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Time-dependent cognitive and somatic symptoms of depression as predictors of new cardiac-related events in at-risk patients: the UPBEAT-UK cohort

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Abstract

Background: evidence suggests that somatic rather than cognitive depressive symptoms are risk factors for recurrent cardiac events in at-risk patients. However, this has never been explored using a time-dependent approach in a narrow time-frame, allowing for an cardiac event-free time-window.

Methods: the analysis was performed on 595 participants (70.6% male, median age 72 (27-98)) drawn from the UPBEAT-UK heart disease patient cohort with 6-monthly follow-ups over 3 years. Depressive symptomatology was measured using the Patient Health Questionnaire-9 (PHQ-9) (four somatic, five cognitive items). New cardiac events (NCE) including cardiac-related mortality were identified by expert examination of patient records. Analyses were performed using Cox Proportional Hazard models with delayed entry, with time-dependent depressive dimensions and covariates measured 12-to-18 months (median: 14.1, IQR: 3.5) prior to the event, with a 12-month cardiac event-free gap.

Results: there were 95 NCE during the follow-up (median time-to-event from baseline: 22.3 months (IQR: 13.4)). Both the somatic (HR: 1.12, 95% CI.: 1.05-1.20, p=0.001) and cognitive dimensions (HR: 1.11, 95% CI.: 1.03-1.18, p=0.004) were time-dependent risk factors for a NCE in the multi-adjusted models. Specific symptoms (poor appetite/overeating for the somatic dimension, hopelessness and feeling like a failure for the cognitive dimension) were also significantly associated.

Conclusion: This is the first study of the association between depressive symptom dimensions and NCEs in at-risk patients using a time-to-event standardized approach. Both dimensions considered apart were independent predictors of a NCE, along with specific items, suggesting regular assessments and tailored interventions targeting specific depressive symptoms may help to prevent new cardiac events in at-risk populations.

Introduction

There is extensive research on the bi-directional relationship between depression and heart disease and the possible mechanisms involved (Ormel and de Jonge, 2011), and it is now widely acknowledged that depression is a risk factor for recurrent cardiac events and cardiac-related deaths in patients with heart disease (Freedland and Carney, 2013, Meijer *et al.*, 2011).

Regarding specific depressive symptoms, a recent meta-analysis suggested that the somatic rather than the cognitive symptom dimension of depression was predictive of recurrent cardiovascular events in at-risk patients (de Miranda Azevedo *et al.*, 2014). For instance, in a study of 2347 acute myocardial infarction (MI) patients, the Patient Health Questionnaire-9 (PHQ-9) somatic symptom dimension only was independently associated with one-year rehospitalisation and to a lesser extent 4-year mortality (Smolderen *et al.*, 2009). In keeping with this, in a cohort of 1019 stable coronary heart disease (CHD) patients, a 14% greater risk of cardiovascular events over 6 years was reported for the PHQ-9 somatic dimension and, more specifically, for fatigue and appetite problems (Hoen *et al.*, 2010b). Several other studies also found somatic symptoms only to be associated with cardiac health status, cardiac prognosis and mortality in MI patients (de Jonge *et al.*, 2006, Martens *et al.*, 2010, Roest *et al.*, 2011). In contrast, some studies identified as potential risk factors for a recurrent event either both dimensions (Denollet *et al.*, 2013, Hoen *et al.*, 2010a) or the cognitive dimension only when depressive symptoms were evaluated postoperatively in coronary bypass patients (Connerney *et al.*, 2010, Tully *et al.*, 2011). However, the diversity of assessment scales and depressive symptom classifications makes study comparison difficult (Carney and Freedland, 2012b).

