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One-year longitudinal study of psychological distress and self-assessed health in survivors of out-of-hospital cardiac arrest

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ABSTRACT

Objectives Few studies have investigated the psychological and health-related outcome after out-of-hospital cardiac arrest (OHCA) over time. This longitudinal study aims to evaluate psychological distress in terms of anxiety and depression, self-assessed health and predictors of these outcomes in survivors of OHCA, 3 and 12 months after resuscitation.

Methods Recruitment took place from 2008 to 2011 and survivors of OHCA were identified through the national Swedish Cardiopulmonary Resuscitation Registry. Inclusion criteria were age ≥18 years, survival ≥12 months and a Cerebral Performance Category score ≤2. Questionnaires containing the Hospital Anxiety and Depression Scale and European Quality of Life 5 Dimensions 3 Level (EQ-5D-3L) were administered at 3 and 12 months after the OHCA. Participants were also asked to report treatment-requiring comorbidities.

Results Of 298 survivors, 85 (29%) were eligible for this study and 74 (25%) responded. Clinically relevant anxiety was reported by 22 survivors at 3 months and by 17 at 12 months, while clinical depression was reported by 10 at 3 months and 4 at 12 months. The mean EQ-5D-3L index value increased from 0.82 (±0.26) to 0.88 (±0.15) over time. There were significantly less symptoms of psychological distress (p=0.01) and better self-assessed health (p=0.003) at 12 months. Treatment-requiring comorbidity predicted anxiety (OR 4.07, p=0.04), while being female and young age predicted poor health (OR 6.33, p=0.04; OR 0.91, p=0.002) at 3 months. At 12 months, being female was linked to anxiety (OR 9.23, p=0.01) and depression (OR 14.78, p=0.002), while young age predicted poor health (OR 0.93, p=0.003).

Conclusion The level of psychological distress and self-assessed health improves among survivors of OHCA between 3 and 12 months after resuscitation. Higher levels of psychological distress can be expected among female survivors and those with comorbidity, while survivors of young age and who are female are at greater risk of poor health.

INTRODUCTION

Out-of-hospital cardiac arrest (OHCA) is a deadly condition that affects approximately 5000 inhabitants in Sweden each year.1 The survival rate of OHCA is low, although great efforts have been made to raise the number of survivors in recent years; consequently, the 30-day survival rate in Sweden has increased from 4.5% in 1992 to 11.4% in 2017.2 When the heart stops, hypoxic brain injury occurs within minutes and is the most critical factor in determining the chance of survival and level of functioning, if resuscitation is successful.3 Following cerebral hypoxia, cardiac arrest survivors are known to be at risk of cognitive and emotional deficits.4–7 Although most survivors with a good neurological outcome return to high function and quality of life,8,9 long-term psychological problems have been reported by up to one-third in Swedish samples.10,11 Furthermore, the ability to pinpoint survivors who need psychiatric rehabilitation will become more important as survival rates continue to increase.

There is already an extensive body of knowledge regarding the outcome after cardiac arrest in different patient populations and time points. Time to awakening, gender, age, role of bystander, use of hypothermia and percutaneous coronary intervention have previously been related to the outcome after OHCA.12–20 Despite the current emphasis on rehabilitation, there is a scientific need for longitudinal studies to evaluate health and psychological outcomes in cardiac arrest survivors.21–24 The median interval between OHCA and rehabilitation is only 6 months in Sweden.25–27 The aim of this study is to evaluate psychological distress in terms of anxiety and depression, self-assessed health and predictors of these outcomes in survivors of OHCA, 3 and 12 months after resuscitation.
cardiac arrest. However, few studies have investigated the psychological and health-related outcome in the same sample over time, and little is known about factors associated with postcardiac arrest improvement in these aspects. The interpretation of the available literature is complicated by high heterogeneity in study outcomes, and knowledge regarding psychological distress and self-assessed health after OHCA in relation to the average population is lacking.

We have previously reported that reduced well-being is experienced by half of OHCA survivors at 3 months, and that female survivors report significantly more anxiety and worse health compared with males. In this study, we will investigate how the psychological and health-related outcomes change in survivors of OHCA with good neurological functioning between 3 and 12 months after resuscitation. Further, we will also evaluate gender differences as well as predictors of psychological distress and self-assessed poor health at these time points.

**METHODS**

**Design and setting**

This was a longitudinal study with self-administered postal questionnaires sent to survivors 3 and 12 months after the OHCA. The setting was a region in Sweden with 1.6 million inhabitants and nine hospitals that treat cardiac arrest. The frequency of OHCA was 850 cases annually and the survival rate approximately 10%.

