

Gender differences in major depressive disorder: findings from the Singapore Mental Health Study

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INTRODUCTION Major depressive disorder (MDD) is one of the most common psychiatric disorders worldwide and has been associated with various sociodemographic risk factors, including age, gender and ethnicity. The present study aimed to establish whether gender-specific differences relating to the prevalence and correlates of MDD exist in the Singapore adult resident population.

METHODS The Singapore Mental Health Study was a population-based, cross-sectional epidemiological study among Singapore citizens and permanent residents aged 18 years and above. Face-to-face interviews were completed with 6,616 respondents between December 2009 and December 2010. Psychiatric conditions were established using version 3.0 of the World Health Organization Composite International Diagnostic Interview (CIDI). In addition, data relating to chronic medical conditions was captured using a modified version of the CIDI checklist for chronic medical conditions.

RESULTS The lifetime prevalence of MDD was higher among women (7.2%) than men (4.3%). MDD was more prevalent among men and women who were divorced/separated and widowed women, as compared to those who were single. Among men, MDD was more prevalent among Indian and other ethnicities as compared to Chinese. Of the 417 respondents with MDD, women had significantly higher odds of having generalised anxiety disorder but lower odds of having high blood pressure, as compared to men.

CONCLUSION The study highlighted key gender-specific correlates of MDD. Given the comorbidities associated with MDD and other psychiatric disorders and/or physical illnesses, these correlates pose additional challenges for care providers.

Keywords: epidemiology, gender differences, mental disorders, prevalence

INTRODUCTION

Epidemiological studies worldwide have consistently reported major depressive disorder (MDD) to be among the most common psychiatric disorders, with an estimated lifetime prevalence in the range of 12% to 16% in Western communities,⁽¹⁻⁴⁾ and much lower in Asia, ranging between 3% and 6%.⁽⁵⁻⁷⁾ MDD can be chronic or recurrent, consequently affecting and impacting individuals for many months, years or even decades. MDD is also associated with significant comorbidity, poor health and mortality.

Certain sociodemographic risk factors, including age, gender and ethnicity, have frequently been associated with MDD. The prevalence of MDD is higher among women compared to men,⁽⁸⁻¹⁰⁾ and is often 1.5–3 times higher among women than men.⁽¹¹⁻¹³⁾ Research has also shown that among women, depression is the leading cause of disease-related disability.⁽¹¹⁾ These findings have been reported in both clinical and general populations and remain evident, irrespective of where the research is conducted and how it is assessed. These gender differences are likely to be a result of a myriad of factors, including biological, social, demographic and/or psychological effects. Gender itself affects many aspects of psychopathology, including prevalence of disorders, expression of symptoms, course of illness, help-seeking behaviour and response to treatment.⁽¹⁴⁾

Singapore is located off the Malaysian peninsula in Southeast Asia and has a resident population (including Singapore

citizens and permanent residents) of 3.8 million people.⁽¹⁵⁾ The Singapore Mental Health Study (SMHS) was a population-based epidemiological study that aimed to establish the prevalence of mental disorders among Singapore residents aged ≥ 18 years. Findings showed that MDD was the most prevalent mental disorder among those examined in the SMHS, which reported a lifetime prevalence of 5.8% and a 12-month prevalence of 2.2%.⁽⁶⁾

Upon further analysis, the SMHS also found that the prevalence of MDD was significantly higher among women, Indians and those who were divorced/separated or widowed. Chronic physical comorbidities were also found to be present in approximately half of all respondents with MDD.⁽¹⁶⁾ Given the high prevalence of MDD among the general adult Singapore population, combined with the significant treatment gap and likelihood of chronic physical comorbidities, the present study aimed to establish whether there were any gender-specific differences relating to the prevalence and correlates of MDD among the adult resident population in Singapore.

METHODS

The SMHS was a cross-sectional epidemiological survey among a representative household sample of Singapore citizens and permanent residents aged ≥ 18 years, who were fluent in English, Mandarin or Malay. Participants were randomly selected from an administrative database that maintains names

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and sociodemographic details, including age, gender, ethnicity and household addresses, of all Singapore residents. There were 2.7 million residents aged ≥ 18 years living in Singapore at the time when the sample was drawn from the sampling frame. A disproportionate stratified sample (based on age group and ethnicity) was used; the three main ethnic groups in Singapore (i.e. Chinese, Malay and Indian) were equally sampled, while older individuals (aged ≥ 65 years) were over-sampled. All participants provided written consent; for those < 21 years, consent was also obtained from a parent or guardian. Residents who were excluded comprised those who were incapable of completing an interview as a result of severe physical or mental conditions, language barriers or living outside the country during the survey period, and those who were not contactable due to an incomplete or incorrect address. Data collection was carried out between December 2009 and December 2010 following approval from the National Healthcare Group's Domain Specific Review Board. During this time, face-to-face interviews were completed with 6,616 respondents, yielding a response rate of 75.9%.