There are various explanations for the widespread evidence in favour of somatic symptoms as risk factors for cardiac events. Firstly, somatic symptoms tend to be more common in depressed patients with heart disease compared to other depressed patients (Martens *et al.*, 2006). They are more likely to overlap with cardiac disease, not only because some symptoms such as fatigue are shared by both conditions, but also because depressed at-risk patients may focus more on their somatic symptoms which they can attribute to their underlying heart condition (Carney and Freedland, 2012a). Secondly, somatic depressive symptoms are likely to be overestimated by self-report questionnaires on which findings almost exclusively rely. This is because no weighing of symptoms is performed as with structured diagnostic interviews. In the only study assessing depression using a diagnostic interview, both somatic and cognitive symptoms were predictive of cardiovascular prognosis in the multi-adjusted analysis (Hoen *et al.*, 2010a). Thirdly, studies performed so far have not addressed the time-frame of the exposure-event matrix. In most studies depressive symptomatology was assessed at a fixed time-point with considerable variability in duration of time between assessment and cardiac onset (Connerney *et al.*, 2010, Hoen *et al.*, 2010b). The absence of a cardiac event-free time-window between depression assessment and outcome could lead to a greater reporting of somatic symptoms, as markers of cardiac disease severity (Meijer *et al.*, 2013).

Depressive symptoms have been shown to fluctuate in the elderly, especially over the long-term, and persistent high or increasing symptom trajectories have been linked to adverse health outcomes, such as dementia (Kaup *et al.*, 2016) and mortality (Andreescu *et al.*, 2008). However, irrespective of symptom change, symptom levels per se can exhibit time-varying effects; this has been shown for example for all-cause and non-cardiovascular disease-specific mortality over a 7-year follow-up (Moise *et al.*, 2018). A time-dependent

approach allows the assessment of risk within a defined time-window, standardising length of time between exposure assessment and the outcome. The potential time-varying effect of the exposure is thereby taken into account. Studies analysing depression as a time-dependent risk factor are scarce (Li *et al.*, 2018, Pequignot *et al.*, 2016). Over a 12-year follow-up, Li *et al.* found that contrary to baseline depressive symptoms, time-dependent symptoms increased the risk of all-cause and cardiovascular mortality (Li *et al.*, 2018). However, no studies so far have examined the different symptom dimensions as recurrent cardiac event risk factors using this approach.

The aim of the current study was to examine PHQ-9 somatic and cognitive symptom dimensions, measured 12 to 18 months prior to the event, as short-term predictors of cardiac-related events and mortality, drawing on data from the UPBEAT-UK cohort of heart disease patients followed up at 6-monthly intervals over 3 years. Individual symptoms and symptom change were also investigated.

Methods

Study design and sample

This study draws on data from the UPBEAT-UK cohort. Participants were identified and recruited from Coronary Heart Disease (CHD) Registers kept by the 16 participating general practices in South East London (Tylee *et al.*, 2011). All register patients (n=2938) were invited to participate by their local GP; 917 agreed to be contacted, of which 803 (87.6%) participated. Each participant had an initial baseline interview and was followed up for three years, undergoing assessments every six months. Recruitment and assessments ran from 2008 to 2011. Written informed consent was obtained from all participants before the initial

assessment. Ethics approval was granted through the Bexley and Greenwich Research Ethics Committee (REC reference number: 07/H0809/38).

The present analysis was performed on 595 patients. From the initial 803 participants, we excluded 83 with an NCE recorded at cohort inclusion or at the 6 or 12 month follow-ups; a further 71 with outcome data missing at all four follow-ups (18, 24, 30 and 36 months), a further 25 with a delay between CHD register entry (corresponding approximately to the first major cardiac event) and recruitment into the cohort less or equal to 6 months, and a further 29 participants with missing data for any of the main covariates.

Measures

Socio-demographic and lifestyle characteristics, including relationship status, smoking, alcohol consumption and body mass index (BMI), were collected at baseline only. The Clinical Interview Schedule Revised (CIS-R) (Lewis *et al.*, 1992) was administered to establish International Classification of Diseases-10 (ICD-10) diagnoses of depression at baseline. Each participant was attributed an Index of Multiple Deprivation (IMD) score according to their postcode (National Statistics, 30 September 2015).

At baseline and each follow-up, severity of chest pain was measured using the Rose Angina Questionnaire (Rose, 1962) and was classified as follows: no angina; chest pain when walking uphill only; chest pain when walking on the level and uphill. Anxiety scores were derived from the anxiety module of the Hospital Anxiety and Depression Scale (HADS-A) (Zigmond and Snaith, 1983). Time-dependent quality of life scores were obtained using the EuroQol EQ-5D scale, excluding the pain/discomfort and depression/anxiety subscales (Rabin and de Charro, 2001), and social problems from the Social Problems Questionnaire (SPQ) (Corney and Clare, 1985).

