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Childhood adversity is associated with hospitalisations and survival following external causes and non-communicable diseases: a 46-year follow-up of a Stockholm birth cohort

Josephine Jackisch ,^{1,2} Ylva B Almquist ¹

► Additional supplemental material is published online only. To view, please visit the journal online (<http://dx.doi.org/10.1136/jech-2022-219851>).

¹Department of Public Health Sciences, Stockholm University, Stockholm, Sweden
²Max Planck Institute for Demographic Research, International Max Planck Research School for Population, Health and Data Science, Rostock, Germany

Correspondence to

Dr Josephine Jackisch, Department of Public Health Sciences, Stockholm University, Stockholm, Sweden; josephine.jackisch@su.se

Received 23 September 2022
Accepted 10 January 2023
Published Online First
3 February 2023

ABSTRACT

Background Childhood adversity indicated by involvement with child welfare services (ICWS) is associated with increased risks of disease and injuries in young adulthood. It is yet unknown whether such risks are limited to external causes and mental and behavioural disorders or whether they extend beyond early adulthood and to non-communicable diseases (NCDs) with later onset. Moreover, it has not been explored whether ICWS associates with decreased survival prospects following hospitalisation.
Methods Based on prospective data for a 1953 Stockholm birth cohort (n=14 134), ICWS was operationalised distinguishing two levels in administrative child welfare records (ages 0–19; ‘investigated’ and ‘placed’ in out-of-home care (OHC)). Hospitalisations and all-cause mortality (ages 20–66) were derived from national registers. Hospitalisation records were categorised into external causes and NCDs, and nine subcategories. Negative binomial regression models were used to estimate differences in hospitalisation risks between those with and without experiences of ICWS and Cox survival models to estimate mortality after hospitalisation.

Results Placement in OHC was associated with higher risks of hospitalisation due to external causes and NCDs and all investigated subcategories except cancers. Risks were generally also elevated among those investigated but not placed. ICWS was further linked to higher mortality risks following hospitalisation.

Conclusion Differential risk of morbidity and differential survival may explain inequalities in mortality following childhood adversity. We conclude that the healthcare sector might play an important role in preventing and mitigating the elevated risks of externally caused morbidity, disease and premature mortality observed among those with a history of ICWS.

INTRODUCTION

Experiences of childhood adversity are highly prevalent and result in high costs not only to the individual but also to all of society.^{1 2} Therefore, childhood adversity constitutes an important public health concern.³ Individuals who experience involvement with child welfare services (ICWS) carry an exceptionally high burden of adverse childhood experiences.^{4 5} Indeed, it has been consistently demonstrated that ICWS is associated with an increased risk of premature mortality.^{6–8}

WHAT IS ALREADY KNOWN

People with experiences of involvement with child welfare services (ICWS) have often high levels of adverse childhood experiences and experience higher risks of hospitalisation in adulthood due to external causes as well as mental and behavioural disorders. As most previous prospective studies have ended their follow-ups by early adulthood, it remains unclear whether there are also increased risks for other non-communicable diseases with a later onset and whether childhood adversity is linked to differences in survival after onset of disease.

WHAT THIS STUDY ADDS

This study demonstrates that adults with experiences of ICWS generally have higher risks of being hospitalised due to external causes and non-communicable diseases. Following an initial hospitalisation, they have lower chances of survival.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

The findings highlight several possible entry points for policy and intervention, which include preventing experiences of childhood adversity, mitigating health-related consequences of such experiences and improving healthcare outcomes in this vulnerable group of individuals.

However, the pathways underlying these differences in mortality are not well known. The current study explores two potential explanations: differential risk of hospitalisation due to external causes and non-communicable diseases (NCDs) and differential survival prospects following such hospitalisations. Such an inquiry may lead us to better understand the healthcare sector's role in mitigating the risk of mortality among individuals with experiences of childhood adversity. Individuals at risk can be identified early through ICWS, which might guide the development of appropriate interventions.