Interviewers from an external survey firm conducted the interviews after undergoing extensive training conducted by research staff at the Institute of Mental Health (IMH), Singapore. Interviewers were taught about ethical aspects of the study, administration of survey measures, and logistical procedures relating to fieldwork and reporting during a three-week intensive training period. Upon passing a detailed evaluation, interviewers were initially closely supervised by IMH staff and field executives from the survey firm. To ensure high-quality data, quality assurance processes were implemented throughout the data collection phase and approximately 20% of each interviewer's cases underwent detailed verification in order to determine any falsification of data. Additional information relating to the methods and procedures employed have been reported in another study.⁽¹⁷⁾

The presence of MDD and other psychiatric disorders was established using the World Health Organization Composite International Diagnostic Interview (CIDI) version 3.0.⁽¹⁸⁾ CIDI 3.0 is a comprehensive, fully structured instrument that assesses mental disorders in terms of 12-month and lifetime prevalence, according to the definitions and criteria outlined by the Diagnostic and Statistical Manual of Mental Disorders, 4th edition (DSM-IV)⁽¹⁹⁾ and the International Classification of Disease, 10th revision (ICD-10).⁽²⁰⁾ The SMHS included the following diagnostic modules: MDD; bipolar disorder; generalised anxiety disorder (GAD); obsessive-compulsive disorder (OCD); and alcohol use disorders (including alcohol abuse and alcohol dependence). Diagnostic hierarchy rules and organic exclusions were applied, where relevant.

Respondents were also asked a series of questions relating to treatment contact. To determine if treatment had ever been sought, respondents were asked whether they had ever 'talked to a medical doctor or other professional' about the disorder. The 'treatment gap' was defined as "*the absolute difference between the true prevalence of a disorder and the treated proportion of individuals affected by the disorder*".⁽²¹⁾

The Sheehan Disability Scale (SDS)⁽²²⁾ was administered and captured functional impairment in three aspects – work/school, social and family life – in the worst month of the past year. Responses were scored on the visual analogue scale (range 0–10), and included the labels none (score 0), mild (score 1–3), moderate (score 4–6), severe (score 7–9) and very severe (score 10).

The depression module in the CIDI includes the Quick Inventory of Depressive Symptomatology – Self-Report (QIDS-SR)⁽²³⁾ which assesses symptom severity in patients with MDD during the worst month of the previous year. Scores from the QIDS-SR were converted into clinical severity scores and categories of the Hamilton Rating Scale for Depression⁽²⁴⁾ based on transformation rules. Categories included none (i.e. not clinically depressed), mild, moderate, severe and very severe. Research has shown very high concordance between the measures.⁽²⁵⁾

A modified version of the CIDI 3.0 checklist for chronic medical conditions was also used. Respondents were read the following statement: "*I'm going to read to you a list of health problems some people have. Has a doctor ever told you that you have any of the following...*". This was followed by a list of 15 chronic conditions that were considered prevalent in Singapore's population. These were then reclassified into the following eight types of physical disorders: (a) respiratory disorders (asthma, chronic lung disease [e.g. chronic bronchitis] or emphysema); (b) diabetes mellitus; (c) hypertension and high blood pressure; (d) chronic pain (arthritis or rheumatism, back problems [including disk or spine] or migraine headaches); (e) cancer; (f) neurological disorders (epilepsy, convulsion or Parkinson's disease); (g) cardiovascular disorders (stroke or major paralysis, heart attack, coronary heart disease, angina, congestive heart failure or other heart diseases); and (h) ulcer and chronic inflamed bowel (stomach ulcer, enteritis or colitis).

Sociodemographic information, including age, gender, ethnicity, education, marital status, income and employment history, was also collected for all respondents. For instruments that were unavailable in Mandarin or Malay, forward translation methods were used to translate these from the English versions.

Statistical analyses were carried out using the Statistical Analysis Software (SAS) System version 9.2 (SAS Institute, Cary, NC, USA). Data was weighted to adjust for oversampling and post-stratified by age and ethnicity distributions between the survey sample and the Singapore resident population in 2007. Descriptive analyses were performed to establish the prevalence of mental disorders and chronic medical conditions, and to describe the sociodemographic characteristics of the study population. We performed multiple logistic regression analyses to examine the odds of having lifetime mental disorders and chronic medical conditions among women when compared to men, after controlling for sociodemographic variables, which included age, ethnicity, marital status, education, employment and income. Analysis of variance and chi-square tests were used to compare the means and rates of continuous and categorical variables between the two groups. Standard errors and significance tests were estimated using the Taylor series linearisation method.