Outcome variable: new cardiac event (NCE).

This was a composite of any cardiac event encompassing: any visit to the rapid access chest pain clinic, A&E (Accident and Emergencies) or emergency hospital admittance, where cardiac-related chest pain was the diagnosis; bypass graft or angioplasty; MI and any cardiovascular cause of death. Events were coded and dated by a medical doctor on the team after examination of GP notes. Mortality data were gathered from case notes and other publically available sources. Cause and date of death were obtained from the practice manager at the different GP surgeries. For the purpose of the analysis, we refer to the first cardiac event noted at the 18, 24, 30 and 36 month follow-ups, which occurred any time during the six months preceding the follow-up.

Exposure variable: somatic and cognitive depression dimensions.

These were derived from the PHQ-9, a screener measuring nine depressive symptoms over the past two weeks, each scored from 0 (not at all) to 3 (nearly every day) (Kroenke *et al.*, 2001). Four questions refer to the somatic dimension (trouble with sleep, feeling tired, poor appetite/overeating, slowness/restlessness) with scores ranging from 0 to 12. Five questions refer to the cognitive dimension (little interest in doing things, feeling depressed/hopeless, like a failure, trouble concentrating, thoughts of death), ranging from 0 to 15. Scores were obtained by summing the individual items for each dimension.

Statistical analysis

All statistical tests were performed using Cox proportional hazard models with delayed entry corresponding to age at baseline, with age as the basic time scale and birth as the time

origin. This method enables a better adjustment for age and is preferred, especially for elderly samples, over the standard model with time since inclusion as the time scale (Lamarca *et al.*, 1998). The assumptions of proportional hazards over time for baseline variables and the linearity of all continuous variables were verified. Results are expressed as hazard ratios (HR) with 95% confidence intervals (CI). For the depression dimensions, HRs represent the increased risk per one-point increase in score on an ordinal scale of 0-12 and 0-15, for the somatic and cognitive dimensions respectively. For each individual symptom, this corresponds to a one-point increase on the 0-3 scale.

Somatic and cognitive dimensions, along with individual symptom items, were analysed separately both as baseline and time-dependent variables. We excluded participants with a NCE at the baseline, 6- and 12-month follow-ups. This ensured a gap of at least 12-months between baseline depression assessment and date of outcome. In the time-dependent analysis, depressive symptom scores and available covariates were taken at the N-3 follow-up, allowing for this 12-month NCE-free gap. The median time between time-dependent measures and event was 14.1 months (Inter-quartile range (IQR): 3.5). Covariates measured at follow-up were: chronic diseases, chest pain, social problems, quality of life and anxiety. All other variables were available at baseline only. For the multivariate models, we selected all baseline or time-dependent covariates associated with NCE with p-values ≤ 0.25 or considered as well-established risk factors in the literature (see Supplementary Table 1). Interactions with sex and chest pain were tested and not significant. Bonferroni correction was used to account for multiple comparisons when considering the individual symptoms on each depression dimension ($p < 0.0125$ and $p < 0.01$ for the somatic and cognitive dimensions, respectively).

In addition, we studied time-dependent change in somatic and cognitive dimension scores since the previous follow-up using the same risk factor to event-time framework. For each subject, we calculated the relative change in score (score at time 't' minus score at time 't-1' divided by score at 't-1') on each dimension. We then compared the two extreme categories of the quartiles of the score distribution, with a greater than 25% decrease (i) or increase in score (ii), to the middle category representing the interquartile range (iii). Significance was set at $p < 0.05$. Statistical analyses were performed using SAS Enterprise Guide Version 7.15 (SAS Institute, Inc. Cary, North Carolina).

Results

Sample description

The characteristics of the cohort have been described in detail elsewhere (Walters *et al.*, 2014). We compared the study sample of 595 participants to the 208 participants excluded from the analysis. There were no differences for the main socio-demographic, lifestyle and health variables except for quality of life ($p=0.01$), anxiety ($p=0.006$) and the cognitive depression dimension ($p=0.0006$), with higher severity scores for those excluded from the analysis.

The study sample was mainly male (70.6%), with a median age of 72 years, ranging from 27 to 98. Chest pain (Grade 1 or 2 angina) was reported at baseline by 24.1% of the sample. Depression was diagnosed in 5.9% of participants and 16.8% had a PHQ-9 score >10 . The median (min-max) scores on the somatic and cognitive dimensions were 2 (0-12) and 0 (0-12), respectively (Table 1).

