

# Age patterns in the prevalence of DSM-IV depressive/anxiety disorders with and without physical co-morbidity

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**Background.** Physical morbidity is a potent risk factor for depression onset and clearly increases with age, yet prior research has often found depressive disorders to decrease with age. This study tests the possibility that the relationship between age and mental disorders differs as a function of physical co-morbidity.

**Method.** Eighteen general population surveys were carried out among household-residing adults as part of the World Mental Health (WMH) surveys initiative ( $n = 42\,697$ ). DSM-IV disorders were assessed using face-to-face interviews with the Composite International Diagnostic Interview (CIDI 3.0). The effect of age was estimated for 12-month depressive and/or anxiety disorders with and without physical or pain co-morbidity, and for physical and/or pain conditions without mental co-morbidity.

**Results.** Depressive and anxiety disorders decreased with age, a result that cannot be explained by organic exclusion criteria. No significant difference was found in the relationship between mental disorders and age as a function of physical/pain co-morbidity. The majority of older persons have chronic physical or pain conditions without co-morbid mental disorders; by contrast, the majority of those with mental disorders have physical/pain co-morbidity, particularly among the older age groups.

**Conclusions.** CIDI-diagnosed depressive and anxiety disorders in the general population decrease with age, despite greatly increasing physical morbidity with age. Physical morbidity among persons with mental disorder is the norm, particularly in older populations. Health professionals, including mental health professionals, need to address barriers to the management of physical co-morbidity among those with mental disorders.

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**Key words:** Age, anxiety disorder, co-morbidity, depressive disorder, physical.

## Introduction

The prevalence of major depressive disorder (MDD) in the general population has frequently been found to decrease with increasing age (Regier *et al.* 1988;

Karel, 1997; Beekman *et al.* 1999; Kessler *et al.* 2003a; Alonso *et al.* 2004; Pirkola *et al.* 2005; Wells *et al.* 2006; Troller *et al.* 2007), and the picture for anxiety disorders is similar (Regier *et al.* 1988; Jorm, 2000; Alonso *et al.* 2004; Wells *et al.* 2006; Troller *et al.* 2007). A number of methodological reasons have been advanced for this, including selective mortality, excluding the institutionalized from the sample and cohort effects (Karel, 1997; Jorm, 2000; Snowdon, 2001;

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Hybels & Blazer, 2002; O'Connor, 2006). With the exception of the cohort effects, these sampling bias explanations are not thought to be sufficient to explain this age-related pattern (Jorm, 2000; Hybels & Blazer, 2002; Troller *et al.* 2007). As to whether the decrease in 12-month or 1-month depressive disorder with age is a cohort or age effect, there is as yet insufficient clear evidence to draw conclusions about this (Jorm, 2000), but it is noteworthy that the same age-related decline in prevalence has now occurred in cross-national surveys conducted in the USA nearly two decades apart (Regier *et al.* 1988; Kessler *et al.* 2003a), which is more supportive of an age-related than a cohort effect.

Complicating the picture, scale measures of depressive symptoms (e.g. the Center for Epidemiologic Studies Depression Scale, CES-D) frequently show an increase with age (Beekman *et al.* 1995; Jorm, 2000; Hybels & Blazer, 2002; Stordal *et al.* 2003; van't Veer-Tazelaar *et al.* 2008), although decreases (Henderson *et al.* 1998), and U-shaped patterns (Newmann, 1989; Kessler *et al.* 1992) have also been observed. Prevalences of 'clinically significant depression' (measured by the CES-D and/or inclusive of subthreshold depression) among the  $\geq 65$  years age group are generally in the range of 10–20% (Beekman *et al.* 1995, 1999; Snowdon, 2001), compared with prevalences of MDD typically less than 3% (Beekman *et al.* 1999; Alonso *et al.* 2004; Wells *et al.* 2006; Troller *et al.* 2007).

The contrast between the age patterning and prevalence estimates of CES-D depressive symptoms *versus* MDD has fueled suspicion about the validity of diagnostic criteria in older persons, and their operationalization in standardized interviews such as the Composite International Diagnostic Interview (CIDI) (Beekman *et al.* 1995; Mulsant & Ganguli, 1999; Snowdon, 2001; Hybels & Blazer, 2002; O'Connor, 2006). It has been suggested that depression may manifest differently with age, with older people being more likely to report somatic symptoms and less likely to report required diagnostic symptoms of depressed mood and anhedonia (Gallo *et al.* 1994; Karel, 1997; Christensen *et al.* 1999; Jorm, 2000; Hybels & Blazer, 2002). It has also been argued that the attribution of depressive symptoms to a co-morbid physical condition increases with age independent of physical health status (Knauper & Wittchen, 1994), with the result that standardized diagnostic interviews may underdiagnose in older people because they exclude symptoms attributed to physical disease (Knauper & Wittchen, 1994; O'Connor, 2006). Knauper & Wittchen (1994) interpret their finding of increasing physical illness symptom attribution with age as a simplifying cognitive heuristic in response to the complex stem questions and probing for physical

causes for every depressive symptom in earlier versions of the CIDI. They suggest that changing the structure of the physical cause probe system so that it occurs after assessment of the complete depressive episode should address this problem. This change has been implemented in the version of the CIDI used in the surveys that form the basis of this report.

What is not in dispute is that the prevalence of chronic physical conditions increases with age. There is also substantial evidence that mental disorders are more common among persons with physical illness (Wells *et al.* 1989; Dew, 1998; Harter *et al.* 2003; Evans *et al.* 2005; Scott *et al.* 2007). Moreover, longitudinal studies show that physical disease is a potent risk factor for depression and anxiety episode onset among the elderly (Schoevers *et al.* 2000; Brilman & Ormel, 2001; de Graaf *et al.* 2002; Krishnan, 2002). In this context, it is perhaps curious to find a decrease in diagnosed depression with increasing age, even taking into account the methodological issues mentioned above.