Multivariate significance was evaluated using the Wald chi-square test based on design-corrected coefficient variance-covariance matrices. Statistical significance was evaluated at the 0.05 level using two-sided tests.

RESULTS

Of the 6,616 respondents who completed the study, there were slightly more women (51.5%) than men (48.5%). The majority of the respondents were Chinese (76.9%) followed by Malay (12.3%) and Indian (8.3%); 2.4% of respondents belonged to other ethnic groups (Table I). The lifetime prevalence of MDD among women was higher than that for men (7.2% vs. 4.3%, $p = 0.003$).

Table I shows the sociodemographic correlates of lifetime MDD by gender among the overall sample ($n = 6,616$). Among women, MDD was less likely among those aged

35–49 years and 50–64 years as compared to respondents aged 18–34 years. MDD was also less likely among women with primary education or below as compared to those with university education. MDD was more likely among women who were divorced/separated or widowed as compared to single women. Among men, MDD was more likely among those of Indian and other ethnicities as compared to Chinese men; it was also less likely among divorced/separated men as compared to single men.

A total of 417 respondents with a lifetime diagnosis of MDD were included in the subsequent analysis. Tables II and III show the prevalence of and odds ratio (OR) for lifetime mental and physical disorders, respectively, among respondents with MDD by gender. After adjusting for demographic variables, multiple logistic regression analysis showed that women with MDD had significantly higher odds of having GAD (adjusted OR 6.6, 95%

Table I. Prevalence and sociodemographic correlates of major depressive disorder (MDD) by gender among overall sample ($n = 6,616$).

Variable	No. (weighted %)			p-value	Men		Women	
	Total ($n = 6,616$)	Men ($n = 3,299$)	Women ($n = 3,317$)		OR (95% CI)	p-value	OR (95% CI)	p-value
Prevalence	417 (5.8%)	170 (4.3%)	247 (7.2%)	0.003				
Age (yr)				0.3106				
18–34	2,293 (31.7)	74 (5.6)	129 (11.5)		Ref		Ref	
35–49	2,369 (34.1)	63 (5.3)	73 (5.5)		0.8 (0.4–1.8)	0.630	0.4 (0.3–0.8)	0.005
50–64	1,542 (23.1)	28 (2.1)	40 (5.0)		0.4 (0.14–1.001)	0.050	0.4 (0.2–0.8)	0.017
≥ 65	412 (11.1)	5 (2.5)	5 (5.3)		0.6 (0.2–2.1)	0.435	0.4 (0.2–1.1)	0.080
Ethnicity				0.7498				
Chinese	2,006 (76.9)	37 (3.8)	77 (7.1)		Ref		Ref	
Malay	2,373 (12.3)	41 (3.7)	68 (5.3)		0.9 (0.5–1.5)	0.603	0.8 (0.5–1.2)	0.257
Indian	1,969 (8.3)	72 (6.6)	90 (9.8)		1.7 (1.1–2.6)	0.027	1.2 (0.8–1.8)	0.272
Other	268 (2.4)	20 (12.0)	12 (15.5)		2.9 (1.4–6.1)	0.004	1.7 (0.8–4.0)	0.194
Marital status*				< 0.0001				
Single	1,825 (28.9)	57 (4.5)	80 (8.6)		Ref		Ref	
Married	4,290 (62.4)	99 (3.7)	128 (5.3)		1.5 (0.6–3.8)	0.356	1.2 (0.7–2.1)	0.454
Divorced/separated	262 (4.2)	13 (16.4)	30 (20.4)		6.5 (2.3–18.9)	0.001	6.7 (3.1–14.2)	< 0.0001
Widowed	237 (4.4)	0 (0)	9 (9.1)		–	–	4.4 (1.8–10.8)	0.001
Education				0.8600				
Pre-primary/primary	1,236 (20.2)	15 (2.7)	32 (3.9)		0.6 (0.2–1.9)	0.368	0.4 (0.1–0.9)	0.049
Secondary	1,975 (27.6)	43 (3.8)	68 (6.1)		0.8 (0.3–2.2)	0.713	0.7 (0.3–1.4)	0.281
Pre-university/JC/ diploma	1,342 (22.4)	44 (5.3)	68 (8.4)		1.0 (0.5–2.3)	0.917	0.8 (0.4–1.3)	0.339
Vocational	721 (7.9)	28 (4.4)	18 (10.9)		0.9 (0.3–2.3)	0.798	0.7 (0.3–1.8)	0.487
University	1,342 (21.9)	40 (5.0)	61 (10.4)		Ref		Ref	
Employment status*				0.9451				
Employed	4,594 (71.0)	13 (4.3)	16 (7.8)		Ref		Ref	
Economically inactive	1,522 (24.5)	13 (3.2)	52 (5.5)		0.8 (0.2–2.5)	0.642	0.9 (0.5–1.6)	0.743
Unemployed	313 (4.5)	11 (8.0)	20 (11.6)		1.5 (0.5–4.3)	0.450	1.9 (0.8–4.4)	0.143
Annual income (SGD)*				0.6641				
< 20,000	3,392 (51.3)	70 (4.6)	134 (6.3)		1.5 (0.6–3.5)	0.390	1.0 (0.5–2.0)	0.924
20,000–49,999	1,924 (31.2)	57 (4.2)	79 (8.8)		1.9 (0.7–5.4)	0.228	1.0 (0.5–2.1)	0.991
≥ 50,000	962 (17.5)	36 (4.2)	24 (9.6)		Ref		Ref	