Of the sample, 42.4% entered the CHD register with a diagnosis of MI, 48.2% with ischemic heart disease, 6.9% with angina, and 2.5% with one of the following indications: heart

failure, chest pain not otherwise specified, arrhythmias or other vascular problems. Median time between CHD register entry and baseline was 8.3 years, ranging from 0.6 (by design, as participants with a delay between CHD register entry and cohort entry less or equal to 6 months were excluded) to 42.8 years. Over the four follow-ups where outcomes were considered in the analysis, there were 95 NCE, 31 at the 18-month, 23 at the 24-month, 12 at the 30-month and 29 at the 36-month follow-up. Median time from baseline to event was 22.3 months (IQR: 13.4, min-max: 12.1-36.2). The most frequent event was a visit to the rapid access chest pain clinic, A&E or emergency hospital admittance (47.4%), followed by cardio-cerebrovascular cause of death (26.3%), bypass graft or angioplasty (24.2%), and MI (2.1%).

Somatic and cognitive symptom dimensions as NCE risk factors

In the multivariate models, both the somatic and cognitive dimensions were significantly associated with a NCE whether measured as time-dependent variables or at baseline (Table 2). All analyses became non-significant when further adjusting for quality of life and anxiety, except for the somatic dimension at baseline ($p=0.04$) (not shown).

Individual symptom items as NCE risk factors

Combining the baseline and time-dependent approaches, all individual somatic symptoms were significantly associated with the onset of a NCE in the multivariate models (Table 3). However, when applying Bonferroni correction, only poor appetite/overeating remained significant. Of the cognitive items, feeling down at baseline and feeling like a failure for the time-dependent approach were significant multiple-comparison corrected predictors.

Time-dependent change in dimension scores as NCE risk factors

For both dimensions, variability in scores over follow-up were low and not significant (F-value=0.32, p=0.93 for the somatic dimension; F-value=1.34, p=0.24 for the cognitive dimension). Approximately one third of participants for the somatic dimension and half for the cognitive dimension presented no change in score between two-successive follow-ups (Figures 1 and 2). Time-dependent change in somatic and cognitive scores were not associated with the risk of a NCE.

Discussion

To the best of our knowledge, this is the first study to examine somatic and cognitive dimensions of depression as time-dependent risk factors for recurrent cardiac events. Conversely to findings from elsewhere, our findings suggest that both dimensions could be important short-term predictors of recurrent cardiac events when measured 12 to 18 months prior to the event, in particular somatic symptom appetite/overeating and cognitive symptoms feeling low and feeling like a failure.

Cognitive and somatic symptom dimensions

Our findings from the baseline and time-dependent analysis suggest both the somatic and cognitive depression dimensions to be predictors of a NCE, whereas most other studies find an effect for the somatic dimension only (de Miranda Azevedo *et al.*, 2014). We adjusted, as elsewhere (Hoen *et al.*, 2010b, Linke *et al.*, 2009, Martens *et al.*, 2010) for history of heart disease taking diagnosis at entry into the CHD register as a proxy, along with the main chronic diseases, smoking and chest pain. However, the associations became largely non-significant when further adjusting for quality of life and anxiety. Despite removing the

pain/discomfort and anxiety/depression scales from the EQ-5D, they could be over-adjusted given the strong associations between chronic diseases and quality of life and the overlap in symptoms between subthreshold depression and anxiety (Braam *et al.*, 2014). In fact, to our knowledge, no other study adjusted for these covariates in their multivariate models.

The findings for the time-dependent analysis are very similar to those using the baseline data. This is not actually surprising given that both analyses examined depressive symptoms as short-term risk factors for a NCE. The main difference between the two approaches is the variability in time-to-event (median (IQR): 22.3 (13.4) in the baseline analysis, and 14.1 (3.5) for the time-dependent analysis). Furthermore, we allowed for an event-free time-window between exposure and event in both approaches, in order to avoid the concomitant onset of the two, with depressive symptoms acting more like a cardiac disease marker than a short-term risk factor.