One possibility is that the relationship between age and depression differs as a function of whether mental disorders are accompanied by physical condition co-morbidity. On the basis of prior research, it might be expected that co-morbid depression (here referring to co-morbidity with a chronic physical and/or chronic pain condition) would increase with age whereas non-co-morbid depression would decrease with age. This paper, based on 18 of the World Mental Health (WMH) surveys (Kessler & Ustun, 2004), examines this possibility by testing whether there is an interaction of age with the presence of physical/pain condition co-morbidity in the association with 12-month depressive or anxiety disorder, and by showing the effect of age in those with 12-month mental disorders disaggregated into those with and without physical/pain co-morbidity. The relative proportion of co-morbid to non-co-morbid depression in each age group is also determined. Relevant to the methodological issues raised above, we show the percentage of depression cases excluded for organic (physical) causes in each age group. Our aims are thus both analytical and descriptive. Describing how mental-physical co-morbidity varies across age groups in general population samples is important for guiding the work of both health-care and mental health-care professionals in appreciating the overall health problems of patients they treat.

## Method

### *Samples*

Eighteen surveys were carried out in 17 countries in the Americas (Colombia, Mexico, the USA), Europe

(Belgium, France, Germany, Italy, The Netherlands, Spain, Ukraine), the Middle East (Israel, Lebanon), Africa (Nigeria, South Africa), Asia (Japan, separate surveys in Beijing and Shanghai in the People's Republic of China) and the South Pacific (New Zealand). All surveys were based on multi-stage, clustered area probability household samples and were carried out face-to-face by trained lay interviewers. Cognitive impairment was not systematically screened for, but given the demands of the lengthy interview, the sample effectively represents the cognitively intact. Sample sizes range from 2372 (The Netherlands) to 12992 (New Zealand), with a total of 85088 respondents. The age ranges reported here include those from 18 years and over, but Mexico and Colombia did not sample beyond 65 years and Beijing and Shanghai did not sample beyond 70 years. Response rates range from 45.9% (France) to 87.7% (Colombia), with a weighted average response rate of 70.8%.

Internal subsampling was used to reduce respondent burden and reduce average interview time by dividing the interview into two parts. Part 1 included the core diagnostic assessment of mental disorders. Part 2 included additional information relevant to a wide range of survey aims, including assessment of chronic physical conditions. All respondents completed Part 1. All Part 1 respondents who met criteria for any mental disorder and a probability sample of other respondents were administered Part 2. Part 2 respondents were weighted by the inverse of their probability of selection for Part 2 of the interview to adjust for differential sampling. Analyses in this article were based on the weighted Part 2 subsample ( $n=42\,697$ ). Additional weights were used to adjust for differential probabilities of selection, adjust for non-response and to match the samples to population sociodemographic distributions.

### *Training and field procedures*

The central WMH staff trained bilingual supervisors in each country. The World Health Organization (WHO) translation protocol was used to translate instruments and training materials. Some surveys were carried out in bi- or multi-lingual form (Belgium; Ukraine, Israel, Nigeria). Other surveys were carried out exclusively in the country's official language. Persons who could not speak these languages were excluded. Quality control protocols, described in more detail elsewhere (Kessler *et al.* 2004), were standardized across countries to check on interviewer accuracy and to specify data cleaning and coding procedures. The institutional review board of the organization that coordinated the survey in each

country approved and monitored compliance with procedures for obtaining informed consent and protecting human subjects.

### *Mental disorder status*

All surveys used the WMH survey version of the WHO-CIDI (now CIDI 3.0; Kessler & Ustun, 2004), a fully structured diagnostic interview, to assess disorders and treatment. Disorders were assessed using DSM-IV definitions and criteria (APA, 1994). CIDI organic exclusion rules were imposed for diagnoses of MDD and panic disorder. The probe for physical illness was applied at the episode level (i.e. after all individual symptom questions had been asked). Respondents were asked whether they considered that their episode of symptoms was *ever* 'the result of physical causes such as physical illness or injury or the use of medication, drugs or alcohol'. If they responded in the affirmative, they were then asked if their episode was *always* the result of physical causes. If they responded in the affirmative again, they were asked to specify the causes, which were recorded as open text. In all countries, a mental health clinician subsequently evaluated the open text responses and coded them as a legitimate organic exclusion or not. An episode of MDD was excluded on organic grounds only if the respondents indicated that their episodes were *always* due to physical causes and if the clinician coded them as legitimate organic exclusions. The percentage of MDD episodes excluded as organic, by age group, for all countries combined was: 4.4% (18–34 years); 6.6% (35–49 years); 8.0% (50–64 years); 8.9% (65–79 years); and 11.1% ( $\geq 80$  years).

This paper includes 12-month anxiety disorders (generalized anxiety disorder, panic disorder and/or agoraphobia, post-traumatic stress disorder, and social phobia) and depressive disorders (dysthymia and MDD). Anxiety and depressive disorders were aggregated into a single category, on the basis of prior findings from the WMH surveys that anxiety disorders and depressive disorders have equal and independent relationships with a wide range of chronic physical conditions (Scott *et al.* 2007).

### *Chronic physical conditions*

Physical conditions were assessed with a standard chronic condition checklist adapted from the US Health Interview Survey (NCHS, 1994). Prior research has demonstrated reasonable correspondence between self-reported chronic conditions, such as diabetes, heart disease and asthma, and general practitioner records (Kriegsman *et al.* 1996).

**Table 1.** World Mental Health Survey questions on chronic physical conditions

The next few questions are about health problems you might have had at any time in your life.