*Data has missing values. CI: confidence interval; JC: junior college; OR: odds ratio; Ref: reference group

Table II. Prevalence of lifetime mental disorders among respondents with major depressive disorder (MDD) by gender and results of multiple logistic regression analysis (n = 417).

Variable	No. (weighted %)		p-value	Women vs. men	
	Men (n = 170)	Women (n = 247)		Adjusted OR (95% CI)	p-value
GAD	6 (1.6)	23 (8.0)	0.001*	6.6 (2.0–21.5)	0.002*
OCD	20 (10.1)	30 (10.6)	0.912	1.0 (0.3–3.0)	0.954
Alcohol abuse	20 (14.2)	11 (4.9)	0.017	0.4 (0.1–1.0)	0.056
Alcohol dependence	5 (1.8)	4 (2.2)	0.750	4.5 (0.4–51.3)	0.222
Any mental disorder	51 (27.1)	77 (28.8)	0.788	0.9 (0.4–1.9)	0.822

Data adjusted by age group, ethnicity, marital status, education, employment and income. *p < 0.05 is statistically significant. CI: confidence interval; GAD: generalised anxiety disorder; OCD: obsessive compulsive disorder; OR: odds ratio

Table III. Prevalence of lifetime chronic physical conditions among respondents with major depressive disorder (MDD) by gender and results of multiple logistic regression analysis (n = 417).

Variable	No. (weighted %)		p-value	Women vs. men	
	Men (n = 170)	Women (n = 247)		Adjusted OR (95% CI)	p-value
Respiratory conditions	33 (15.2)	41 (12.7)	0.599	1.0 (0.4–2.4)	0.957
Diabetes mellitus	18 (11.5)	15 (4.4)	0.051	0.5 (0.01–2.2)	0.379
High blood pressure	29 (26.5)	27 (12.7)	0.033*	0.2 (0.1–0.7)	0.006*
Chronic pain	34 (24.2)	86 (33.6)	0.188	1.6 (0.7–3.7)	0.223
Cancer†	4 (0.9)	1 (0.1)	0.065	–	–
Neurological conditions*	3 (3.7)	9 (5.4)	0.698	–	–
Cardiovascular disease	11 (3.9)	6 (5.0)	0.724	0.1 (0.01–1.1)	0.061
Ulcer	7 (3.4)	5 (1.7)	0.405	0.4 (0.04–2.6)	0.289
Any chronic physical condition	87 (52.4)	127 (47.4)	0.507	0.9 (0.5–1.8)	0.778

Data adjusted by age group, ethnicity, marital status, education, employment and income. *p < 0.05 is statistically significant. †Estimates were not reported due to small sample size. CI: confidence interval; OR: odds ratio

confidence interval [CI] 2.0–21.5; p = 0.002, Table II) but lower odds of having high blood pressure (OR 0.2, 95% CI 0.1–0.7; p = 0.006, Table III) as compared to men.

Table IV shows the age of onset, severity and treatment gap among people with MDD by gender. Women with lifetime MDD tended to have a slightly later age of onset of MDD. Severity of impairment, based on MDD over the past 12 months and according to SDS and HAM-D, showed that women had less severe impairment when compared to men. Women also had a lower treatment gap compared to men (67.6% vs. 75.3%, p = 0.290). However, none of these differences were statistically significant.

DISCUSSION

A number of gender differences were observed among respondents with MDD. Firstly, the prevalence of MDD was higher among women (7.2%) compared to men (4.3%), a finding that has been consistently reported in psychiatric epidemiology. The prevalence of MDD among women in these studies is typically reported to be 1.5–3 times higher than that observed in men,⁽¹¹⁾ which is consistent with our findings. While the exact reason for such gender differences in relation to MDD prevalence is not known, it is likely to be a myriad of social, behavioural, psychological and biological factors that possibly interact with one another. More specifically, risk factors (e.g. biological susceptibility resulting

Table IV. Age of onset, severity and treatment gap among respondents with major depressive disorder (MDD) by gender.

