSA: (1) one-factor model in which all (both state and trait, as well as both cognitive and somatic) items were forced to load on a single higher order factor (Model 1); (2) two-factor model in which items across the state and trait scales loaded on either cognitive or somatic factors (Model 2); (3) two-factor model in which items loaded on either state or trait factors (Model 3); and (4) four-factor model in which State-Cognitive (STICSA-SC), State-Somatic (STICSA-SS), Trait-Cognitive (STICSA-TC), and Trait-Somatic factors (STICSA-TS) were directly modeled (Model 4).

The fit of each model was assessed by means of the following goodness-of-fit indices: (1) the Satorra–Bentler chi-squared (SB $\chi^2$) statistic and its degree of freedom; (2) the root mean square error of approximation (RMSEA) and its 90% confidence interval (90% CI); (3) the Non-Normed Fit Index (NNFI); (4) the comparative fit index (CFI); and (5) the standardized root mean square residuals (SRMR). In the present study, we considered NNFI, and CFI values of 0.90 and above to reflect adequate fit and values of 0.95 and above to indicate excellent fit. RMSEA and SRMR values of 0.08 or less were considered to reflect adequate fit; values
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Table 2. Fit indices for the structural models (N = 396)

<table>
<thead>
<tr>
<th>MODEL</th>
<th>SB ( \chi^2 )</th>
<th>df</th>
<th>NNFI</th>
<th>CFI</th>
<th>SRMR</th>
<th>RMSEA</th>
<th>90% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>One factor (Model 1)</td>
<td>3,992.55*</td>
<td>819</td>
<td>0.92</td>
<td>0.92</td>
<td>0.085</td>
<td>0.099</td>
<td>0.096/0.100</td>
</tr>
<tr>
<td>Two factor, cognitive-somatic (Model 2)</td>
<td>2,879.00*</td>
<td>818</td>
<td>0.95</td>
<td>0.95</td>
<td>0.073</td>
<td>0.080</td>
<td>0.077/0.083</td>
</tr>
<tr>
<td>Two factor, state-trait (Model 3)</td>
<td>3,929.59*</td>
<td>818</td>
<td>0.92</td>
<td>0.92</td>
<td>0.084</td>
<td>0.098</td>
<td>0.095/0.10</td>
</tr>
<tr>
<td>Four-factor (Model 4)</td>
<td>2,713.13*</td>
<td>813</td>
<td>0.95</td>
<td>0.95</td>
<td>0.070</td>
<td>0.077</td>
<td>0.074/0.080</td>
</tr>
</tbody>
</table>

*p < 0.001.

Note. SB \( \chi^2 \) = satorra and bentler chi-squared test; df = degrees of freedom, NNFI = Non-Normed fit index; CFI = comparative fit index; SRMR = standardized root mean square residual; RMSEA = root-mean-square error of approximation; 90% CI = 90% confidence interval of RMSEA.

...of 0.06 or less were considered to reflect good fit (Browne and Cudek, 1993; Hu and Bentler, 1999).

To investigate the psychometric properties of the STICSA, we assessed internal consistency of its scales using Cronbach’s \( \alpha \) indices. The concurrent and discriminant validity of the STICSA and its factor subscales were examined through Pearson’s correlation coefficients. We compared the pairs of correlation coefficients in the analysis of discriminant validity using the approach recommended by Meng et al. (1992). A multivariate general linear model (GLM) was used to assess the effect of age (middle-aged adults vs. older adults) and sex (males vs. females) on STICSA scores. Wilks’ lambda (\( \lambda \)) and the partial eta squared (\( \eta^2 \)) are reported as measures of multivariate effect and effect size, respectively.

Results

Confirmatory factor analysis

Goodness-of-fit statistics for all tested structural models are presented in Table 2. The SB \( \chi^2 \) goodness-of-fit tests were significant for each of the CFA models (SB \( \chi^2 \) ranged from 3,992.55, df = 819, to 2,713.13, df = 813, \( p < 0.001 \)). When we investigated the nested models with scaled difference in \( \chi^2 \) tests (Satorra and Bentler, 1994; Crawford and Henry, 2003; Brown, 2006), Model 4 demonstrated significantly better fit compared to Model 1 \([\Delta \chi^2 (6, N = 396) = 935.95, p < 0.001]\), and both of the two-factor models \([\Delta \chi^2 (5, N = 396) = 130.28, \text{ and Model 3, } \Delta \chi^2 (5, N = 396) = 823.03 (p < 0.001)\]).