Have you ever had any of the following?<sup>a,b</sup>

- Arthritis or rheumatism
- Chronic back or neck problems<sup>c</sup>
- Frequent or severe headaches<sup>c</sup>
- Any other chronic pain<sup>c</sup>
- A stroke
- A heart attack

Did a doctor or other health professional ever tell you that you had any of the following illnesses?<sup>b</sup>

- Heart disease
- High blood pressure<sup>c</sup>
- Asthma
- Any other chronic lung disease such as COPD or emphysema
- Diabetes or high blood sugar<sup>c</sup>
- An ulcer in your stomach or intestine<sup>c</sup>
- HIV infection or AIDS
- Tuberculosis<sup>c</sup>
- Epilepsy or seizures
- Cancer

COPD, Chronic obstructive pulmonary disease.

<sup>a</sup> In Nigeria, Lebanon, China and Ukraine the first block of questions (from arthritis to heart attack) was asked only in a 12-month format: 'Have you had any of the following in the past 12 months?', and the follow-up question (below) was not asked.

<sup>b</sup> Participants had to respond to each condition (yes/no/don't know/refused).

<sup>c</sup> If these conditions were endorsed by the participant, the following question was asked: 'Did you still have (the condition) or receive any treatment for (it/them) at any time during the past 12 months?'

For all countries except Nigeria, Lebanon, China and Ukraine, chronic physical conditions were screened for by asking participants in the Part 2 subsample if they had ever had arthritis, rheumatism, chronic back or neck problems, frequent or severe headaches, other chronic pain, seasonal allergies, stroke, heart attack, and whether they had ever been told by a doctor they had had heart disease, high blood pressure, asthma, tuberculosis, chronic obstructive pulmonary disease (COPD), diabetes, ulcer, HIV/AIDS, epilepsy or cancer (Table 1). For problems that could have remitted, participants were asked if they still had the condition(s) in the past 12 months, except in Nigeria, Lebanon, China and Ukraine (Table 1). The 12-month prevalence of these latter conditions is used in this paper. For the analyses reported here, the physical conditions were aggregated

and included: stroke, heart attack, heart disease, asthma, COPD, diabetes, ulcer, HIV/AIDS, epilepsy, tuberculosis and cancer. The pain conditions were also aggregated and included: arthritis, chronic back/neck problems, frequent or severe headaches and other chronic pain.

### Analysis methods

Prevalences were estimated for four groups, those with: (i) a 12-month depressive and/or anxiety disorder with a co-morbid pain condition, (ii) a 12-month depressive and/or anxiety disorder with a co-morbid physical condition, (iii) a 12-month depressive and/or anxiety disorder without a co-morbid physical or pain condition, and (iv) a physical and/or pain condition without co-morbid depressive/anxiety disorder. Groups (i) and (ii) are not mutually exclusive. Group (iv) is included to provide context for the mental disorder estimates.

Odds ratios for the effect of age (reference level 18–34 years) in each of the four mental and/or physical condition combinations specified above were calculated for all countries combined, in logistic regression models controlling for gender. The interaction of age with physical or pain co-morbidity (present *versus* absent) in predicting 12-month depressive and/or anxiety disorder was assessed in a logistic regression model on the pooled dataset, controlling for gender.

A separate set of analyses using those with a 12-month depressive and/or anxiety disorder (i.e. cases) as denominator calculated the percentage with a co-morbid physical and/or pain condition, by age group, for all countries combined. All analyses were run with SUDAAN version 8.0.1 (SUDAAN, 2002) to adjust for clustering and weighting.

## Results

### Sample characteristics

The combined sample of those who completed the longer version of the interview (Part 1+Part 2) including the physical condition checklist was 42 697. The Part 2 sample in each country ranged in size from the smaller Asian surveys in Japan (887), Beijing (914) and Shanghai (714) to the larger samples in New Zealand (7312), the USA (5692), Israel (4859) and South Africa (4315). The proportion of the sample that was age 60 or greater was higher in the developed countries than the developing countries, and the percentage with 12 or more years of education was also generally higher in the developed countries. Further detail on sample characteristics is provided elsewhere (Scott *et al.* 2007).

**Table 2.** Percentage (and 95% confidence interval) in each of the four mental and/or physical condition groups, by country

Country	Depressive/anxiety disorder with pain condition <sup>a</sup>	Depressive/anxiety disorder with physical condition <sup>a</sup>	Depressive/anxiety disorder without physical or pain condition <sup>a</sup>	Physical and/or pain condition without depressive/anxiety disorder <sup>a</sup>
Colombia	4.8 (3.9–5.8)	3.1 (2.4–4.1)	4.4 (3.6–5.4)	32.1 (29.3–34.9)
Mexico	3.6 (2.9–4.3)	1.3 (1.0–1.7)	2.9 (2.4–3.5)	27.7 (24.4–31.3)
USA	10.3 (9.4–11.3)	7.2 (6.5–8.0)	5.1 (4.6–5.8)	44.2 (42.4–45.9)
Japan	1.7 (1.1–2.5)	1.4 (0.9–2.2)	1.7 (1.2–2.3)	43.1 (37.7–48.7)
Beijing	2.2 (1.5–3.2)	1.0 (0.6–1.6)	1.2 (0.7–2.2)	41.7 (37.2–46.3)
Shanghai	1.6 (0.8–3.1)	1.2 (0.4–3.4)	0.6 (0.2–1.4)	46.0 (40.8–51.3)
New Zealand	7.0 (6.4–7.7)	5.2 (4.6–5.7)	5.0 (4.5–5.5)	45.4 (43.6–47.3)
Belgium	4.9 (3.6–6.7)	2.6 (1.8–3.8)	2.9 (2.0–4.3)	46.1 (42.5–49.8)
France	7.2 (5.8–8.8)	3.6 (2.8–4.6)	3.3 (2.3–4.7)	50.6 (46.1–55.1)
Germany	2.9 (2.2–3.9)	1.6 (1.2–2.2)	2.2 (1.6–2.9)	41.1 (37.9–44.5)
Italy	3.4 (2.8–4.2)	1.7 (1.3–2.4)	1.6 (1.2–2.2)	47.5 (44.3–50.6)
Netherlands	5.1 (3.9–6.7)	3.8 (2.5–5.5)	2.9 (1.9–4.5)	39.4 (34.8–44.2)
Spain	3.1 (2.5–3.8)	1.5 (1.2–1.9)	2.3 (1.8–2.9)	41.3 (38.6–44.1)
Ukraine	11.9 (10.1–13.9)	7.9 (6.6–9.5)	1.9 (1.2–3.1)	54.0 (50.8–57.2)
Lebanon	2.2 (1.4–3.6)	0.6 (0.2–1.6)	1.8 (0.9–3.7)	30.3 (26.0–35.1)
Nigeria	0.7 (0.5–1.1)	0.3 (0.2–0.7)	0.8 (0.4–1.5)	33.0 (30.3–35.9)
Israel	5.1 (4.4–5.8)	3.5 (3.0–4.1)	2.6 (2.2–3.1)	41.7 (40.3–43.1)
South Africa	7.7 (6.8–8.7)	4.2 (3.5–4.9)	3.4 (2.6–4.5)	45.9 (43.2–48.5)
All countries	5.9 (5.6–6.2)	3.8 (3.6–4.0)	3.3 (3.1–3.5)	42.2 (41.6–42.9)