Theoretically, experiences of childhood adversity—and, particularly, those so severe that they lead to ICWS—can be assumed to ‘get under the skin’, changing the epigenome and other biological and psychological response systems.^{9 10} This inscription of social experiences within the body, from in-utero



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To cite: Jackisch J, Almquist YB. *J Epidemiol Community Health* 2023;**77**:209–215.

to death, is sometimes referred to as ‘embodiment’.¹¹ Through the process of embodiment, those exposed to childhood adversity might be more susceptible to disease development or more likely to experience more severe consequences following disease (including death).

Thus far, morbidity among individuals with experiences of ICWS has primarily been examined in terms of external causes or mental and behavioural disorders.^{12–14} Previous prospective longitudinal studies have, for example, noted increased risks for developing anxiety, depression, substance use disorders and self-harm^{13 15–19} up until early midlife among those placed in out-of-home care (OHC; foster family or institutional care).

Evidence is scarcer concerning risks of diseases with a later age at onset, including cardiovascular diseases (CVDs), cancers, respiratory diseases and diabetes. Alongside mental and behavioural disorders, they represent the main types of NCDs with an extensive individual and societal burden. These five subcategories—most of which are considered preventable—are together estimated to account for 90% of deaths in the European region.²⁰

Looking into the broader literature on childhood adversity, retrospective studies have indicated increased risks of developing NCDs among those with adverse childhood experiences.^{2 21–27} There is, however, a paucity of prospective studies. Those that do exist tend to show inconsistent results, for example, regarding the risks for cancer and diabetes, which might be due to relatively limited follow-ups.^{21 28–30}

Study aims

With the aim to explore the role of childhood adversity in relation to hospitalisation risks and survival prospects following hospitalisation, the present study draws on longitudinal prospective data from a Swedish cohort born in 1953 (n=14 134). Childhood adversity is approximated by experiences of ICWS (ages 0–19). Two groups of ICWS are differentiated, indicating the severity of ICWS experiences: individuals who were investigated but not placed, and individuals for whom the involvement resulted in placement in OHC. Following an outcome-wide approach to epidemiology,³¹ we include hospitalisations due to external causes and NCDs as well as all-cause mortality (ages 20–66). More specifically, this study investigates the following hypotheses:

1. Individuals with experiences of ICWS are at higher risk of being hospitalised due to external causes (including self-harm) and NCDs (including mental and behavioural disorders, CVDs, cancers, respiratory diseases and diabetes) when compared with individuals without experiences of ICWS.
2. Individuals with experiences of ICWS have a shorter time to death after initial hospitalisation due to external causes and NCDs compared with their majority population peers.
3. The relationships in hypotheses 1 and 2 follow a gradient that reflects the severity of ICWS experienced.

METHODS

Study population

The data material is the Stockholm Birth Cohort Multigenerational study (SBC Multigen). The cohort is defined as all children born in 1953 and living in the greater Stockholm metropolitan area in 1963, who could be probability matched to follow-up data from nationwide registers (n= 14,608).³² In this study, we restricted the analysis to those who did not die or emigrate without returning before age 20 (n=14,509). The analytical sample includes everyone without missing values

on the covariates (n=14,134). Missing observations in family education at age 10 were retained as a separate category.

Measurements

Involvement with child welfare services

In this cohort, ICWS has been shown to be a viable proxy for childhood adversity because of its predictive power for long-term health outcomes such as premature mortality.⁷ Other prospectively measured indicators of childhood adversity (e.g., accumulative experiences of household dysfunction) do not independently of ICWS predict mortality or educational attainment.³³

The SBC Multigen contains child welfare records from municipal social registers covering the period between 1953 and 1972 (ages 0–19). Besides placement in OHC, these data include referrals to child welfare services that did not result in the child being taken out of the family but rather cases that were dismissed or resulted in warnings or in-home services. This offers an interesting alternative comparison group, encompassing individuals who have displayed signs of adversity severe enough to come to the attention of the child welfare services but not severe enough to justify a placement in OHC. Based on this, we distinguished three mutually exclusive ICWS groups following the severity of ICWS: (1) those without any record in the child welfare register (‘without ICWS’; who are used as reference group in the main analysis), (2) child welfare investigations not resulting in OHC placement (‘investigated’), (3) placement in OHC, including foster and institutional care (‘placed’). The highest order took precedence in coding these groups.