Since the utility of the \( \chi^2 \) goodness-of-fit test has been questioned, probably due to its sensitivity to large sample sizes (Hu and Bentler, 1998; Kahn, 2006; Kline, 2011), we calculated additional fit indices to support the fit of the models. According to other goodness-of-fit indices, Model 1 and Model 3 generally provided inadequate fit to the observed response data. In contrast, Model 4 yielded adequate-to-excellent fit across all fit indices. A similar pattern was found for Model 2, which demonstrated adequate fit to the data. Taken together, these results support both the state-trait and cognitive-somatic distinctions implied by the STICSA item pool and instruction sets. Further, the four-factor model seemed to best fit the data, showing the lowest RMSEA value (RMSEA = 0.077; 90% CI: 0.074–0.080), closest to the criterion value of best fit of 0.05 (Schermelleh-Engel et al., 2003).

Reliability and validity analysis

All subscales of the STICSA had high internal consistency, with Cronbach’s \( \alpha \) coefficients of 0.87 and 0.86, respectively for the STICSA-TS and the STICSA-TC, and 0.90 and 0.86, respectively for the STICSA-SS and the STICSA-SC. Cronbach’s \( \alpha \) coefficients were equal to 0.92 and 0.91, respectively for the STICSA-S and the STICSA-T. In addition, the average inter-item correlations ranged between 0.38 (for the STICSA-TC and the STICSA-SS) and 0.45 (for the STICSA-SC), with STICSA-SC exhibiting an average inter-item correlation of 0.39.

Concurrent and discriminant validity of the STICSA

To investigate the concurrent and discriminant validity of the STICSA, we computed correlations among the STICSA dimensions and with measures of depression and health-related quality of life (see Table 3).

Relationships between cognitive and somatic anxiety, and depression

Compared to the somatic scale, the STICSA cognitive scale displayed a significantly higher correlation with the GDS both for the trait scale \((Z = 3.77, p < 0.01)\) and for the state scale \((Z = 5.41, p < 0.01)\). No significant differences were detected in the magnitude of the correlations among the STICSA cognitive scale with the TDI compared to the somatic scale, for state \((Z = -1.42, p = 0.08)\) or trait anxiety \((Z = -1.44, p = 0.08)\).
Table 3. Correlations among the STICSA, measures of depression, and the SF-12 composite scores (N = 396)

<table>
<thead>
<tr>
<th>MEASURE</th>
<th>1</th>
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<th>3</th>
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<th>7</th>
<th>8</th>
<th>9</th>
<th>10</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. STICSA-Trait</td>
<td>0.92**</td>
<td>0.91**</td>
<td>0.84**</td>
<td>0.73**</td>
<td>0.80**</td>
<td>0.51**</td>
<td>0.63**</td>
<td>−0.29**</td>
<td>−0.48**</td>
<td></td>
</tr>
<tr>
<td>2. STICSA-Trait, Somatic</td>
<td>0.67**</td>
<td>0.80**</td>
<td>−0.65**</td>
<td>0.44**</td>
<td>0.52**</td>
<td>−0.31**</td>
<td>−0.43**</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>3. STICSA-Trait, Cognitive</td>
<td>0.73**</td>
<td>0.52**</td>
<td>0.82**</td>
<td>0.49**</td>
<td>0.64**</td>
<td>−0.22**</td>
<td>−0.45**</td>
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<tr>
<td>4. STICSA-State</td>
<td>0.91**</td>
<td>0.91**</td>
<td>0.49**</td>
<td>0.57**</td>
<td>−0.32**</td>
<td>−0.44**</td>
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<tr>
<td>5. STICSA-State, Somatic</td>
<td>0.68**</td>
<td>0.43**</td>
<td>0.43**</td>
<td>−0.32**</td>
<td>−0.38**</td>
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<tr>
<td>6. STICSA-State, Cognitive</td>
<td>0.48**</td>
<td>0.61**</td>
<td>−0.26**</td>
<td>−0.44**</td>
<td></td>
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<tr>
<td>7. TDI</td>
<td>0.56**</td>
<td>−0.39**</td>
<td>−0.54**</td>
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<tr>
<td>8. GDS</td>
<td>−0.34**</td>
<td>−0.51**</td>
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<tr>
<td>9. PCS</td>
<td>0.13*</td>
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<td>10. MCS</td>
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</table>

* p < 0.01; ** p < 0.05.

Note. STICSA = state-trait inventory for cognitive and somatic anxiety; TDI = teate depression inventory; GDS = geriatric depression scale; PCS = SF-12 physical composite score; MCS = SF-12 mental composite score.