<sup>a</sup> 12-month DSM-IV depressive and/or anxiety disorders; physical disorders included stroke, heart attack, heart disease, asthma, chronic obstructive pulmonary disease (COPD), diabetes, ulcer, HIV/AIDS, epilepsy, tuberculosis and cancer; pain conditions included: arthritis, 12-month chronic back/neck problems, 12-month frequent or severe headaches and/or other 12-month chronic pain.

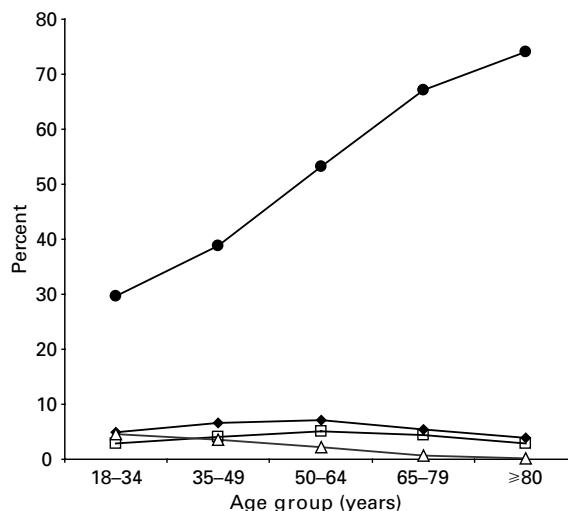
### Mental and/or physical condition groups by country

The prevalence of the four groups is shown in Table 2 for each of the contributing countries, all ages combined. The table illustrates two key features: there is a good deal of variability in prevalences across countries, but despite this variability, there is a pattern of mental disorder being more prevalent with pain or physical condition co-morbidity than without it. Physical/pain conditions unaccompanied by mental co-morbidity are a good deal more prevalent than when accompanied by mental co-morbidity, in all countries.

### Mental and/or physical condition groups by age

#### Prevalence by age

The age patterns in the prevalences of the four mental and/or physical condition groups are depicted in Fig. 1. The prevalence of depressive/anxiety disorders in the absence of physical/pain co-morbidity decreases with age. By contrast, the prevalence of physical/pain conditions in the absence of depressive/anxiety disorder increases sharply with



**Fig. 1.** Percentage with the four mental and/or physical condition combinations, by age group, all countries combined ( $n = 42\,697$ ). —●—, Depression/anxiety with chronic pain; —□—, depression/anxiety with physical and/or pain condition; —△—, depression/anxiety without physical condition; —●—, Physical and/or pain condition without depression/anxiety.

**Table 3.** Effect of age (reference level 18–34 years) in predicting the four mental and/or physical condition combinations, controlling for gender, all countries combined ( $n = 42\ 697$ )

Age group years ( $n$ )	Depressive/anxiety disorder with pain condition <sup>a</sup>	Depressive/anxiety disorder with physical condition <sup>a</sup>	Depressive/anxiety disorder without physical or pain condition <sup>a</sup>	Physical and/or pain condition without depressive/anxiety disorder <sup>a</sup>
18–34 (14 818)	1.0	1.0	1.0	1.0
35–49 (13 236)	1.3 (1.2–1.5)*	1.4 (1.3–1.6)*	0.8 (0.7–0.9)*	1.5 (1.4–1.6)*
50–64 (9006)	1.5 (1.3–1.6)*	1.9 (1.6–2.2)*	0.5 (0.4–0.5)*	2.7 (2.7–2.9)*
65–79 (4631)	1.0 (0.9–1.2)	1.6 (1.3–1.8)*	0.2 (0.1–0.2)*	4.8 (4.4–5.3)*
≥80 (959)	0.7 (0.5–1.0)*	0.9 (0.6–1.3)	0.0 (0.0–0.1)*	6.6 (5.5–8.0)*

Values are odds ratios (95% confidence intervals).

<sup>a</sup> 12-month DSM-IV depressive and/or anxiety disorders; physical disorders included stroke, heart attack, heart disease, asthma, chronic obstructive pulmonary disease (COPD), diabetes, ulcer, HIV/AIDS, epilepsy, tuberculosis and cancer; pain conditions included: arthritis, 12-month chronic back/neck problems, 12-month frequent or severe headaches and/or other 12-month chronic pain.