Hospitalisation

From the Swedish National Patient Register, we derived data on in-patient care between 1973 and 2019 (ages 20–66), that is, cases of disease that required hospitalisation (at least one overnight stay at the hospital). A total of 11 cause groups were selected based on both primary and secondary diagnoses and classified according to the 10th revision of the International Classification of Diseases and the corresponding codes in the eighth and ninth revisions (see online supplemental table 1). We identified two major cause groups: external causes and NCDs. External causes include hospitalisations due to causes outside the body, for example, injuries, poisoning, accidents, violent assaults. We separated out the subcategory of (intentional and unintentional) self-harm. Within NCDs, we distinguished five subcategories: CVDs, cancers, respiratory diseases, diabetes and mental and behavioural disorders. Under mental and behavioural disorders, we further separate three subcategories: anxiety, depression and substance use disorders. Besides those subcategories, NCDs as an overall category also include in situ and benign neoplasms; diseases of the blood; other endocrine and metabolic disorders; musculoskeletal disorders; diseases of the nervous, digestive or genitourinary system and diseases of the eye, ear and skin.

Death

Mortality data were drawn from the National Cause of Death Register as death by any cause between 1 January 1973 and 31 December 2019 (ages 20–66). Since chronic diseases tend to co-occur, cause-specific mortality might include uncertainty in the registered causes of death. For this reason and to ensure sufficient statistical power, we preferred using all-cause mortality.

Covariates

ICWS is associated with the socioeconomic conditions of the family, and such factors have also been shown to influence adulthood morbidity and mortality. Therefore, we included the following covariates: born out of wedlock; mother's age at birth; occupational status of family at birth (usually assessed through father's occupation but, if unavailable, based on the mother's occupation) and family education at age 10.

Sex

All analyses are reported separately for men and women, based on the biological sex assigned at birth.

Statistical analysis

Follow-up started on 1 January 1973 and ended at death, emigration or end-of follow-up on 31 December 2019. Within this observation window, 519 people got censored at the date of emigration, and 1635 people died.

Risk of hospitalisation

We analysed hospitalisation between ages 20 and 66 for each major cause group and subcategory, stratified by sex and ICWS group. First, we calculated hospitalisation prevalence (expressed as %), deriving 95% CIs from predicted margins based on logistic regression analysis. Next, we calculated incidence rates of hospitalisation per 1000 person-years at risk. Incidence measures the occurrence of new cases during a certain period and, thus, signifies a person's probability of being hospitalised with a disease.

To further examine the first hypothesis, we conducted multivariate modelling to predict frequencies of hospitalisation within each cause group (thereby taking repeated hospitalisations into account). Model fit statistics for the count variable were derived for the covariate-adjusted Poisson, zero-inflated Poisson and negative binomial models (online supplemental table 2). The negative binomial models fit the data best for all outcomes and were, thus, used to examine the differences in the risk of hospitalisation by ICWS groups. We report adjusted incidence rate ratios (IRRs) with 95% CIs.

Time to death

The second hypothesis concerns differential survival after hospitalisation. To investigate this, we used Cox survival models estimating the chances of survival on a calendar time scale from the date of the first cause-specific hospitalisation to death from all causes. The few tied failure times were dealt with by the Efron method. We adjusted these models for covariates and for age at first cause-group-specific hospitalisation, reporting HRs over the follow-up period with 95% CIs.

In order to counteract the likelihood of false-positive chance findings due to multiple outcomes, we used Bonferroni correction to correct α (the significance level considered) to <0.004 . However, in the tables, we reported CIs at a 95% level to not confuse readers, whereas $p < 0.004$ is marked with a bold face. Data management was conducted in R (V.4.1.0), and descriptives and analyses were made in Stata (V.17.0) using commands 'nbreg' and 'stcox'. Plots were produced in R.