The association between time-dependent depression and cardiac prognosis has seldom been investigated (Li *et al.*, 2018, Pequignot *et al.*, 2016). In a Chinese cohort of 1999 subjects aged 55 years and above, the time-dependent depressive symptom score increased the risk of all-cause and cardiovascular mortality over a 12-year follow-up, whereas no association was found with symptom score at baseline (Li *et al.*, 2018). The difference could be explained by the longer follow-up period than in our study, with a greater difference in inter-subject variability in time-to-event. In the Three-City Study including 9294 persons aged 65 years and above followed up for over 10 years, time-dependent number of visits with a high level of depressive symptoms was associated with cardiovascular disease mortality risk, which increased 1.28-fold for each additional visit (Pequignot *et al.*, 2016). However, neither

of these studies using the Centre for Epidemiologic Studies-Depression Scale considered somatic and cognitive symptom dimensions separately.

Many behavioural and biological mechanisms have been suggested to explain the link between depressive symptomatology and recurrent cardiac events. Inflammation is thought to play a key role, in particular in somatic and late-onset depressive symptoms (Meijer *et al.*, 2013). Ormel and de Jonge suggested a differential effect on acute coronary syndromes in CHD patients, depending on the type of symptoms (Ormel and de Jonge, 2011). Their integrative model rests on three core hypotheses. The first is that depression in CHD patients consists of a mix of two types of depression, cognitive/affective and somatic depression, each having a characteristic symptom expression and aetiology. The second is that the effects on CHD depend on the type and duration of depression. The third hypothesis is that the most plausible pathways mediating these effects are behavioural. They argue that for both cognitive-affective and somatic depressive dimensions, persistent (chronic or recurrent) symptoms would be expected to be associated with recurrent cardiac events, but not brief symptom episodes (remission within 6 months) that are likely to be confounded by the severity of underlying cardiac symptoms and associated physiological abnormalities. Moreover, they suggest a different pathway for each symptom dimension, with behavioural factors alone mediating the cognitive pathway, whereas both behavioural and other factors such as inflammation would mediate the somatic symptom pathway.

Greater fluctuations over time of behavioural factors (e.g. lifestyle, medication adherence) compared to biological factors could explain why studies considering depressive symptoms at a fixed time-point find no association for the cognitive dimension. The latter present considerable variability, both between and within studies, in duration of time between baseline depression assessment and outcome. In the present study, median time to event

was relatively short (22.3 months (IQR: 13.7)). Other studies report average times to event ranging from 2.5 years (sd=0.9) (de Jonge *et al.*, 2006) to 6.1 years (sd=2.0) (Hoen *et al.*, 2010b).

Of the few studies reporting an association for the cognitive dimension measured at a fixed time-point, Hoen *et al.* reported that both the somatic and cognitive dimensions were associated with cardiovascular prognosis in a study of 468 MI patients with a relatively short mean follow-up of 2.5 years (sd=0.8), (Hoen *et al.*, 2010a). The authors attributed this to a more thorough assessment of symptoms thanks to the use of a structured diagnostic interview. This suggests that an overestimation of somatic symptoms with self-report instruments coupled with longer and varying times-to-event could account for previous negative findings for the cognitive dimension. Another study showed a link between the cognitive-affective dimension of the BDI administered at baseline and recurrent MI and mortality in 1823 patients with a mean follow-up of 2.1 years (sd=1.1); this was found for the under 70 age-group only in a sample of patients with post-MI depression (Denollet *et al.*, 2013). This highlights the complexity of the relationship between NCE and depressive symptoms that could be modulated by both an age and depression level threshold effect.

Time-dependent symptom change over time

We also examined for each dimension, the association between time-dependent symptom change and the risk of a NCE. Neither of the associations were significant. Of the few studies investigating change over time (Roest *et al.*, 2013, Scafato *et al.*, 2012), only Roest *et al.* with the ENRICHD study examined separate symptom dimensions. They reported that over 6 months improvement in somatic symptoms but not cognitive symptoms was associated with a reduced risk of recurrent MI and mortality; however, change was not considered as a time-

dependent variable allowing for considerable variability in time-to-event over the 2.4 year (standard deviation (sd)=2.9) follow-up period (Roest *et al.*, 2013).