\*  $p < 0.05$ .

age. The two mental–physical co-morbidity groups display a similar age-related pattern with prevalence climbing slightly from younger to middle age and then reducing somewhat in the older groups.

#### Effects of age, controlling for gender

The effects of age on the four mental and/or physical condition groups after controlling for gender are provided in Table 3. These data clarify the age curves shown in Fig. 1, without the distraction of the differing prevalences across the four groups. The greatest change across the lifespan occurs in the decreasing odds of depressive/anxiety disorder without physical/pain co-morbidity. The two mental–physical co-morbid groups show slightly different patterns in that the odds of mental disorder co-morbid with pain are the same in the 65–79 years age group as the youngest group, but the odds of mental disorder co-morbid with physical condition are higher in the 65–79 years age group compared with the youngest group. However, all types of mental disorder (with or without co-morbidity) decline in the oldest age group. The lack of substantive difference between the mental disorder groups in age patterns is confirmed in finding that the interaction of age and physical disorder co-morbidity (present *versus* absent) in the association with 12-month depression/anxiety was not significant ( $p = 0.40$ ).

#### Proportion of mental disorder cases with physical/pain co-morbidity, by age

Mental disorders in the aggregate (i.e. both with and without physical/pain condition co-morbidity)

**Table 4.** Percentage with any 12-month depressive or anxiety disorder (cases), and proportion of cases with physical and/or pain condition co-morbidity, by age group, all countries combined ( $n = 42\ 697$ )

Age group years ( $n$ )	Percentage with any 12-month depressive or anxiety disorder (i.e. with or without physical/pain co-morbidity) <sup>a</sup> [% (95% CI)]	Proportion of cases (those with any 12-month depressive or anxiety disorder) with physical/pain co-morbidity <sup>a</sup> (%)
18–34 (14 818)	10.7 (10.2–11.3)	56.9
35–49 (13 236)	11.4 (10.8–12.0)	68.6
50–64 (9006)	10.4 (9.7–11.1)	78.9
65–79 (4631)	6.8 (6.1–7.5)	89.2
≥80 (959)	4.5 (3.5–5.9)	95.3

CI, Confidence interval.

<sup>a</sup> 12-month DSM-IV depressive and/or anxiety disorders; physical disorders included stroke, heart attack, heart disease, asthma, chronic obstructive pulmonary disease (COPD), diabetes, ulcer, HIV/AIDS, epilepsy, tuberculosis and cancer; pain conditions included: arthritis, 12-month chronic back/neck problems, 12-month frequent or severe headaches and/or other 12-month chronic pain.

decline in prevalence with age (Table 4). The proportion of cases with physical or pain condition co-morbidity is greater than 50% at all ages, increases in a monotonic fashion with age, and comprises the vast majority of the oldest age group with mental disorders (Table 4).

## Discussion

When mental disorders were disaggregated into those with and without physical/pain condition co-morbidity, those without co-morbidity decreased monotonically with age, whereas those with co-morbidity peaked in the middle years and then decreased in older age. However, no significant difference was found in the relationship between mental disorders and age as a function of physical/pain co-morbidity. In the aggregate, depressive and anxiety disorders decreased with age. The overlap of mental and physical conditions was asymmetrical; among those with mental disorders, physical/pain co-morbidity was more common than not, at all ages, and rises with age. By contrast, physical/pain conditions were more likely to occur without mental co-morbidity than with it, at all ages.

The results of this study are consistent with other general population surveys using standardized diagnostic measures in showing lower prevalences of depression and anxiety disorders in older age groups relative to younger age groups. One of the criticisms of this kind of study is that it samples from the non-institutionalized population and, effectively, given the demands of the CIDI, from a cognitively intact population. This is undeniable, but these sampling issues probably do not explain the decline in mental disorder prevalence with age. US data indicate that rates of institutionalization are fairly low among those aged 65–79 years, who form the majority of the older population (3% in those aged 65–69; 5% in those 70–74; 7% in those 75–79) (Siegler *et al.* 2002), and similar figures have been quoted for Australia (Troller *et al.* 2007). Troller *et al.* (2007) have shown (a) that inclusion of those living in aged-care facilities would have minimal impact on mental disorder prevalences among those aged  $\geq 65$  years, and (b) that prevalences are unaffected whether those with mild cognitive impairment are included or excluded from analysis.

It also seems improbable that the application of organic exclusion criteria can explain the decline in mental disorder prevalence with age. First, because the probes for organic causes were applied at the episode and not the symptom level, thus obviating the possibility that older persons would use organic attribution as a cognitive heuristic for dealing with symptom-related probes (Knauper & Wittchen, 1994). Although this does not obviate the possibility that older persons may be more likely to attribute entire episodes to organic causes (independent of actual health status), at present we are not aware of evidence that this occurs, and in each of the WMH surveys mental health clinicians judged the legitimacy of

all such attributions. Second, although the data we present on the rates of organic exclusions show a clear increase with age, the proportion of cases excluded for organic causes is not high enough to explain the decline in mental disorder prevalence with age. For example, if the excluded cases were all reincluded, estimates of mental disorder in the youngest and oldest age groups would rise to 11.2% and 5.0% respectively, compared with estimates of 10.7% and 4.5% with organic exclusions applied (and in fact this would be an overestimate of the impact of organic exclusions because the mental disorder estimates in this paper combine depressive and anxiety disorders). Heithoff (1995) showed a similar result with Epidemiologic Catchment Area (ECA) data.