RESULTS

Characteristics of the study population are shown in table 1. ICWS was relatively prevalent in the cohort: 26.8% of men and 14.5% of women had such experiences in ages 0–19. The prevalence of OHC was 9.2% among men and 8.4% among women. Those with experiences of ICWS were more likely to have been born out of wedlock, have teenage mothers, have fathers with a manual occupational background and were less likely to have parents with secondary or postsecondary educational degrees.

Table 2 describes the absolute prevalence and incidence rates of hospitalisation. For all subcategories, men and women placed in OHC have higher prevalence and incidence rates than those without experiences of ICWS. The investigated children's estimates fall between those two groups with few exceptions. Moreover, we found a generally higher median number of hospitalisations (frequency) and a lower median age at first hospitalisation (onset) for both ICWS groups ('investigated' and 'placed'), except for cancers and respiratory diseases (online supplemental table 3).

Table 1 Sample characteristics, stratified by sex and ICWS group

	Men				Women			
	Without ICWS	Investigated	Placed	Total	Without ICWS	Investigated	Placed	Total
	%	%	%	%	%	%	%	%
Born out of wedlock (yes)	4.32	7.91	20.88	6.55	4.37	12.80	23.02	6.45
Mother's age at birth								
Teenager	3.66	4.89	10.23	4.51	3.54	5.92	9.45	4.18
20–35 years of age	80.43	80.03	75.57	79.89	80.66	78.20	75.77	80.10
>35 years of age	15.91	15.08	14.20	15.60	15.79	15.88	14.78	15.71
Occupational status of father at birth								
Non-manual worker	57.25	39.61	34.52	52.03	55.01	38.63	33.68	52.22
Manual worker	42.75	60.39	65.48	47.97	44.99	61.37	66.32	47.78
Family education at age 10								
Primary	64.23	82.48	87.36	69.60	66.27	81.75	87.46	68.99
Secondary	19.14	9.78	7.10	16.37	18.17	9.00	7.73	16.74
Tertiary	10.92	3.99	2.41	8.91	10.06	3.32	1.20	8.90
Missing	5.71	3.75	3.13	5.12	5.50	5.92	3.61	5.37
Total, row per cent	73.2	17	9.2	100	85.5	6.1	8.4	100
Total, n	5273	1227	704	7204	5926	422	582	6930

ICWS, involvement with child welfare services.

Table 2 Hospitalisations in ages 20–66: prevalence (%) and incidence rates (hospitalisations per 1000 person-years), stratified by sex and ICWS group