We chose to express change in terms of the relative change in score but this may have biased our findings as nearly half of subjects were in the 'no change' category, in particular for the cognitive dimension due to a clustering of scores at 0 for approximately 50% of participants at each follow-up. Furthermore, the 'no change' category included the middle quartiles with both persistently high and low scores. This low level of change has previously been reported for this cohort (Palacios *et al.*, 2016). A more sophisticated approach to change taking into account trajectories over the entire follow-up (Palacios *et al.*, 2018), was not possible given the time-dependent approach used in the analysis.

Individual depressive symptoms

To our knowledge, this is the first time individual depressive symptom items are examined as time-dependent variables. Of the somatic symptoms, poor appetite/overeating was the most robust predictor of a NCE. Hoen *et al.* in a study of 1019 patients with stable CHD over a relatively long follow-up period (mean 6.1 years (SD=2.0)), found that only two of PHQ-9 symptoms – namely somatic items appetite problems and fatigue – measured as continuous variables at a fixed time-point were associated with recurrent cardiovascular events (Hoen *et al.*, 2010b). When using a structured diagnostic interview, the same authors reported that all baseline somatic symptoms except for sleeping difficulties were predictors of cardiovascular prognosis, along with the cognitive symptoms lack of interest and suicidal ideation (Hoen *et al.*, 2010a). Denollet *et al.* (2013) examined individual cognitive symptoms and found that only hopelessness remained significantly associated with recurrent MI and mortality

(Denollet *et al.*, 2013). This is in keeping with our findings regarding cognitive items feeling down and feeling like a failure.

Limitations

There are several limitations to our study. Firstly, we used a self-report instrument to measure depressive symptoms which may have led to an overestimation of somatic symptoms (Hoen *et al.*, 2010a). On the other hand, the PHQ-9 has been suggested as a useful screening tool in cardiac patients, as it covers a wide range of symptoms with a clear and validated distinction between the somatic and cognitive dimensions (Carney and Freedland, 2012a, de Jonge *et al.*, 2007). It has also been shown to have a superior diagnostic accuracy to the HADS-D in CHD patients (Haddad *et al.*, 2013). Secondly, although we were able to adjust for a wide range of variables, we did not have data on medication use and could not adjust for antidepressant or cardiac-related medication and treatment compliance. A further limitation is the heterogeneity of cardiac events both at entry into the CHD register and as outcome variable (Freedland and Carney, 2013). Indeed NCE included events ranging from rapid access chest pain clinic for cardiac-related chest pain to cardio-cerebrovascular cause of death. The exclusion of events at the first three time-points and the limited number of events per category of NCE did not allow us to investigate each type of event separately. Finally, we were not able to show any effect of change in scores between follow-ups, mainly because symptom scores fluctuated very little over the follow-up. This in turn can be explained by the clustering of scores at the lower end of the distribution due to the selection of a healthy sub-sample, as well as the short duration of follow-up (36 months).

Strengths

The main strength of our study is the design regarding the delay between depression assessment and the NCE. The time-dependent approach allowed us to standardise for time-to-event, which is crucial when examining symptoms that are known to fluctuate considerably over time. Furthermore, in order to distinguish 'depression' as a short-term risk factor for a NCE from 'depression as a non causal variable risk marker' (Meijer *et al.*, 2013) we chose to leave an event-free period of 12-months between symptom assessment and the cardiac onset period. This reduced the likelihood of over-reporting somatic symptoms as a marker of cardiac disease.

We also removed patients who entered the register in the six months preceding baseline, in an attempt to measure depressive symptomatology as a risk factor for a NCE rather than the opposite, although we cannot fully rule out that baseline symptoms could be the consequence of past cardiac events or worsening of symptoms. Finally, our study was based on a relatively large sample of patients from CHD registers with regular 6-monthly follow-ups over 3 years, with full datasets. NCE cases were certified by an expert clinical review of all patient records with exact dates for events.

Conclusion

We showed that both the somatic and cognitive depressive symptom dimensions measured 12 to 18 months earlier were significantly and positively associated with a greater risk of a NCE. Thus, cognitive symptoms should not be disregarded as short-term risk factors for recurrent cardiac events as suggested by most previous research. More regular assessment of depressive symptoms, with a special attention given to specific symptoms such as eating

problems and feeling hopeless or like a failure, could help predict and prevent future cardiac events.