**Study population**

The study population was identified through the national Swedish Cardiopulmonary Resuscitation Registry (SCRR). Every person who suffered from an OHCA and for whom resuscitation was initiated should be included in the SCRR. To ensure complete coverage, there was a specially trained nurse at each hospital who could identify missing cases and match patients treated for OHCA with data from the SCRR.

Recruitment took place from 1 January 2008 to 31 December 2011 and every adult survivor of OHCA in the region was assessed for inclusion. The study inclusion criteria were age ≥18 years, survival ≥12 months and a Cerebral Performance Category (CPC) score ≤2 at discharge, denoting good neurological functioning.

Exclusion criteria were cardiac arrest due to trauma, attempted suicide, intoxication or abuse. Survivors with other severe illness and survivors treated but not residing in the region were also excluded.

**Procedure and questionnaire**

Eligible survivors received a postal questionnaire in Swedish 3 months after the OHCA. Survivors who responded to the questionnaires and gave a written consent were included. Participants at 3 months then received a second questionnaire at 12 months. A reminder was sent if no response was received after 3 weeks. The questionnaire consisted of the Hospital Anxiety and Depression Scale (HADS) and European Quality of Life 5 Dimensions 3 Level (EQ-5D-3L). Study participants were also asked to report treatment-requiring comorbidity.

The HADS is a 14-item screening instrument developed to detect psychological distress in terms of clinically significant anxiety and/or depression. It is divided into two subscales, seven items regarding anxiety (HADS-anxiety) and seven items regarding symptoms of depression (HADS-depression). Each item has four levels, scored 0–3, with a maximum score of 21 in each subscale. Scoring >7 in one subscale indicates mild to moderate problems and a score >10 indicates severe problems. Although the HADS has never been thoroughly evaluated in a cardiac arrest population, it is considered a reliable measurement for depression and anxiety in patients with somatic illness and is frequently used in cardiovascular research.

EQ-5D-3L is designed to measure subjective health and includes the EQ-5D-3L descriptive system and a Visual Analogue Scale (EQ VAS). The descriptive system comprises five dimensions (mobility, self-care, usual activities, pain/discomfort and anxiety/depression); each dimension has three levels (no problems, moderate problems, severe problems) denoting a score of 1–3. The total score in the EQ-5D-3L descriptive system can be converted to an index value which ranges between −0.594 and 1.00, calculated with the UK EQ-5D index tariff. An index value of 1.00 indicates full health, while negative values represent health states worse than death. EQ VAS is visualised as a continuum of value scale where the patient assesses his or her current health on a scale from 0 to 100, with endpoints labelled ‘best imaginable health state’ and ‘worst imaginable health state’. EQ-5D-3L is commonly used in cardiovascular research. However, the use of EQ-5D-3L in OHCA populations has been questioned as the instrument has shown a limited ability to differentiate between survivors of good health.

**Statistics and data analysis**

Statistical analyses were executed with IBM SPSS Statistics V.24.0. All p values were two-sided and interpreted at a 0.05 significance level. Non-parametric tests were used for all comparisons as all dependent variables were non-normally distributed. The χ² test was applied for nominal values and the Mann-Whitney U-test for ordinal values. Wilcoxon signed ranks test was used to explore the change between 3 and 12 months. Predictors of psychological distress (HADS) and self-assessed poor health (EQ-5D-3L index values) were investigated using multivariate binary logistic regression models. The outcome was dichotomised as good or bad with regard to the mean scores in two Swedish reference populations, matched to gender and age-group. A bad outcome was considered a score 2 SD below the mean of the reference population. Reference values regarding the HADS were drawn from a random sample gathered 1997 in Jämtland County, Sweden (n=624). Similarly, reference values regarding the EQ-5D-3L index values were drawn from a random sample gathered 2001 in Stockholm County, Sweden.
The predictors were included based on clinical reasoning and are presented in Table 1. With regard to the sample size, only six variables were evaluated and correlations between them were examined using Spearman’s rank correlation; only variables with a Spearman’s r ≤0.7 were accepted. Univariate analysis was conducted separately between each individual predictor. The significant level of the univariate analysis was 0.25 and predictors with a p value >0.25 were excluded from further analysis. Hosmer and Lemeshow tests and receiver operating characteristics curves were performed to ensure the reliability of the logistic regression models.

**Patient and public involvement**

None of the included patients were involved in the design or conduct of this study and no patient opinion regarding the subject has been obtained. The results will be reported to the Health and Medical Care Committee of the Regional Executive Board, Region Västra Götaland, Sweden.