	Hospitalisations among men ages 20–66 (n=7204)				Hospitalisations among women ages 20–66 (n=6930)			
	Without ICWS (n=5273)		Placed (n=704)		Without ICWS (n=5926)		Placed (n=582)	
	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)	Estimate (95% CI)
External causes (n cases=4448)								
Prevalence	30.5 (29.3 to 31.7)	44.0 (41.2 to 46.8)	46.0 (42.3 to 49.7)	27.3 (26.1 to 28.4)	34.6 (30.1 to 39.1)	40.0 (35.5 to 43.5)		
Incidence rate	7.7 (7.4 to 8.1)	12.6 (11.6 to 13.7)	14.5 (13.0 to 16.1)	6.6 (6.3 to 7.0)	9.0 (7.6 to 10.5)	10.5 (9.3 to 12.0)		
Self-harm (n cases=504)								
Prevalence	2.0 (1.7 to 2.4)	6.4 (5.1 to 7.8)	8.9 (6.8 to 11.1)	2.8 (2.4 to 3.2)	7.3 (4.9 to 9.8)	10.3 (7.8 to 12.8)		
Incidence rate	0.5 (0.4 to 0.6)	1.5 (1.2 to 1.8)	2.3 (1.8 to 2.9)	0.6 (0.5 to 0.7)	1.7 (1.2 to 2.4)	2.4 (1.9 to 3.1)		
NCDs (n cases=10097)								
Prevalence	67.4 (66.1 to 68.7)	76.0 (73.7 to 78.4)	78.8 (75.8 to 81.9)	71.7 (70.7 to 72.8)	78.0 (74.0 to 81.9)	84.3 (81.4 to 87.3)		
Incidence rate	22.2 (21.5 to 22.9)	27.8 (26.1 to 29.6)	35.0 (32.3 to 38.0)	26.6 (25.8 to 27.4)	35.8 (32.2 to 39.9)	42.1 (38.6 to 46.0)		
Mental and behavioural disorders (n cases=2013)								
Prevalence	11.3 (10.3 to 12.1)	23.6 (21.3 to 26.0)	37.1 (33.5 to 40.6)	10.2 (9.4 to 10.9)	21.8 (17.9 to 25.7)	29.9 (26.2 to 33.6)		
Incidence rate	2.6 (2.4 to 2.8)	5.8 (5.2 to 6.5)	10.8 (9.6 to 12.2)	2.3 (2.2 to 2.5)	5.5 (4.5 to 6.8)	8.0 (6.9 to 9.2)		
Anxiety (n cases=509)								
Prevalence	2.6 (2.2 to 3.0)	4.4 (3.4 to 5.5)	8.5 (6.4 to 10.6)	3.2 (2.7 to 3.6)	5.0 (2.9 to 7.1)	8.4 (6.2 to 10.7)		
Incidence rate	0.6 (0.5 to 0.7)	1.0 (0.8 to 1.3)	2.0 (1.6 to 2.6)	0.7 (0.6 to 0.8)	1.1 (0.8 to 1.7)	2.0 (1.5 to 2.6)		
Depression (n cases=596)								
Prevalence	3.1 (2.6 to 3.6)	4.6 (3.4 to 5.7)	8.9 (6.8 to 11.1)	3.8 (3.3 to 4.2)	6.2 (3.9 to 8.5)	11.0 (8.5 to 13.5)		
Incidence rate	0.7 (0.6 to 0.8)	1.1 (0.8 to 1.4)	2.3 (1.8 to 2.9)	0.8 (0.7 to 1.0)	1.5 (1.0 to 2.1)	2.5 (2.0 to 3.2)		
Substance use disorders (n cases=1176)								
Prevalence	5.8 (5.2 to 6.5)	20.4 (18.1 to 22.6)	28.4 (25.1 to 31.7)	4.0 (3.5 to 4.5)	15.4 (12.0 to 18.8)	20.3 (17.0 to 23.5)		
Incidence rate	1.3 (1.2 to 1.5)	4.9 (4.4 to 5.6)	7.9 (6.9 to 9.0)	0.9 (0.8 to 1.0)	3.6 (2.9 to 4.6)	5.0 (4.2 to 6.0)		
CVDs (n cases=3725)								
Prevalence	29.0 (27.8 to 30.2)	35.9 (33.3 to 38.6)	33.8 (30.3 to 37.3)	21.1 (20.1 to 22.2)	27.3 (23.0 to 31.5)	28.4 (24.7 to 32.0)		
Incidence rate	6.8 (6.5 to 7.1)	8.7 (8.0 to 9.5)	8.6 (7.6 to 9.7)	4.8 (4.6 to 5.1)	6.3 (5.3 to 7.6)	6.7 (5.8 to 7.8)		
Cancers (n cases=1942)								
Prevalence	11.5 (10.6 to 12.3)	12.2 (10.4 to 14.1)	13.5 (11.0 to 16.0)	15.7 (14.7 to 16.6)	16.8 (13.3 to 20.4)	16.7 (13.6 to 19.7)		
Incidence rate	2.5 (2.3 to 2.7)	2.7 (2.3 to 3.2)	3.2 (2.6 to 3.9)	3.5 (3.3 to 3.7)	4.0 (3.2 to 5.0)	3.9 (3.2 to 4.7)		
Respiratory diseases (n cases=1630)								
Prevalence	10.1 (9.3 to 10.9)	14.3 (12.4 to 16.3)	15.9 (13.2 to 18.6)	10.8 (10.0 to 11.6)	13.0 (9.8 to 16.2)	21.1 (17.8 to 24.5)		
Incidence rate	2.3 (2.1 to 2.5)	3.4 (3.0 to 4.0)	4.0 (3.3 to 4.8)	2.4 (2.3 to 2.6)	3.2 (2.5 to 4.1)	5.2 (4.4 to 6.2)		
Diabetes (n cases=624)								
Prevalence	4.7 (4.2 to 5.3)	6.8 (5.4 to 8.3)	6.7 (4.8 to 8.5)	3.2 (2.7 to 3.6)	4.5 (2.5 to 6.5)	6.0 (4.1 to 7.9)		
Incidence rate	1.1 (1.0 to 1.2)	1.5 (1.2 to 1.9)	1.6 (1.2 to 2.1)	0.7 (0.6 to 0.8)	1.0 (0.7 to 1.6)	1.3 (0.9 to 1.8)		
CVDs, cardiovascular diseases; ICWS, involvement with child welfare services; NCDs, non-communicable diseases.								