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Conflict of interest

None

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Table 1. Sample description at baseline (N=595)

Variable	N	%
Age at cohort entry*	595	72 [27-98]
Sex (Male)	420	70.6
Education (> 10 years)	309	51.9
Ethnicity (White)	530	89.1
Living alone (Yes)	211	35.5
Smoking status		
Never	178	29.9
Ex	330	55.5
Current	87	14.6
Alcohol (units per week)		
Doesn't drink	160	26.9
0-10 units	286	48.1
>10 units	149	25.0
Body mass index		
Normal	133	22.4
Overweight	272	45.7
Obese	190	31.9
Chronic diseases:		
Diabetes	147	24.7
Hypertension	326	54.8
Arthritis	98	16.5
COPD	65	10.9
active Cancer	67	11.3
CKD	113	19.0
Asthma	42	7.1
Chest pain		
No angina	451	75.9
Grade 1 angina	86	14.5
Grade 2 angina	58	9.6
Social problems		
No problems	281	47.2
1 problem	197	33.1
2 to 7 problems	117	19.7
Diagnosis at registry entry		
Myocardial infarction	252	42.4
Ischemic heart disease	287	48.2
Other	56	9.4
Depression diagnosis (CIS-R)	595	5.9
PHQ-9 score (>10)	595	16.8
PHQ-9 somatic dimension*	595	2 [0-12]
PHQ-9 cognitive dimension**	595	0 [0-12]
Age registry entry*	595	60.6 [25.6-97.2]
Quality of life score*	595	3 [3-8]
Anxiety score*	595	4 [0-21]
IMD*	593	16.9 [1.8-62.4]

COPD: chronic obstructive pulmonary disease; CKD: chronic kidney disease; IMD: Index of Multiple Deprivation
*median [min-max]

**standardised to a scale of 0-12 for comparative purposes

Table 2. Associations between baseline and time-dependent depressive symptom dimensions and new cardiac event at follow-up (95 events/595)

	Baseline analysis		Time-dependent analysis	
	HR (95% CI)	p	HR (95% CI)	p
Univariate model ¹				
Somatic dimension	1.13 (1.05-1.21)	0.0005	1.12 (1.04-1.19)	0.001
Cognitive dimension	1.09 (1.02-1.16)	0.01	1.09 (1.02-1.16)	0.01
Multivariate model ²				
Somatic dimension	1.13 (1.06-1.22)	0.0007	1.12 (1.05-1.20)	0.001
Cognitive dimension	1.09 (1.02-1.17)	0.01	1.11 (1.03-1.18)	0.004

¹ adjusted for sex

² further adjusted for diagnosis at entry (MI no/yes), smoking, and time-dependent chest pain, hypertension, COPD, active cancer

Table 3. Association between baseline and time-dependent PHQ-9 items and NCE (multivariate Cox Model¹)

n/N=95/595	Baseline depression		Time-dependent Depression	
	HR (95% CI)	p	HR (95% CI)	p
Somatic dimension items (item, n°):				
Trouble with sleep (3)	1.18 (0.98-1.41)	0.08	1.23 (1.03-1.46)	0.03
Feeling tired or having little energy (4)	1.26 (1.03-1.53)	0.02	1.20 (0.99-1.44)	0.06
Poor appetite or overeating (5)	1.40 (1.15-1.71)	0.0008	1.31 (1.07-1.60)	0.009
Slowness or restlessness (8)	1.38 (1.07-1.77)	0.01	1.30 (1.03-1.65)	0.03
Cognitive dimension items (item, n°):				
Little interest/pleasure in doing things (1)	1.31 (1.05-1.63)	0.02	1.26 (1.01-1.57)	0.04
Feeling down, depressed or hopeless (2)	1.37 (1.07-1.74)	0.01	1.32 (1.05-1.66)	0.02
Feeling like a failure (6)	1.27 (0.98-1.65)	0.07	1.39 (1.08-1.79)	0.01
Trouble concentrating (7)	1.15 (0.90-1.49)	0.3	1.25 (0.99-1.58)	0.06
Thoughts that better off dead (9)	1.25 (0.88-1.75)	0.2	1.29 (0.87-1.91)	0.2

¹adjusted for sex, diagnosis at entry (MI no/yes), smoking, and time-dependent chest pain, hypertension, COPD, active cancer