**RESULTS**

There were 3082 cases of OHCA in the region during the study period, of which 298 were alive at 30 days. One hundred and fifty survivors fulfilled the inclusion criteria for a 3-month follow-up and 94 responded. Of the respondents at 3 months, 85 were included at 12 months and 74 responded again (87% response rate); in the group that did not receive a second questionnaire, five had died and four were lost due to logistical errors (figure 1).

The representativeness of the study population has been investigated in a prior study, and there were no significant differences regarding age, gender, cardiac arrest circumstances or in-hospital interventions between respondents and non-respondents. However, there were differences between participants and the survivors that only responded once. Double respondents were significantly younger (Z=2.316, p=0.021), had a lower frequency of ventricular fibrillation or ventricular tachycardia as initial rhythm (χ²=5.067, p=0.032). Background characteristics and self-reported comorbidity of the participants are presented in Table 1.

Mild to moderate anxiety (≥7 in HADS-A) was reported by 22 (30%) survivors at the 3-months follow-up and by 17 (23%) at 12 months. Mild to moderate depression (≥7 in HADS-D) was reported by 10 (14%) survivors at 3 months and by four (5%) at 12 months. A score indicating severe anxiety (≥10 in HADS-A) was reported by 12 survivors at 3 months and by eight at 12 months, while severe depression (≥10 in HADS-D) was reported by three at 3 months and one at 12 months. Overall, survivors of OHCA reported a significantly improved psychological and health-related outcome over time (Table 2). About half of the survivors reported less psychological distress in the HADS at 12 months (51% decrease in anxiety and 46% in symptoms of depression); although one-third reported an increase in psychological distress (30% increase in anxiety and 28% in symptoms of depression). Two-thirds of female survivors reported more psychological distress at the 12-month follow-up, corresponding to one-fifth among male survivors. Comparably, half of the survivors reported better self-assessed health at 12 months in EQ-5D-3L, whereas 41% reported the same health state at 3 and 12 months. Among the survivors who remained unchanged, 73% reported an EQ-5D-3L index value of 1.00, denoting full health at both 3 and 12 months.
time points. However, only one survivor entered full health at both times in EQ VAS (figure 2).

Binary logistic regression analyses were preformed to find predictors of psychological distress and self-assessed poor health (table 3). The presence of treatment-requiring comorbidity was found to predict more anxiety at 3 months after resuscitation (OR 4.07, p=0.04), but there was no significant OR regarding comorbidity at 12 months; instead, being female was found to predict more anxiety (OR 9.23, p=0.01) and more symptoms of depression (OR 14.78, p=0.002) at this time. Being female was also found to predict self-assessed poor health at 3 months (OR 6.33, p=0.04), while young age was found to predict poor health at 3 and 12 months, respectively (OR 0.91, p=0.002; OR 0.93, p=0.003).

**DISCUSSION**

One objective of this study was to investigate how the psychological and health-related outcomes change between 3 and 12 months in survivors of OHCA with good neurological functioning. We found an encouraging progression as survivors at large showed less psychological distress and better self-assessed health over time. In comparison, participants in this study even surpassed the average level of depression and self-assessed health in the Swedish population at their 12-month follow-up. However, we also found that even though about half of OHCA survivors report a decrease in psychological distress between 3 and 12 months, almost one-third report increased psychological distress; hence, our results indicate that the psychological outcome after OHCA may differ considerably on an individual level.

As stated earlier, few prior longitudinal studies have investigated the psychological and health-related outcomes after cardiac arrest. In accordance with our results, two small sample studies have previously reported a decrease of anxiety and depressive symptoms in cardiac arrest survivors over time. In contrast, a rather large

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**Figure 1** Flowchart of study participants. CPC, Cerebral Performance Category; OHCA, out-of-hospital cardiac arrest. 1) Administrative reasons include survivors documented with an incorrect social security number; not able to be identified within 3 months or not residing in the region. OHCA-circumstances include trauma, attempted suicide, drowning and intoxication as cause of the cardiac arrest or severe illness afterwards.
A prospective study found no change in anxiety or depressive symptoms between 1 and 12 months. It is possible that the heterogeneity in psychological outcomes observed in this study could explain a part of the discrepancies in the available literature. As regards self-assessed health, the improvements seen in this study confirm the results of two previous longitudinal studies. However, these previous studies found that the greatest health-related improvements occurred in the early stages of recovery, within 3 months after cardiac arrest. Moreover, although the ‘minimal clinically important difference’ in the HADS and EQ-5D-3L among survivors of cardiac arrest is unknown, estimates from other patient populations indicate that the improvements seen in this study might not be clinically noticeable.

