Anxiety and depressive disorders are associated with delusional-like experiences: a replication study based on a National Survey of Mental Health and Wellbeing

Sukanta Saha,1 James Scott,1,2,3,4 Daniel Varghese,5 John McGrath1,4,6

ABSTRACT

Objectives: There is growing evidence that delusional-like experiences (DLE) are associated with common mental disorders. In particular, a National Mental Health Survey conducted in Australia during 2007 reported an association between DLE and both anxiety disorder and major depressive disorder (MDD). However, the previous study did not examine this association with respect to subtypes of anxiety disorder nor with severity of MDD. The aim of this study was to examine the associations between DLE and both anxiety disorder and MDD in more detail based on an independent population sample.

Design: Cross-sectional study.

Setting: Subjects were drawn from the Australian Survey of Mental Health and Wellbeing 1997 using a stratified multistage area sampling of persons living in private dwellings in all States and Territories of Australia.

Participants: Approximately 13,600 private dwellings were initially selected with one person aged 18 years or older from each dwelling invited to participate. In total, 10,641 individuals participated in the survey.

Primary and secondary outcome measures: The Composite International Diagnostic Interview was used to identify individuals with DLE and Diagnostic and Statistical Manual of Mental Disorder, Fourth Edition (DSM IV) lifetime diagnoses of anxiety disorders and MDD. The influence of various anxiety disorders and MDD on DLE was assessed with logistic regression.

Results: Having a lifetime diagnosis of either any anxiety disorder or MDD was significantly associated with the endorsement of DLE. The association was found for each of the main anxiety disorders when examined separately. There was a dose—response relationship between increasing severity of MDD and higher odds of DLE endorsement.

Conclusions: DLE are associated with a wide range of anxiety disorders and are more prevalent in those with MDD. Understanding the relationship between DLE, anxiety disorders and depression may provide insights into shared pathways that underpin both psychotic disorders and common mental disorders.

ARTICLE SUMMARY

Article focus

- The study was undertaken in order (1) to examine the association between DLE and (a) broadly defined anxiety disorders and (b) MDDs; (2) to explore the association between DLE and a range of specific anxiety disorders and (3) to examine if severity of MDD influenced the risk of endorsement of DLE.

Key message

- Having a lifetime diagnosis of either any anxiety disorder or MDDs was significantly associated with the endorsement of DLE.
- The association was found for each of the main anxiety disorders when examined separately.
- There was a dose—response relationship between increasing severity of MDD and higher odds of DLE endorsement.

Strengths and limitations of this study

- The data were drawn from the nationally representative sample from the Australia general population.
- Cross-sectional study.

INTRODUCTION

There is now robust evidence indicating that hallucinations and delusional-like experiences (DLE) are common in the general population. In recent years, the field has focused on the demographic and clinical correlates of hallucinations and DLE.1–10 Of particular interest, there is a growing body of evidence reporting an association between DLE endorsement and common mental disorders, such as anxiety disorders and major depressive disorder (MDD).
example, panic attacks during adolescence were significantly associated with increased levels of DLE among young adults. In the NEMESIS study, subjects with obsessive–compulsive symptoms were more likely to develop incident psychotic symptoms 3 years later. Conversely, a Swiss-based cohort reported that young adults with psychotic-like experiences were significantly more likely to later develop common mental disorders, such as anxiety disorders and MDD. A German community-based study found an association between social phobia, social anxiety and DLE, while a US primary-care-based sample reported that those who reported psychotic-like experiences were more likely to have generalised anxiety disorders and panic disorders.

Trauma exposure with or without post-traumatic stress disorder has been associated with DLE. Several Australian studies have found significant associations between DLE and broadly defined anxiety disorders; however, to date, these studies did not report on subtypes of anxiety disorders. In light of the evidence linking DLE with a wide range of different types of anxiety disorders, the evidence suggests that DLE are non-specifically associated with anxiety disorders.

With respect to depression, several studies have found that individuals with depression are significantly more likely to endorse DLE. Studies also show that DLE requiring clinical care were progressively more likely to occur with greater levels of affective dysregulation (depressive symptoms and hypomanic symptoms). Importantly, there was a significant association between severity of depressive symptoms and persistence of psychotic symptoms.