Variable	No. (weighted %)		p-value
	Men (n = 170)	Women (n = 247)	
Mean age of onset (yr)	27.5	28.3	0.709
Severity of role impairment*			
By SDS score†			0.061
None	1 (0.4)	6 (7.0)	
Mild	5 (4.2)	12 (16.3)	
Moderate	31 (48.8)	47 (37.0)	
Severe	26 (40.8)	27 (32.3)	
Very severe	4 (5.9)	8 (7.5)	
By HAM-D score‡,§			0.589
None	1 (0.6)	2 (4.0)	
Mild	4 (2.3)	4 (7.4)	
Moderate	6 (9.6)	7 (6.1)	
Severe	21 (39.9)	28 (35.8)	
Very severe	25 (47.5)	46 (46.6)	
Treatment gap	122 (75.3)	152 (67.6)	0.290

*Severity of role impairment was only measured among respondents with MDD during the previous 12 months (n = 181). †14 cases were missing. ‡37 cases were missing. §Transformation rules developed for the Quick Inventory of Depressive Symptomatology – Self-Report were used to convert scores into clinical severity categories of the Hamilton Rating Scale for Depression (HAM-D). SDS: Sheehan Disability Scale

from hormonal mechanisms, women being more likely and open to seeking help, and social and cultural influences) that have been deemed to lead to stress and added difficulties among women have been explored. However, these have yet to be determined with any certainty.⁽²⁶⁾

While these gender gaps occur among patients with MDD and other mental disorders, it appears that the gap is narrowing in some countries.^(27,28) Consequently, when explaining gender differences in mental disorders, there has been a shift toward the role of typical stressors, coping resources, and the opportunities available to men and women for expressing psychological distress.⁽⁸⁾ This needs to be further explored, particularly as gender roles have changed over time in many parts of the world, including Singapore.

In the overall sample, women with MDD were less likely to be older (35–49 years and 50–64 years compared to 18–34 years). This is consistent with previous research, which has shown that younger age is a risk factor for MDD,⁽²⁹⁾ although this finding is not gender specific. MDD was also more likely among respondents belonging to the younger age group (age 18–34 years) in the overall sample;⁽¹⁶⁾ when the sample was split by gender, it became evident that women were driving this association. Possible reasons that may account for these differences between the genders at a younger age include psychological characteristics; for instance, neuroticism may result in more vulnerable responses to life events.⁽³⁰⁾ While women are at greater risk of MDD at a younger age, compared to men, this may only partly account for their preponderance in rates of adult MDD.⁽²⁶⁾ In cases where age-related recall bias is less likely to be an influence, it is not surprising that the women with MDD are more likely to be younger, as MDD has been known to have an early age of onset irrespective of gender.^(1,29,31)

Unsurprisingly, being divorced or separated was also a risk factor for MDD among both men and women when compared to single respondents, and this is likely explained by the nature of such a stressful life event. The cross-sectional nature of this study did not allow us to determine whether MDD caused divorce/separation or vice versa. However, it is thought to be bidirectional⁽³²⁾ and could also be explained by a number of social processes. For example, people with a mental illness may be less likely to get married or more likely to experience more marital difficulties, which may result in divorce or separation.⁽³³⁾ Widowed women also had higher odds of MDD than their single counterparts, as the death of a spouse may have adversely affected their mental health.⁽³⁴⁾ Contrary to some findings that marriage was a ‘buffer’ or protective factor for depression,^(3,35) others have found that married women were more likely to have higher rates of depression compared to those who were divorced or separated;⁽³⁶⁾ however, we did not find this association among our sample.

Interestingly, we found that, among women, respondents with less education (pre-primary or primary school education only) had lower odds of MDD than those with university-level education. The association of education with MDD status and risk is debatable. Several studies have found that

MDD risk did not differ by education,^(3,9) while a recent study indicated a lower depression rate among those with more education, with the association between higher education and improved mental health being significantly more prominent among women compared to men.⁽³⁵⁾ While we are unable to ascertain the exact reason behind our findings, it is important to highlight that the number of men ($n = 15$) and women ($n = 32$) with MDD who had pre-primary or primary education in our study was quite small, and therefore this finding should be viewed with caution.