A second objective was to evaluate predictors of psychological distress self-assessed poor health after OHCA. Since it is already known that cardiac arrest survivors generally report few psychological problems and good health, it is difficult to define a threshold for a bad outcome among those who perform well neurologically. In this study, the outcome was dichotomised as good or bad in comparison to mean scores from the average Swedish population. Prior studies have concluded that many survivors reach a psychological state and health comparable to the average population, although no study has previously investigated predictors of this outcome. In addition, there is also a lack of knowledge regarding factors associated with postcardiac arrest improvement over time.

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Comparison of mean scores at 3 and 12 months in the HADS and EQ-5D-3L (n=74)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean (SD)</td>
</tr>
<tr>
<td>HADS</td>
<td></td>
</tr>
<tr>
<td>HADS-anxiety</td>
<td>5.6 (±4.8)</td>
</tr>
<tr>
<td>HADS-depression</td>
<td>3.4 (±3.5)</td>
</tr>
<tr>
<td>Total score</td>
<td>9.0 (±7.8)</td>
</tr>
<tr>
<td>EQ-5D-3L</td>
<td></td>
</tr>
<tr>
<td>EQ VAS (n=67)</td>
<td>73 (±18)</td>
</tr>
<tr>
<td>Index value</td>
<td>0.82 (±0.26)</td>
</tr>
</tbody>
</table>

*P < 0.05 indicates statistical significance.

EQ-5D-3L, European Quality of Life 5 Dimensions 3 Level; HADS, Hospital Anxiety and Depression Scale; VAS, Visual Analogue Scale.

Figure 2 Scatter plot with individual reported outcomes in the HADS and EQ-5D-3L at 3 months (x-axis) and 12 months (y-axis). Dots above the diagonal lines indicate a higher score at the 12-month follow-up. Scores from seven participants are missing in EQ VAS. EQ-5D-3L, European Quality of Life 5 Dimensions 3 Level; HADS, Hospital Anxiety and Depression Scale; VAS, Visual Analogue Scale.
We found that comorbidity is associated with higher levels of anxiety compared with the average population at 3 months after OHCA. However, we did not find a significant association between comorbidity and anxiety or depression at the 12-month follow-up. In contrast, being female was strongly associated with psychological distress at 1 year after resuscitation, but this was not significant at 3 months. We also found a significant association between being female and self-assessed poor health at 3 months, while young age was associated with self-assessed poor health at both 3 and 12 months.

Studies investigating prognostic factors regarding the psychological or health-related outcomes after cardiac arrest are scarce. However, receiving an ICD has been linked to higher levels of psychological distress, but this effect has not been shown in a cardiac arrest population. We did not find any significant association regarding ICD in our regression models either. Young age and being female have previously been associated with anxiety among cardiac arrest survivors, but there were no significant association between psychological distress and age in our study population. Nevertheless, we did find a considerably higher likelihood of depression among female survivors of OHCA at 12 months, which has not previously been reported. Also novel is the likelihood of a better self-assessed health among survivors of higher age.

The European Resuscitation Council has provided guidelines for postcardiac arrest rehabilitation since 2015 and has pointed out the need of a structured follow-up care, including physical, neurological and emotional screening. Still, far from all cardiac arrest survivors in Sweden are offered rehabilitation or neuropsychological evaluation, despite recommendations from the Swedish Resuscitation Council. Instead, survivors in Sweden are often followed based on the aetiology of the cardiac arrest with little to no focus on emotional sequelae. Our results indicate that even if the majority of OHCA survivors live without psychological distress and with good self-assessed health, there are individuals that experience major problems. Consequently, it should be expected that some survivors will benefit from psychiatric rehabilitation, and survivors of young age, who are female, and those with treatment-requiring comorbidity should be screened early on.

**Strengths and limitations**

This study has several strengths. During the study period of 4 years, a specially trained nurse was placed at every hospital in a large region of Sweden. With this approach, we were able to collect data from virtually every case of OHCA. The study design also allowed us to match non-respondents with data collected from the SCRR at the time of their hospitalisation, thus enabling comparisons between respondents and non-respondents. Consequently, we were able to confirm that the study population represents the group at large in these aspects. Another strength is that the outcomes in this study could be compared with

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**Table 3** Predictors of psychological distress and self-assessed poor health (*n*=70)