While longitudinal studies are required to explore the temporal sequence between depression, anxiety and DLE, we had the opportunity to replicate our previous findings with respect to the cross-sectional association between DLE and (1) broadly defined anxiety disorders and (2) MDD. Based on our previous studies, we predicted that those with anxiety disorder or major depression disorder would be more likely to endorse DLE. In addition, we were able to explore the association between DLE and a range of specific anxiety disorders. Furthermore, we were able to examine if severity of MDD influenced the risk of endorsement of DLE—we predicted that those with more severe MDD would be more likely to endorse DLE compared with those with milder forms of MDD.

METHODS

Participants

The data were drawn from the 1997 National Survey of Mental Health and Wellbeing conducted in Australia by the Australian Bureau of Statistics from a representative sample (random stratified multistage area sampling) of persons living in private dwellings in all States and Territories of Australia. Details of the survey methodology were published elsewhere. In brief, approximately 13,600 private dwellings were initially selected with one person aged 18 years or older from each dwelling invited to participate. In total, 10,641 individuals participated in the survey, representing a response rate of 78%. Interviews were carried out by trained interviewers from the Australian Bureau of Statistics, a statutory body responsible for conducting such surveys using ethical protocols that include written informed consent.

Assessment of DLE and DSM-IV diagnoses

Mental disorders were assessed by a modified version of the Composite International Diagnostic Interview (CIDI), which yielded diagnoses of Diagnostic and Statistical Manual of Mental Disorder, Fourth Edition (DSM IV) disorders. Briefly, within the CIDI, there are three items related to identifying individuals who may be psychotic (‘items: ‘screening items’). For those who endorsed the screen item, a follow-up item was used to further explore the delusional-like nature of the experiences (‘probe items’). Full details of the screen and probe items are provided in online appendix 1. The items covered the following features of psychotic disorders: delusions of control, thought interference and passivity (question 1 and 1a); delusions of reference or persecution (question 2 and 2a) and grandiose delusions (question 3 and 3a). There was no item to assess hallucinations.

Based on CIDI-derived DSM-IV criteria, we identified subjects who had lifetime diagnoses of (1) an anxiety disorder and (2) MDD. Anxiety disorders included panic disorder with or without agoraphobia, social phobia, generalised anxiety disorder, obsessive–compulsive disorder and agoraphobia without panic disorder. For those with MDD, allocation to subtypes was based on the total number of particular ‘depressive’ symptoms with the duration of at least 2 weeks. Full details of the symptom list and related rules to deal with multiple episodes can be found in the full report. In brief, mild MDD was characterised by the presence of at least four symptoms, moderate MDD with at least six symptoms and severe MDD with at least eight symptoms. These subtypes of MDD were mutually exclusive.

To ascertain trauma exposure, the CIDI elicits responses from 10 questions pertaining to past exposure to traumatic events. Details of the trauma variables have been published previously by our group. In keeping with our previous analyses, individuals who screened positively for schizophrenia (ie, respondents who reported ‘yes’ to the item ‘Had been told at any time by a psychiatrist that they had schizophrenia’) were excluded from the analyses (n=87), leaving a total of 10,554 subjects for this study.

Statistical analysis

To examine the association between DLE and both anxiety disorders and MDD, logistic models were fitted to the data while adjusting for various confounding factors. Because sex and age are associated with DLE, we included these as covariates in the main analyses. In keeping with our previous studies, we included a range of CIDI-derived potential confounding
Table 1  Descriptive statistics of delusional-like experiences (screen items), anxiety disorder and major depressive disorder (n=10 554)

<table>
<thead>
<tr>
<th>Exposure</th>
<th>Sample N (%)</th>
<th>Delusional-like experiences endorsement, n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>No (88.44)</td>
</tr>
<tr>
<td>Total sample</td>
<td>10 554 (100.00)</td>
<td>9278</td>
</tr>
<tr>
<td>Anxiety and depressive disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No anxiety disorders</td>
<td>9974 (95.13)</td>
<td>8900 (85.16)</td>
</tr>
<tr>
<td>Any anxiety disorders: lifetime*</td>
<td>580 (4.87)</td>
<td>378 (4.29)</td>
</tr>
<tr>
<td>No major depressive disorder</td>
<td>9903 (94.66)</td>
<td>8834 (84.76)</td>
</tr>
<tr>
<td>Any major depressive disorder: lifetime†</td>
<td>651 (5.34)</td>
<td>444 (4.77)</td>
</tr>
</tbody>
</table>

*Anxiety disorders based on Composite International Diagnostic Interview (CIDI) DSM IV diagnosis.
†Major depressive disorder based on CIDI DSM IV diagnosis.