These data do not, however, inform with regard to the possibility that depression may manifest differently with age, which might mean that the declining prevalence we observe is an artifact of the fact that DSM-IV criteria become less valid with increasing age. The research on this is not consistent. Gallo *et al.* (1994), for example, find a negative association between age and reports of anhedonia, whereas Christensen *et al.* (1999) find the opposite (see also Karel, 1997). A further consideration is the fact that these criticisms of diagnostic validity on the grounds of age differences in symptom manifestation and symptom attribution largely apply to depression diagnoses, yet anxiety disorders show the same decline with age (Alonso *et al.* 2004; Wells *et al.* 2006; Troller *et al.* 2007). Nevertheless, it is also clear that fully structured lay-conducted interviews such as the CIDI produce considerably lower estimates of DSM-based depressive disorders among older populations than semi-structured interviews conducted by clinicians (Skoog, 2004). The source of this discrepancy in prevalences, and its implications for conclusions about the validity of either the fully structured or semi-structured interviews, remains to be clarified.

In sum, this study demonstrates that despite the extremely prevalent reports of chronic pain and physical conditions among older age groups, the vast majority in the older general population do not have CIDI-diagnosed mental disorders. Methodological explanations notwithstanding, there are possible substantive explanations for this also, including age-related changes in expectations of declining health status (Karel, 1997; Siegler *et al.* 2002). Another possibility is that if, as some researchers have speculated, older persons have reduced psychological and social vulnerability to depression (Karel, 1997; Henderson *et al.* 1998; Blazer & Hybels, 2005), this may compensate for the increased prevalence of physical illness, or increase individuals' ability to cope with it. British researchers, observing a sharp decrease in prevalence

of depressive and anxiety disorders at retirement age (Villamil *et al.* 2006), have suggested that it may be explained by reducing societal demands and expectations associated with the statutory retirement age. This cannot explain all of the decrease in mental disorder prevalence observed in the current report, as it continues to reduce in the  $\geq 80$  years group relative to the 65–79 years age group (see also Troller *et al.* 2007), but different factors may be responsible for decreasing mental disorder prevalence at different ages, and retirement may well be an important factor in the younger old that helps to offset the psychological impact of increasing physical morbidity.

Considering the population with mental disorders, however, a different perspective emerges: physical condition co-morbidity is the rule, not the exception, regardless of age. Past research has emphasized the need for improved detection and treatment of mental disorders in primary care (Ormel *et al.* 1991; Katon *et al.* 1992; Coyne *et al.* 2002). The present study highlights the need for health professionals, including mental health professionals, to address barriers to adequately manage physical co-morbidity among those with mental disorders. Research among mental health professionals has identified barriers such as a lack of explicit allocation of responsibility for medical treatment, lack of service delivery integration, and pessimistic attitudes among treatment providers as to whether improved physical health is possible, or a priority, among those with mental illness (Brown *et al.* 2000; Friedli & Dardis, 2002; Hyland *et al.* 2003). Further research among primary-care providers and hospital physicians may be warranted to identify whether there are barriers (e.g. attitudinal, or time pressure) to a treatment focus on physical morbidity once a mental disorder has been diagnosed.

Several limitations of this study should be noted, in addition to the sampling limitations already discussed. First, physical and pain conditions were ascertained by a standard checklist, rather than physician's examination, which contrasts with the detailed assessment of mental disorders. While acknowledging the limitation of self-report, methods research indicates that self-report of diagnosis (which was the measure for most of the physical conditions) generally shows good agreement with medical records data (Kehoe *et al.* 1994; NCHS, 1994; Kriegsman *et al.* 1996). A second, related issue is that the relative prevalences of mental and physical conditions observed here, and their degree of overlap, are influenced by which disorders we have chosen to include/exclude. Not all mental disorders were included, although we included most of the common disorders and those most closely associated with physical co-morbidity. We also grouped together a larger number

of physical/pain conditions. While this needs to be borne in mind when interpreting the results, it seems likely that the kind of asymmetry we observe here (a lower proportion of the population with physical conditions having mental disorder co-morbidity relative to the proportion of the population with mental disorder who have physical co-morbidity) is characteristic of general populations. Third, we have not included any data on disability, so although on a pure frequency basis it appears as if physical morbidity outweighs mental morbidity in older populations, this does not take into account the complicating and disabling contribution that depression makes to the morbidity of medical conditions (Kessler *et al.* 2003b; Buist-Bouwman *et al.* 2005; Scott *et al.* in press).

In conclusion, this study provides a global, population perspective on the age patterning of CIDI-diagnosed depression and anxiety disorders, for the first time disaggregated into those with and without physical and pain co-morbidity. No significant difference was found in the relationship between mental disorders and age as a function of physical/pain co-morbidity. In the aggregate, depressive and anxiety disorders decreased with age, a result that cannot be explained by organic exclusion criteria. Physical/pain co-morbidity among those with mental disorders is normative and increases with age, which suggests that barriers to the adequate management of mental-physical co-morbidity remain.

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#### Declaration of Interest

None.