In our study, the risk of MDD was higher among men of Indian ethnicity when compared to Chinese men. Currently, we are unable to elucidate why Indian men are at higher risk of MDD. This could be due to a biological vulnerability that has yet to be explained, or environmental factors, including acculturation, resulting in the internalisation of disorders such as depression among ethnic minority groups.⁽³⁷⁾ In Singapore, Indians are a minority ethnic group compared to the Chinese and Malays. Notably, previous research, conducted largely in the United States, found a lower-than-expected prevalence of depression among minority ethnic groups compared to white Americans.^(38–40) These ethnic or racial disparities may be a result of methodological procedures, including inconsistencies and discrepancies in how depression is measured, as well as how racial or ethnic groups are classified and the use of different controls as confounding factors. Caution is required when interpreting findings based on ethnic or racial groups, as comparisons are filled with challenges and complexities. Ethnic or racial categories are “*at best approximations of societally defined groupings to which individuals are assigned based largely on skin colour, country of origin or ... language or dialect spoken*”,⁽⁴¹⁾ and therefore may mask wide-ranging heterogeneity and complicate more granular differences.

Results from our study showed that among the subsample of respondents with MDD, there were a number of gender differences relating to other psychiatric comorbidities. For example, GAD was significantly more prevalent among women compared to men, and women with MDD were 6.2 times more likely to have a comorbid GAD diagnosis during their lifetime. Similar to MDD, psychiatric epidemiology has consistently found that an anxiety disorder is significantly more likely to occur in women during their lifetime, as compared to men.^(42–44) It is important to note, however, that in our subsample of respondents with MDD, there was only a small number of men with GAD, which resulted in a wide CI.

Among those with MDD, men were significantly more likely to have high blood pressure compared to women ($p < 0.006$). Hypertension prevalence, as reported in a worldwide systematic review, tends to occur more frequently among men than women in most countries.⁽⁴⁵⁾ Various explanations have been proposed for gender differences in hypertension among respondents with MDD, including social factors. For example, research has shown that men are less likely to perceive themselves as being at risk of developing various health problems, and the two genders

generally have different opinions on healthy behaviours.^(46,47) Yang and Reckelhoff also attribute these gender differences to sex hormones, suggesting that premenopausal women are comparatively protected against hypertension in comparison to postmenopausal women and men.⁽⁴⁸⁾ The cross-sectional nature of this study, however, did not allow any causal relationships to be established.

In addition to the psychiatric and physical comorbidities observed among respondents with MDD, we also explored gender differences by age of onset, impairment severity and treatment gap. Women with lifetime MDD tended to have a slightly later age of onset of MDD, but less severe impairment based on SDS and HAM-D. Women also had a lower treatment gap compared to men. However, none of these differences were statistically significant. A cross-sectional epidemiological study, conducted by Gili et al, which explored gender differences in disability among those with MDD, also found no statistically significant difference in functioning between men and women, suggesting that the relationship between depression and functioning or impairment is not gender-dependent.⁽⁴⁹⁾

Our findings should be viewed in the context of some limitations. First, depressive symptoms were assessed based on self-report and may be subject to various biases. Furthermore, it has been suggested that women may be more willing to admit their depressive symptoms or experiences to an interviewer than men.⁽⁵⁰⁾ Second, we were unable to establish the cause-and-effect relationship of mental and physical disorders due to the cross-sectional nature of the study. Third, people residing in nursing homes, prisons and hospitals were excluded from the survey. Lastly, due to respondent burden, we were unable to include all psychiatric disorders or physical illnesses that may co-occur with MDD, which might have resulted in an underestimation of MDD comorbidities.

These limitations notwithstanding, the strengths of our study are the use of a well-established instrument for collecting information, a large sample size and a high response rate that provides confidence in the results and improves generalisability. This study has highlighted key gender-specific predictors and risk factors for MDD. Given the comorbidities with MDD and other psychiatric disorders and/or physical illnesses, these correlates pose additional challenges for care providers, and also emphasise the importance of early detection and screening of conditions such as hypertension, particularly among men with MDD. Furthermore, the finding that MDD is more prevalent among women, particularly in the younger age group, suggests the need to involve and educate parents and teachers about the signs of depression. The importance of establishing early detection and screening systems in various settings, including educational institutions and general medical practices, is key and could include innovative measures, such as the Internet or telephone-based therapy.