Table 2  Association between delusional-like experiences, and anxiety disorders and major depressive disorder (n=10 554)

<table>
<thead>
<tr>
<th>Disorders</th>
<th>Delusional-like experiences</th>
<th>Probe items</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Screen items</td>
<td>Probe items</td>
</tr>
<tr>
<td>Model 1§</td>
<td>Model 2†</td>
<td>Model 1§</td>
</tr>
<tr>
<td>Anxiety disorders: lifetime‡</td>
<td>3.88 (2.92 to 5.16)§</td>
<td>2.43 (1.91 to 3.09)§</td>
</tr>
<tr>
<td>Major depressive disorder: lifetime†</td>
<td>3.63 (2.75 to 4.79)§</td>
<td>2.17 (1.65 to 2.86)§</td>
</tr>
</tbody>
</table>

§Significance: p<0.001.
†Model 2 was adjusted for age, sex, marital status, migrant status, income, employment status, educational status, any alcohol use/dependence disorders, any drug use/dependence disorders and any traumatic life events (in model 2, anxiety disorders were adjusted for major depressive disorder and vice versa).
‡Anxiety and depressive disorders were based on Composite International Diagnostic Interview DSM IV diagnosis.

The sample was weighted to adjust for differential probabilities of selection within households, oversampling of population subgroups and non-response to match census population distribution on a number of geographic and socio-demographic variables. The initial weights were calibrated against known population estimates. Replicate weight variables were developed using the Jack-knife procedure of replication (ie, the analysis was repeated after one subject was dropped and then the SE was derived from the distribution of results from all ‘minus one’ resamples). Analyses were performed using Proc Surveylogistic, which is designed to analyse complex survey sample using SAS (V.9.3; SAS Institute). χ² Test-for-linear trend was used to assess dose–response relationships between the exposure variables and DLE.

RESULTS
Of the 10 554 subjects surveyed, 11.6% (n=1276) positively endorsed one or more DLE items (table 1). There was a weak effect of women being more likely to endorse DLE than men (OR 1.05, 95% CI 1.04 to 1.05). The prevalence of lifetime diagnosis of any anxiety disorder was 4.9% (n=580), and the prevalence of lifetime depressive disorders was 5.3% (n=651).

As predicted, the main analyses showed that those with any anxiety disorder and participants who had lifetime diagnosis of MDD were significantly more likely to endorse DLE. Those with anxiety disorders were two to three times more likely to endorse both DLE screen and probe items (table 2), and those with a diagnosis of MDD were also two to three times more likely to endorse DLE screen and probe items.

Concerning the subtypes of anxiety disorders, each disorder was significantly associated with DLE screen items, and there were no marked differences in the effect sizes between the different disorders (table 3). There was a dose–response relationship between the severity of the MDD and DLE in which severe depression showed twice the odds of endorsement of DLE screen items compared with a diagnosis of mild MDD with a significant linear trend (χ²=44.19, p<0.0001). Broadly similar (but less precise) associations were also found for probe items.
### Table 3: Association between delusional-like experiences, and different individual exposure to lifetime anxiety disorders and major depressive disorder (n=10,554)