#### References

- Alonso J, Angermeyer MC, Bernert S, Bruffaerts R, Brugha TS, Bryson H, de Girolamo G, Graaf R, Demyttenaere K, Gasquet I, Haro JM, Katz SJ, Kessler RC, Kovess V, Lepine JP, Ormel J, Polidori G, Russo LJ, Vilagut G, Almansa J, Arbabzadeh-Bouchez S, Autonell J, Bernal M, Buist-Bouwman MA, Codony M, Domingo-Salvany A, Ferrer M, Joo SS, Martinez-Alonso M, Matschinger H, Mazzi F, Morgan Z, Morosini P, Palacin C, Romera B, Taub N, Vollebergh WA (2004). Prevalence of mental disorders in Europe: results from the European Study of the Epidemiology of Mental Disorders (ESEMeD) project. *Acta Psychiatrica Scandinavica Supplementum* **420**, 21–27.
- APA (1994). *DSM-IV: Diagnostic and Statistical Manual of Mental Disorders*, 4th edn. American Psychiatric Association: Washington, DC.
- Beekman ATF, Copeland JRM, Prince MJ (1999). Review of community prevalence of depression in later life. *British Journal of Psychiatry* **174**, 307–311.
- Beekman ATF, Deeg DJH, Van Tilburg T, Smit JH, Hooijer C, Van Tilburg W (1995). Major and minor depression in later life: a study of prevalence and risk factors. *Journal of Affective Disorders* **36**, 65–75.
- Blazer DG, Hybels CF (2005). Origins of depression in later life. *Psychological Medicine* **35**, 1241–1252.
- Brilman E, Ormel J (2001). Life events, difficulties and onset of depressive episodes in later life. *Psychological Medicine* **31**, 859–869.
- Brown S, Inskip H, Barraclough B (2000). Causes of the excess mortality of schizophrenia. *British Journal of Psychiatry* **177**, 212–217.
- Buist-Bouwman MA, de Graaf R, Vollebergh WAM, Ormel J (2005). Comorbidity of physical and mental disorders and the effect on work-loss days. *Acta Psychiatrica Scandinavica* **111**, 436–443.
- Christensen H, Jorm AF, Mackinnon AJ, Korten AE, Jacomb PA, Henderson AS, Rodgers B (1999). Age differences in depression and anxiety symptoms: a structural equation modelling analysis of data from a general population sample. *Psychological Medicine* **29**, 325–339.
- Coyne JC, Thompson R, Klinkman MS, Nease JR (2002). Emotional disorders in primary care. *Journal of Consulting and Clinical Psychology* **70**, 798–809.
- de Graaf R, Bijl RV, Ravelli A, Smit F, Vollebergh WAM (2002). Predictors of first incidence of DSM-III-R psychiatric disorders in the general population: findings from the Netherlands Mental Health Survey and Incidence Study. *Acta Psychiatrica Scandinavica* **106**, 303–313.
- Dew MA (1998). Psychiatric disorder in the context of physical illness. In *Adversity, Stress and Psychopathology* (ed. B. P. Dohrenwend), pp. 177–218. Oxford University Press: New York.
- Evans DL, Charney DS, Lewis L, Golden JM, Ranga Rama Krishnan K, Nemeroff CB, Bremner JD, Carney RM, Coyne JC, Delong MR, Frasurre-Smith N, Glassman AH, Gold PW, Grant I, Gwyther L, Ironson G, Johnson RL, Kanner AM, Katon WJ, Kaufmann PG, Keefe FJ, Ketter T,

- Laughren TP, Leserman J, Lyketsos CG, McDonald WM, McEwan BS, Miller AH, Musselman D, O'Connor C, Petitto JM, Pollock BG, Robinson RG, Roose SP, Rowland J, Sheline Y, Sheps DS, Simon G, Spiegel D, Stunkard A, Sunderland T, Tibbits P, Valvo WJ (2005). Mood disorders in the medically ill: scientific review and recommendations. *Biological Psychiatry* **58**, 175–189.
- Friedli L, Dardis C (2002). Not all in the mind: mental health service user perspectives on physical health. *Journal of Mental Health Promotion* **1**, 36–46.
- Gallo JJ, Anthony JC, Muthén BO (1994). Age differences in the symptoms of depression: a latent trait analysis. *Journal of Gerontology* **49**, 251–264.
- Harter MC, Conway KP, Merikangas KR (2003). Associations between anxiety disorders and physical illness. *European Archives of Psychiatry and Clinical Neuroscience* **253**, 313–320.
- Heithoff K (1995). Does the ECA underestimate the prevalence of late-life depression? *Journal of the American Geriatrics Society* **43**, 2–6.
- Henderson AS, Jorm AF, Korten A, Jacomb P, Christensen H, Rodgers B (1998). Symptoms of depression and anxiety during adult life: evidence of a decline in prevalence with age. *Psychological Medicine* **28**, 1321–1328.
- Hybels CF, Blazer D (2002). Epidemiology and geriatric psychiatry. In *Textbook in Psychiatric Epidemiology* (ed. M. T. Tsuang and M. Tohen), pp. 603–628. Wiley-Liss Inc.: New York.
- Hyland B, Judd F, Davidson S, Jolley D, Hocking B (2003). Case managers' attitudes to the physical health of their patients. *Australian and New Zealand Journal of Psychiatry* **37**, 710–714.
- Jorm AF (2000). Does old age reduce the risk of anxiety and depression? A review of epidemiological studies across the adult life span. *Psychological Medicine* **30**, 11–22.
- Karel MJ (1997). Aging and depression: vulnerability and stress across adulthood. *Clinical Psychology Review* **17**, 847–879.
- Katon W, Von Korff M, Lin E, Bush T, Ormel J (1992). Adequacy and duration of antidepressant treatment in primary care. *Medical Care* **30**, 67–76.
- Kehoe R, Wu S-Y, Leske MC, Chylack LT (1994). Comparing self-reported and physician reported medical history. *American Journal of Epidemiology* **139**, 813–818.
- Kessler R, Berglund P, Demler O, Jin R, Koretz D, Merikangas KR, Rush AJ, Walters EE, Wang PS (2003a). The epidemiology of major depressive disorder. Results from the National Comorbidity Survey Replication (NCS-R). *Journal of the American Medical Association* **289**, 3095–3105.
- Kessler R, Foster C, Webster PS, House JS (1992). The relationship between age and depressive symptoms in two national surveys. *Psychology and Aging* **7**, 119–126.
- Kessler RC, Berglund P, Chiu WT, Demler O, Heeringa S, Hiripi E, Jin R, Pennell B-E, Walters EE, Zaslavsky A, Zheung H (2004). The US National Comorbidity Survey Replication (NCS-R): design and field procedures. *International Journal of Methods in Psychiatric Research* **13**, 69–92.
- Kessler RC, Ormel J, Demler O, Stang PE (2003b). Comorbid mental disorders account for the role impairment of commonly occurring chronic physical disorders: results from the National Comorbidity Survey. *Journal of Occupational and Environmental Medicine* **45**, 1257–1266.
- Kessler RC, Ustun B (2004). The World Mental Health (WMH) Survey Initiative version of the World Health Organization (WHO) Composite International Diagnostic Interview (CIDI). *International Journal of Methods in Psychiatric Research* **13**, 93–121.
- Knäuper B, Wittchen H-U (1994). Diagnosing major depression in the elderly: evidence for response bias in standardized diagnostic interviews? *Journal of Psychiatric Research* **28**, 147–164.
- Kriegsman DM, Penninx BW, Van Eijk JT, Boeke AJ, Deeg DJ (1996). Self-reports and general practitioner information on the presence of chronic diseases in community dwelling elderly. *Journal of Clinical Epidemiology* **49**, 1407–1417.
- Krishnan KRR (2002). Biological risk factors in late life depression. *Biological Psychiatry* **52**, 185–192.
- Mulsant BH, Ganguli M (1999). Epidemiology and diagnosis of depression in late life. *Journal of Clinical Psychiatry* **60** (Suppl. 20), 9–15.
- NCHS (1994). Evaluation of National Health Interview Survey diagnostic reporting. *Vital and Health Statistics, Series 2* **120**, 1–116.
- Newmann JP (1989). Aging and depression. *Psychology and Aging* **4**, 150–165.
- O'Connor DW (2006). Do older Australians truly have low rates of anxiety and depression? A critique of the 1997 National Survey of Mental Health and Wellbeing. *Australian and New Zealand Journal of Psychiatry* **40**, 623–631.
- Ormel J, Koeter M, van den Brink W, van de Willige G (1991). Recognition, management and course of anxiety and depression in general practice. *Archives of General Psychiatry* **48**, 700–706.
- Pirkola SP, Isometsä E, Suvisaari J, Aro H, Joukamaa M, Poikolainen K, Koskinen S, Aromaa A, Lönnqvist JK (2005). DSM-IV mood, anxiety and alcohol use disorders and their comorbidity in the Finnish general population. Results from the Health 2000 study. *Social Psychiatry and Psychiatric Epidemiology* **40**, 1–10.
- Regier DA, Boyd JH, Burke JDJ, Rae DS, Myers JK, Kramer M, Robins LN, George LK, Karno M, Locke BZ (1988). One-month prevalence of mental disorders in the United States. *Archives of General Psychiatry* **45**, 977–986.
- Schoevers RA, Beekman ATF, Deeg DJ, Geerlings MI, Jonker C, Van Tilburg W (2000). Risk factors for depression in later life; results of a prospective community based study (AMSTEL). *Journal of Affective Disorders* **59**, 127–137.
- Scott KM, Bruffaerts R, Tsang A, Ormel J, Alonso J, Angermeyer MC, Benjet C, Bromet E, de Girolamo G, de Graaf R, Gasquet I, Gureye O, Haro JM, He Y, Kessler RC, Levinson D, Mneimneh ZN, Oakley Browne MA, Posada-Villa J, Stein DJ, Takeshima T, Von Korff M