Relationships between cognitive and somatic anxiety, and physical well-being

The STICSA somatic scale displayed a significantly higher correlation with the SF-12 PCS compared with the cognitive scale for trait anxiety (Z = 3.77; p < 0.01), but not for state anxiety (Z = 1.57; p = 0.06).

Relationship of cognitive and somatic anxiety relative to relationship among

Depression measures
All anxiety dimension correlations with depression were weaker than the correlation between measures of depression (r = 0.56), with the sole exception of the cognitive dimension of the STICSA and the GDS.

Relationship of anxiety dimensions with well-being
The correlations among the STICSA subscales and the SF-12 composite scores were all negative, indicating that more severe anxiety was associated with lower quality of life. Comparison of the STICSA with the mental and physical health subscales of the SF-12 revealed that STICSA dimensions/subscales were significantly more correlated with the SF-12 MCS than with the SF-12 PCS (Z between −2.02 [p < 0.05] and −2.99 [p < 0.001]), except for the STICSA-SS (Z = −0.99; p = 0.16), suggesting the STICSA measures anxiety independently from physical health (Table 3).

 Differences between sex and age classes

Descriptive statistics for the STICSA by sex (males vs. females) and age group (middle-aged vs. older adults) are listed in Table 4. Multivariate analyses revealed a significant effect of age (Wilks’ λ = 0.97, F(5, 384) = 3.76, p < 0.05, partial η² = 0.05) and sex (Wilks’ λ = 0.93, F(5, 384) = 5.74, p < 0.01, partial η² = 0.07), but no interaction (Wilks’ λ = 0.98, F(5, 384) = 1.93, p = 0.09, partial η² = 0.02). Men and women differed on all the dimensions of the STICSA (p < 0.001), with women reporting higher scores (partial η² ranging from 0.03 to 0.05). Age groups differed only on the STICSA-SS, with older adults reporting higher scores than middle-aged adults (partial η² = 0.02).

Discussion
Some anxiety measures have acceptable psychometric evidence for measuring anxiety in older adults, although the majority may not necessarily differentiate from depression sufficiently well (Dennis et al., 2007; Therrien and Hunsley, 2012). Given the recent establishment of the STICSA as a promising instrument in the assessment of anxiety for its ability to differentiate anxiety from depression, our general aim was to investigate whether it showed psychometric properties sufficient to justify its use in older adults.

In our sample of middle-aged and older adults, the original four-factor structure yielded adequate-to-excellent fit according to established standards in the field (Hu and Bentler, 1999). These findings provide support for the distinctions between state and trait forms of anxiety and between cognitive and somatic anxiety, extending previous findings in younger samples (Gros et al., 2007; Ree et al., 2008; Gros et al., 2010; Van Dam et al., 2013). All dimensions of the STICSA have been found to have excellent internal consistency as reported in previous psychometric analyses carried out in younger samples (Gros et al., 2007; Ree et al., 2008). As regards the investigation on the concurrent and discriminant
STICSA and the TDI. Previous work has suggested that the cognitive and somatic subscales of the STICSA would have to be higher than either subscales’ correlation with two depression measures, the GDS and the TDI. Previous work has suggested that the STICSA shows promise discriminating anxiety from depression (Gros et al., 2007). In the present study, the correlation among depression measures was higher than correlations between anxiety and depression with the exception of cognitive anxiety (trait and state) and overall trait anxiety with the GDS. Not surprisingly, cognitive anxiety may be associated with measures of depression more strongly than somatic anxiety. While items derived from cognitive dimensions of anxiety (e.g. “I think the worst will happen,” “I feel agonized over my problems”) likely overlap with aspects also common in depressive states, items derived from somatic dimensions of anxiety (e.g. “My heart beats fast,” “My muscles are tense”) likely tap the presence of physiological hyperarousal, which is more uniquely characteristic of anxiety. Despite some overlap with depressive symptoms, cognitive anxiety is an important component of many anxiety disorders (Van Dam et al., 2013). Correlations between cognitive anxiety and the GDS were above 0.60 (0.61–0.64), and they were higher than the correlation between the two measures of depression (0.56). However, these effect sizes were comparable to those reported by Gros et al. (2007), who obtained correlations of 0.59–0.64 between cognitive anxiety and depression, as measured with the Depression Anxiety Stress scales (DASS; Lovibond and Lovibond, 1995). While these figures denote a partially limited ability to discriminate cognitive anxiety symptoms from depression, Gros et al. (2007) reported correlations even higher between the STAI and depression (ranging between 0.64 and 0.68).