<table>
<thead>
<tr>
<th>Anxiety disorders</th>
<th>Screen items</th>
<th>Probe items</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Number (%), SE</td>
<td>OR (95% CI)</td>
</tr>
<tr>
<td>Panicking disorder</td>
<td>124 (1.02, 0.12)</td>
<td>4.56 (2.51 to 8.33)</td>
</tr>
<tr>
<td>General anxiety</td>
<td>311 (2.57, 0.23)</td>
<td>3.69 (2.57 to 5.29)</td>
</tr>
<tr>
<td>Obsessive-compulsive disorder</td>
<td>77 (0.69, 0.12)</td>
<td>5.19 (2.69 to 10.03)</td>
</tr>
<tr>
<td>Agoraphobia without panic disorder</td>
<td>60 (0.49, 0.06)</td>
<td>5.18 (2.72 to 9.85)</td>
</tr>
<tr>
<td>Social phobia</td>
<td>160 (1.35, 0.14)</td>
<td>4.14 (2.81 to 6.11)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Major depressive disorder</th>
<th>Screen items</th>
<th>Probe items</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mild</td>
<td>297 (2.52, 0.20)</td>
<td>2.96 (1.82 to 4.82)</td>
</tr>
<tr>
<td>Moderate</td>
<td>190 (1.52, 0.14)</td>
<td>3.29 (1.81 to 6.01)</td>
</tr>
<tr>
<td>Severe</td>
<td>164 (1.29, 0.12)</td>
<td>5.73 (3.96 to 8.30)</td>
</tr>
</tbody>
</table>

**Trend**

χ²=111.83, p<0.0001

χ²=44.19, p<0.0001

χ²=21.19, p<0.0001

χ²=6.04, p<0.001

**Models**

†Model 1 = adjusted for age, sex, marital status, migrant status, income, employment status, educational status, any alcohol use/dependence disorders, any drug use/dependence disorders and any traumatic life events (in model 2, anxiety disorders were adjusted for major depressive disorder and vice versa).

‡Significance: p<0.001 (shown in bold).

### DISCUSSION

In the secondary analysis, when we conducted the modelling using two DLE items (C1 and C2), the pattern of significant association for major anxiety and depressive disorders remained unchanged (data not shown).

Individuals with a lifetime diagnosis of MDD or an anxiety disorder were significantly more likely to report DLE compared with those without these disorders. We found that each subtype of anxiety disorder was associated with DLE, and there were no marked differences in the effect sizes for these associations (the CIs around these findings, weighted with the exposure to DLE. Our new findings demonstrated that trauma exposure was associated with DLE. Our new findings and additional weight to the conclusion that a range of disorders with these associations for the DLE overlapped. Based on these findings, associations with major anxiety and depressive disorders remained unchanged (data not shown).

The mechanisms linking DLE with anxiety disorder and MDD remain unclear. However, there is evidence to suggest that shared familial factors may contribute to these findings. In the current study, we were not able to examine the temporal sequence between the variables of interest—for example, we do not know if anxiety or depressive symptoms preceded the onset of the DLE or vice versa. Additional research is required to explore these particular research questions. For instance, we do not know if anxiety or depressive symptoms preceded the onset of the DLE or vice versa. Additional research is required to explore these particular research questions.

As predicted, there was also a dose–response relationship between severity of MDD and DLE. All associations remained significant when adjusted for associated comorbidity with anxiety, alcohol and illicit substance misuse and any traumatic life events, indicating that the associations are independent of comorbid psychiatric illnesses and selected environmental and cognitive risk factors.
There is now robust and consistent evidence indicating that those with anxiety disorders and MDD have an increased risk of DLE. For example, clinicians involved in the care of those with primary diagnoses of anxiety disorder or depression may not routinely enquire about DLE. In light of the association between DLE and suicidal ideation/behaviour, the presence of these experiences may suggest that clinical care plans place greater emphasis on the detection and management of suicidal ideation. A recent study based on adolescents found that most individuals (57%–80% depending on age) who reported psychotic-like experiences (eg, hallucinations and/or DLE) had at least one diagnosable non-psychotic psychiatric disorder. We agree with these authors, who note that psychotic symptoms appear to be important risk markers for a wide range of non-psychotic mental health disorders.

**Contributors**

JM, SS and JS have directly participated in the planning and execution of the study. SS analysed the data. All authors have critically read and approved the final version submitted.

**Funding**

This research received no specific grant from any funding agency in public, commercial or not-for-profit sectors.

**Competing interests**

None.

**Patient consent**

Obtained.

**Ethics approval**

We obtained data from the Australian Bureau of Statistics (ABS), which is a government organisation. So, we are not aware about the exact approval authority. However, it is understood that the ABS has followed all the ethical standards to conduct this national survey.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

**Data sharing statement**

The data are available from the Australian Bureau of Statistics.

**REFERENCES**