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REFERENCES

1. Kessler RC, Berglund P, Demler O, et al; National Comorbidity Survey Replication. The epidemiology of major depressive disorder: results from the National Comorbidity Survey Replication (NCS-R). *JAMA* 2003; 289:3095-105.
2. Kessler RC, Berglund P, Demler O, et al. Lifetime prevalence and age-of-onset distributions of DSM-IV disorders in the National Comorbidity Survey Replication. *Arch Gen Psychiatry* 2005; 62:593-602.
3. Hasin DS, Goodwin RD, Stinson FS, Grant BF. Epidemiology of major depressive disorder: results from the National Epidemiologic Survey on Alcoholism and Related Conditions. *Arch Gen Psychiatry* 2005; 62:1097-106.
4. Alonso J, Angermeyer MC, Bernert S, et al; ESEMeD/MHEDEA 2000 Investigators, European Study of the Epidemiology of Mental Disorders (ESEMeD) Project. Prevalence of mental disorders in Europe: results from the European Study of the Epidemiology of Mental Disorders (ESEMeD) project. *Acta Psychiatr Scand Suppl* 2004; (420):21-7.
5. Gu L, Xie J, Long J, et al. Epidemiology of major depressive disorder in mainland china: a systematic review. *PLoS One* 2013; 8:e65356.
6. Chong SA, Abidin E, Vaingankar JA, et al. A population-based survey of mental disorders in Singapore. *Ann Acad Med Singapore* 2012; 41:49-66.
7. Orui M, Kawakami N, Iwata N, Takeshima T, Fukao A. Lifetime prevalence of mental disorders and its relationship to suicidal ideation in a Japanese rural community with high suicide and alcohol consumption rates. *Environ Health Prev Med* 2011; 16:384-9.
8. Seedat S, Scott KM, Angermeyer MC, et al. Cross-national associations between gender and mental disorders in the World Health Organization World Mental Health Surveys. *Arch Gen Psychiatry* 2009; 66:785-95.
9. Patten SB, Wang JL, Williams JV, et al. Descriptive epidemiology of major depression in Canada. *Can J Psychiatry* 2006; 51:84-90.
10. Regier DA, Narrow WE, Rae DS, et al. The de facto US mental and addictive disorders service system. Epidemiologic catchment area prospective 1-year prevalence rates of disorders and services. *Arch Gen Psychiatry* 1993; 50:85-94.
11. Kessler RC. Epidemiology of women and depression. *J Affect Disord* 2003; 74:5-13.
12. Weissman MM, Olfson M. Depression in women: implications for health care research. *Science* 1995; 269:799-801.
13. Burt VK. Women and depression: Special considerations in assessment and management. In: Lewis-Hall F, Williams TS, Panetta JA, Herrera JM, eds. *Psychiatric Illness in Women: Emerging Treatments and Research*. Washington, DC: American Psychological Publishing Inc, 2002: 113-6.
14. Afifi M. Gender differences in mental health. *Singapore Med J* 2007; 48:385-91.
15. Department of Statistics, Singapore. *Population Trends 2013*. Available at: http://www.singstat.gov.sg/Publications/publications_and_papers/population_and_population_structure/population2013.pdf. Accessed June 5, 2014.
16. Chong SA, Vaingankar J, Abidin E, Subramaniam M. The prevalence and impact of major depressive disorder among Chinese, Malays and Indians in an Asian multi-racial population. *J Affect Disord* 2012; 138:128-36.
17. Subramaniam M, Vaingankar J, Heng D, et al. The Singapore Mental Health Study: an overview of the methodology. *Int J Methods Psychiatr Res* 2012; 21:149-57.
18. Kessler RC, Üstün TB. The World Mental Health (WMH) Survey Initiative Version of the World Health Organization (WHO) Composite International Diagnostic Interview (CIDI). *Int J Methods Psychiatr Res* 2004; 13:93-121.
19. American Psychiatric Association. *Diagnostic and statistical manual of mental disorders (4th ed)*. Washington, DC: American Psychiatric Association, 1994.
20. World Health Organization. *The ICD-10 Classification of Mental and Behavioural Disorders: Clinical Descriptions and Diagnostic Guidelines*. Geneva: World Health Organization, 1992.
21. Kohn R, Saxena S, Levav I, Saraceno B. The treatment gap in mental health care. *Bull World Health Organ* 2004; 82:858-66.
22. Sheehan DV, Harnett-Sheehan K, Raj BA. The measurement of disability. *Int Clin Psychopharmacol* 1996; 11 Suppl 3:89-95.
23. Rush AJ, Carmody T, Reimtz PE. The Inventory of Depressive Symptomatology (IDS): Clinician (IDS-C) and Self-Report (IDS-SR) ratings of depressive symptoms. *Int J Methods Psychiatr Res* 2000; 9:45-59.
24. Hamilton M. A rating scale for depression. *J Neurol Neurosurg Psychiatry* 1960; 23:56-62.
25. Rush, AJ, Trivedi MH, Ibrahim HM, et al. The 16-Item Quick Inventory of Depressive Symptomatology (QIDS), clinician rating (QIDS-C), and self-report (QIDS-SR): a psychometric evaluation in patients with chronic major depression. *Biol Psychiatry* 2003; 54:573-83.
26. Piccinelli M, Wilkinson G. Gender differences in depression. Critical review. *Br J Psychiatry* 2000; 177:486-92.
27. Joyce PR, Oakley-Browne MA, Wells JE, Bushnell JA, Hornblow AR. Birth cohort trends in major depression: increasing rates and earlier onset in New Zealand. *J Affect Disord* 1990; 18:83-9.
28. Wickramaratne PJ, Weissman MM, Leaf PJ, Holford TR. Age, period and cohort effects on the risk of major depression: results from five United States communities. *J Clin Epidemiol* 1989; 42:333-43.
29. Blazer DG, Kessler RC, McGonagle KA, Swartz MS. The prevalence and