Also, the elevated correlation between the GDS and the cognitive scales of the STICSA could be due to the fact that the GDS only measures cognitive and affective aspects of depression (not somatic/physical).

According to our second hypothesis, the STICSA would have to be more strongly correlated with mental health functioning than physical health functioning, as measured with the SF-12. We found trait somatic anxiety was more correlated with mental health functioning than with physical health functioning, as assessed by the SF-12. This finding supports the hypothesis that the STICSA might assess anxiety at least partially independently from the presence of physical health limitations, despite a heavy reliance on somatic items. Consistent with two depression measures, the GDS and the TDI. Previous work has suggested that the cognitive and somatic subscales of the STICSA would have to be higher than either subscales’ correlation with two depression measures, the GDS and the TDI. Previous work has suggested that the STICSA shows promise discriminating anxiety from depression (Gros et al., 2007). In the present study, the correlation among depression measures was higher than correlations between anxiety and depression with the exception of cognitive anxiety (trait and state) and overall trait anxiety with the GDS. Not surprisingly, cognitive anxiety may be associated with measures of depression more strongly than somatic anxiety. While items derived from cognitive dimensions of anxiety (e.g. “I think the worst will happen,” “I feel agonized over my problems”) likely overlap with aspects also common in depressive states, items derived from somatic dimensions of anxiety (e.g. “My heart beats fast,” “My muscles are tense”) likely tap the presence of physiological hyperarousal, which is more uniquely characteristic of anxiety. Despite some overlap with depressive symptoms, cognitive anxiety is an important component of many anxiety disorders (Van Dam et al., 2013). Correlations between cognitive anxiety and the GDS were above 0.60 (0.61–0.64), and they were higher than the correlation between the two measures of depression (0.56). However, these effect sizes were comparable to those reported by Gros et al. (2007), who obtained correlations of 0.59–0.64 between cognitive anxiety and depression, as measured with the Depression Anxiety Stress scales (DASS; Lovibond and Lovibond, 1995). While these figures denote a partially limited ability to discriminate cognitive anxiety symptoms from depression, Gros et al. (2007) reported correlations even higher between the STAI and depression (ranging between 0.64 and 0.68).

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### Table 4. Descriptive statistics broken down by sex (men vs. women) and age class (middle-aged vs. older adults)

<table>
<thead>
<tr>
<th></th>
<th>MALES VERSUS FEMALES M±SD</th>
<th>F(1)</th>
<th>P</th>
<th>PARTIAL η²</th>
<th>MIDDLE-AGED VERSUS OLDER ADULTS M±SD</th>
<th>F(1)</th>
<th>P</th>
<th>PARTIAL η²</th>
<th>WHOLE SAMPLE M±SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>STICSA-trait</td>
<td>34.23±10.23</td>
<td>25.86</td>
<td>&lt;0.001</td>
<td>0.06</td>
<td>37.58±11.03</td>
<td>0.21</td>
<td>0.65</td>
<td></td>
<td>37.93±12.13</td>
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<tr>
<td></td>
<td>versus 41.55±12.76</td>
<td></td>
<td></td>
<td></td>
<td>38.18±12.73</td>
<td></td>
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<tr>
<td>STICSA-trait, cognitive</td>
<td>17.29±5.41</td>
<td>21.92</td>
<td>&lt;0.001</td>
<td>0.05</td>
<td>18.75±6.12</td>
<td>1.03</td>
<td>0.31</td>
<td></td>
<td>19.21±6.84</td>
</tr>
<tr>
<td></td>
<td>versus 21.08±7.54</td>
<td></td>
<td></td>
<td></td>
<td>19.46±7.20</td>
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<td></td>
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<tr>
<td>STICSA-state</td>
<td>17.00±5.77</td>
<td>20.57</td>
<td>&lt;0.001</td>
<td>0.05</td>
<td>18.86±6.23</td>
<td>0.05</td>
<td>0.83</td>
<td></td>
<td>18.77±6.45</td>
</tr>
<tr>
<td></td>
<td>versus 20.49±6.63</td>
<td></td>
<td></td>
<td></td>
<td>18.72±6.59</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>STICSA-state,</td>
<td>30.81±9.28</td>
<td>18.09</td>
<td>&lt;0.001</td>
<td>0.05</td>
<td>32.14±10.46</td>
<td>3.36</td>
<td>0.07</td>
<td></td>
<td>33.77±11.81</td>
</tr>
<tr>
<td></td>
<td>versus 36.68±13.24</td>
<td></td>
<td></td>
<td></td>
<td>34.74±12.43</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>STICSA-state,</td>
<td>15.04±5.06</td>
<td>17.19</td>
<td>&lt;0.001</td>
<td>0.04</td>
<td>15.49±5.51</td>
<td>7.17</td>
<td>0.01</td>
<td>0.02</td>
<td>16.65±6.64</td>
</tr>
<tr>
<td></td>
<td>versus 18.22±7.57</td>
<td></td>
<td></td>
<td></td>
<td>17.29±7.11</td>
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</tr>
<tr>
<td>STICSA-state,</td>
<td>15.68±5.35</td>
<td>11.94</td>
<td>0.001</td>
<td>0.03</td>
<td>16.55±5.71</td>
<td>1.33</td>
<td>0.25</td>
<td></td>
<td>17.03±6.24</td>
</tr>
<tr>
<td></td>
<td>versus 18.34±6.74</td>
<td></td>
<td></td>
<td></td>
<td>17.30±6.50</td>
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</table>