- (2007). Depression-anxiety relationships with chronic physical conditions: results from the World Mental Health surveys. *Journal of Affective Disorders* **103**, 113–120.
- Scott KM, Von Korff M, Alonso J, Angermeyer MC, Bromet E, Fayyad J, de Girolamo G, Demyttenaere K, Gasquet I, Gureje O, Haro JM, He Y, Kessler RC, Levinson D, Medina Mora ME, Oakley Browne MA, Ormel J, Posada-Villa J, Watanabe M, Williams D** (in press). Mental–physical comorbidity and its relationship with disability: results from the World Mental Health surveys. *Psychological Medicine*.
- Siegler IC, Bastian LA, Steffens DC, Bosworth HB, Costa PT** (2002). Behavioural medicine and aging. *Journal of Consulting and Clinical Psychology* **70**, 843–851.
- Skoog I** (2004). Psychiatric epidemiology of old age: the H70 study – the NAPE lecture 2003. *Acta Psychiatrica Scandinavica* **109**, 4–18.
- Snowdon J** (2001). Is depression more prevalent in old age? *Australian and New Zealand Journal of Psychiatry* **35**, 782–787.
- Stordal E, Mykletun A, Dahl AA** (2003). The association between age and depression in the general population: a multivariate examination. *Acta Psychiatrica Scandinavica* **107**, 132–141.
- SUDAAN** (2002). SUDAAN Inc., Research Triangle Institute, Research Triangle Park: NC.
- Troller JN, Anderson TM, Sachdev PS, Brodaty H, Andrews G** (2007). Age shall not weary them: mental health in the middle-aged and the elderly. *Australian and New Zealand Journal of Psychiatry* **41**, 581–589.
- van't Veer-Tazelaar PJ, van Marwijk HWJ, Jansen APD, Rijmen F, Kostense PJ, van Oppen P, van Hout HPJ, Stalman WAB, Beekman ATF** (2008). Depression in old age (75+), the PIKO study. *Journal of Affective Disorders* **106**, 295–299.
- Villamil E, Huppert FA, Melzer D** (2006). Low prevalence of depression and anxiety is linked to statutory retirement ages rather than personal work exit: a national survey. *Psychological Medicine* **36**, 999–1009.
- Wells JE, Oakley Browne MA, Scott KM, McGee MA, Baxter J, Kokaua J** (2006). Prevalence, interference with life and severity of DSM-IV disorders in Te Rau Hinengaro: The New Zealand Mental Health Survey. *Australian and New Zealand Journal of Psychiatry* **40**, 845–854.
- Wells KB, Golding JM, Burnam MA** (1989). Affective, substance use, and anxiety disorders in persons with arthritis, diabetes, heart disease, high blood pressure, or chronic lung conditions. *General Hospital Psychiatry* **11**, 320–327.