- distribution of major depression in a national community sample: the National Comorbidity Survey. *Am J Psychiatry* 1994; 151:979-86.
30. Wilhelm K, Parker G, Hadzi-Pavlovic D. Fifteen years on: evolving ideas in researching sex differences in depression. *Psychol Med* 1997; 27:875-83.
 31. Andrade L, Caraveo-Anduaga JJ, Berglund P, et al. The epidemiology of major depressive episodes: results from the International Consortium of Psychiatric Epidemiology (ICPE) Surveys. *Int J Methods Psychiatr Res* 2003; 12:3-21.
 32. Bulloch AG, Williams JV, Lavorato DH, Patten SB. The relationship between major depression and marital disruption is bidirectional. *Depress Anxiety* 2009; 26:1172-7.
 33. Green RG. The influence of divorce prediction variables on divorce adjustment: an expansion and test of Lewis' and Spanier's theory of marital quality and marital stability. *J Divorce* 1983; 7:67-81.
 34. Carey RG. Weathering widowhood: problems and adjustment of the widowed during the first year. *Omega J Death Dying* 1979; 10:163-74.
 35. Van de Velde S, Bracke P, Levecque K. Gender differences in depression in 23 European countries. Cross-national variation in the gender gap in depression. *Soc Sci Med* 2010; 71:305-13.
 36. Parker G, Brotchie H. Gender differences in depression. *Int Rev Psychiatry* 2010; 25:429-36.
 37. Anderson ER, Mayes LC. Race/ethnicity and internalizing disorders in youth: a review. *Clin Psychol Rev* 2010; 30:338-48.
 38. Breslau J, Kendler KS, Su M, Gaxiola-Aguilar S, Kessler RC. Lifetime risk and persistence of psychiatric disorders across ethnic groups in the United States. *Psychol Med* 2005; 35:317-27.
 39. Breslau J, Aguilar-Gaxiola S, Kendler KS, et al. Specifying race-ethnic differences in risk for psychiatric disorder in a USA national sample. *Psychol Med* 2006; 36:57-68.
 40. Kessler RC, McGonagle KA, Zhao S, et al. Lifetime and 12-month prevalence of DSM-III-R psychiatric disorders in the United States. Results from the National Comorbidity Survey. *Arch Gen Psychiatry* 1994; 51:8-19.
 41. Kaplan JB, Bennett T. Use of race and ethnicity in biomedical publication. *JAMA* 2003; 289:2709-16.
 42. Angst J, Dobler-Mikola A. The Zurich Study. V. Anxiety and phobia in young adults. *Eur Arch Psychiatry Neurol Sci* 1985; 235:171-8.
 43. Bruce SE, Yonkers KA, Otto MW, et al. Influence of psychiatric comorbidity on recovery and recurrence in generalized anxiety disorder, social phobia, and panic disorder: a 12-year prospective study. *Am J Psychiatry* 2005; 162:1179-87.
 44. Regier DA, Narrow WE, Rae DS. The epidemiology of anxiety disorders: the Epidemiologic Catchment Area (ECA) experience. *J Psychiatr Res* 1990; 24 Suppl 2:3-14.
 45. Kearney PM, Whelton M, Reynolds K, Whelton PK, He J. Worldwide prevalence of hypertension: a systematic review. *J Hypertens* 2004; 22:11-9.
 46. Flynn J, Slovic P, Mertz CK. Gender, race and perception of environmental health risks. *Risk Anal* 1994; 14:1101-8.
 47. White A. How men respond to illness. *Men's Health J*; 1:18-9.
 48. Yang XP, Reckelhoff JF. Estrogen, hormonal replacement therapy and cardiovascular disease. *Curr Opin Nephrol Hypertens* 2011; 20:133-8.
 49. Gili M, Castro A, Navarro C, et al. Gender differences on functioning in depressive patients. *J Affect Disord* 2014; 166:292-6.
 50. Young MA, Fogg LF, Scheftner WA, Keller MB, Fawcett JA. Sex differences in the lifetime prevalence of depression: does varying the diagnostic criteria reduce the female/male ratio? *J Affect Disord* 1990; 18:187-92.