**Multivariate statistics:** Sex: Wilks’ λ = 0.93; F(5;384) = 5.74; p < 0.01; partial η² = 0.07. Age: Wilks’ λ = 0.95; F(5;384) = 3.76; p < 0.05; partial η² = 0.05. Sex * Age: Wilks’ λ = 0.98; F(5;384) = 1.93; p = 0.09; partial η² = 0.02.
with this pattern of correlations, the multivariate GLM indicated a partial independence of the STICSA from age: although older adults are affected more frequently by medical conditions, multivariate analyses indicated a significant, but weak effect of age, with middle-aged adults and older adults showing meaningful differences only for state somatic anxiety. These results are important because the detection of anxiety in older adults is generally complicated by the high frequency of medical disorders present in this age group. Several studies suggest that between 80% and 86% of adults aged 65 years and older have at least one significant medical condition (Dawson et al., 1986; Naughton et al., 2007). Since somatic anxiety may be misidentified as a physical symptom and/or may be resultant from a medical condition, appropriate identification of somatic anxiety in the elderly is critical (Therrien and Hunsley, 2012).

As anxiety appears to have meaningful differences in demographics, between-group analyses for sex were conducted to explore the extent of these differences. Sex had a significant impact on all dimensions of the STICSA in the present sample, with effect sizes greater for trait anxiety than for state anxiety. Compared with males, females reported higher scores on all dimensions of the STICSA, consistent with increased risk among women for an anxiety disorder in older age (Wolitzky-Taylor et al., 2010), as well as with higher scores on the STAI trait subscale among older women relative to male peers (Bergua et al., 2012). Higher STICSA scores for females, compared with males, have also been reported in younger samples (Van Dam et al., 2013).

Limitations and strengths

Several limitations of this study should be addressed in future studies. First, we did not screen the sample for the presence of anxiety disorders thorough diagnostic interviews, and did not assess the ability of the STICSA to discriminate between individuals with anxiety disorders and healthy controls. Second, we did not assess the validity of the STICSA relative to the STAI or other existing measures of anxiety (especially those specifically devised to be used with older adults) (e.g. the Geriatric Mental State Examination; Copeland et al., 1976). Third, we did not assess whether scores on the STICSA were affected by social desirability or the presence of memory impairment and cognitive deficits.

Despite these limitations, our study has several strengths: it is the first study investigating the psychometric properties of the STICSA in a sample of older adults and the first study evaluating the Italian version of this measure, to our knowledge. Furthermore, we also administered to our sample the SF-12, which shed light on the ability of the STICSA in detecting anxiety symptoms while simultaneously assessing for the presence of physical health symptoms.

Conclusion

In conclusion, the STICSA is a promising measure of general anxiety, which provides a more specific assessment of cognitive and somatic anxiety symptoms among older adults. Together, the present findings support its reliability and validity as a measure of state and trait anxiety in a geriatric population. Nevertheless, our data suggest that the STICSA may have some limitations in discriminating between anxiety and depression. Further studies to investigate its properties relative to other geriatric anxiety measures are necessary. The STICSA may be particularly attractive for researchers and clinicians working with older adults because of its relatively weak association with physical health.

Conflict of interest

None.

Description of authors’ roles

M. Balsamo and M. Innamorati contributed equally to the design of the study and supervision of data collection. M. Balsamo, M. Innamorati, and N.T. Van Dam all contributed to drafting of the manuscript. M. Balsamo, M. Innamorati, and L. Carlucci performed statistical analyses. All authors have contributed to and have approved the final manuscript.

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