<table>
<thead>
<tr>
<th>Predictors</th>
<th>3 months</th>
<th>12 months</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>OR (95% CI)</td>
<td>SE</td>
</tr>
<tr>
<td>HADS-anxiety</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>0.99 (0.95 to 1.03)</td>
<td>0.02</td>
</tr>
<tr>
<td>Comorbidity</td>
<td>4.07 (1.07 to 17.00)</td>
<td>0.73</td>
</tr>
<tr>
<td>Being female</td>
<td>4.94 (0.91 to 26.83)</td>
<td>0.86</td>
</tr>
<tr>
<td>Hypothermia</td>
<td>1.25 (0.43 to 3.62)</td>
<td>0.54</td>
</tr>
<tr>
<td>HADS-depression</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>0.99 (0.94 to 1.04)</td>
<td>0.03</td>
</tr>
<tr>
<td>Comorbidity</td>
<td>1.45 (0.33 to 6.44)</td>
<td>0.76</td>
</tr>
<tr>
<td>Being female</td>
<td>3.71 (0.90 to 15.37)</td>
<td>0.78</td>
</tr>
<tr>
<td>Hypothermia</td>
<td>1.16 (0.37 to 3.61)</td>
<td>0.58</td>
</tr>
<tr>
<td>ICD</td>
<td>1.52 (0.40 to 5.69)</td>
<td>0.68</td>
</tr>
<tr>
<td>EQ-5D-3L</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>0.91 (0.86 to 0.97)</td>
<td>0.03</td>
</tr>
<tr>
<td>Comorbidity</td>
<td>1.19 (0.28 to 5.01)</td>
<td>0.73</td>
</tr>
<tr>
<td>Being female</td>
<td>6.33 (1.03 to 38.81)</td>
<td>0.93</td>
</tr>
</tbody>
</table>

*P < 0.05 indicates statistical significance.

EQ-5D-3L, European Quality of Life 5 Dimensions 3 Level; HADS, Hospital Anxiety and Depression Scale; ICD, implantable cardioverter defibrillator.
the average Swedish population. Knowledge regarding factors of importance for a psychological state and self-assessed health in line with the normal population after OHCA in Sweden is missing. This information will help to generalise the overall picture of OHCA survival and provide guidance regarding which survivors who need psychiatric rehabilitation. Furthermore, the response rate was very high at 12 months.

The main limitations of this study are the retrospective design and the restrictive inclusion of survivors with good neurological functioning (CPC-score ≤2), which limits the applicability on OHCA survivors more broadly. However, we made the assessment that inclusion of survivors with a severe cerebral dysfunction would complicate the interpretability of the results. Furthermore, it is not possible to differentiate between patients with a CPC-score of 1 and 2. However, as mentioned above, we lack knowledge of the minimal clinically important difference regarding the HADS and EQ-5D-3L in this population. The strict inclusion is also reflected by the relatively small sample size, resulting in a low count of events per variable in the logistic regression (five variables were evaluated in 70 participants), why the significance of these results should be interpreted with caution. Another methodical limitation in this study is the use of EQ-5D-3L to measure self-assessed health. A recent study concluded that EQ-5D-3L might underestimate the health in OHCA survivors, yielding high ceiling effects, which was apparent in the EQ-5D-3L descriptive system in this study as well. Although EQ-5D showed higher interpretability with a more varied outcome in our study population, it is known that generic measures of health often fail to capture the true complexity of the health state in survivors of cardiac arrest. Finally, some survivors were excluded due to severe illness and a few non-respondents mentioned poor health as the reason for not participating. Therefore, it is possible that the participants in this study had a better outcome than survivors more generally.

CONCLUSIONS

On a group level, there is a significant improvement in the psychological and health-related outcomes among survivors of OHCA between 3 and 12 months after resuscitation. Higher levels of psychological distress can be expected among female survivors and those with treatment-requiring comorbidity, while young and female survivors are at risk of a self-assessed poor health. These findings should be considered in the follow-up after OHCA, although future research is required to identify further predictors regarding the long-term outcome and chance of improvement postcardiac arrest.

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Contributors All authors have made a substantial contribution to the conception and design of the study. AV analysed the data and wrote the manuscript with input from all authors. KSS and AA initiated and planned the study, collected the data, assisted in the interpretation of the data and took part in drafting the manuscript. JH took part in the planning of the study, contributed with external expertise and have continuously revised the manuscript. DJ helped to analyse the data and have continuously revised the manuscript.

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Competing interests None declared.

Patient consent for publication Not required.

Ethics approval Ethical permission was granted by the Regional Ethics Review Board, Western Sweden (no. 465–7). Written consent was obtained from all participants.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement Due to ethical restrictions, data are available on request. Interested researchers may submit a request for data to the authors (adam.viktorisson@gu.se). According to Swedish regulation: http://www.epn.se/en/start/ regulations/, the permission to use data is only for what has been applied for and then approved by the Ethical board. To not follow the regulations is seen as scientific misconduct.

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