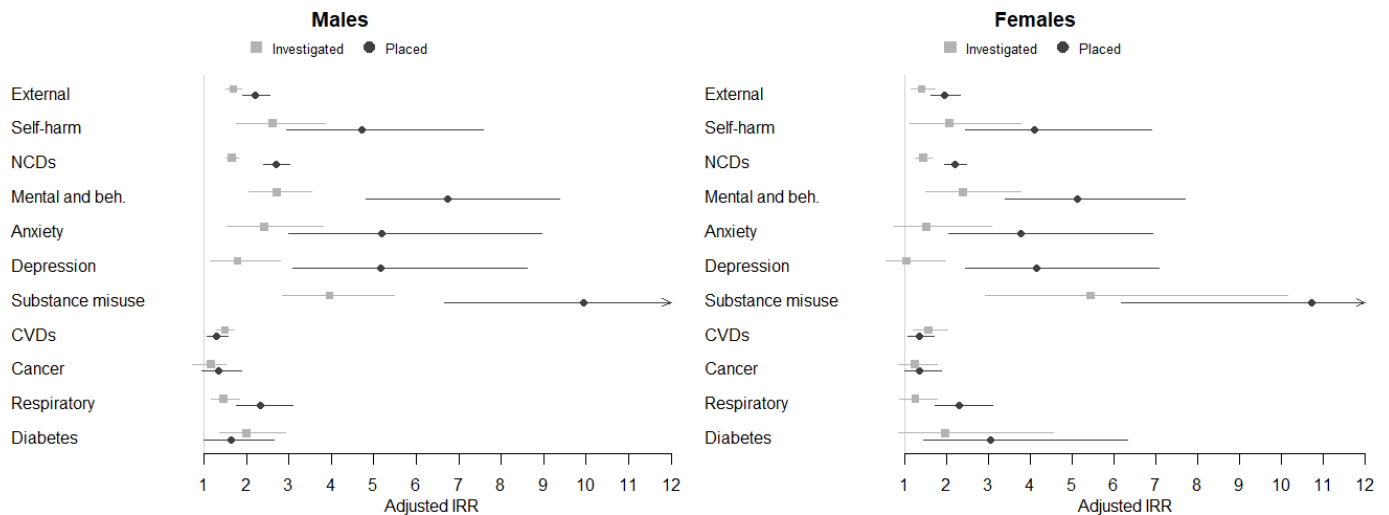


Figure 1 The association between ICWS and hospitalisation, stratified by sex and cause group. Results from the adjusted negative binomial regression models. 'Without ICWS' is the reference group (IRR=1.00). All models are adjusted for being born out of wedlock, mother's age at birth, occupational status at birth, and family education at age 10. CVDs, cardiovascular diseases; IRR, incidence rate ratio; NCDs, non-communicable diseases.

The results from the adjusted negative binomial regression of experiences of ICWS on cause-group-specific hospitalisation are plotted in figure 1. Individuals with ICWS were more likely to be hospitalised due to external causes (IRRs ranging from 1.7 to 2.2) and NCDs (IRRs: 1.7–2.7). Hospitalisation risks were particularly pronounced in the subcategory of mental and behavioural disorders (IRRs: 2.7–6.7), with those who experienced ICWS being more likely to be hospitalised due to substance use disorders (IRRs: 4.0–10.7) and anxiety (IRRs: 2.4–5.2). They also have some increased risks of hospitalisation due to self-harm (IRRs: 2.6–4.7), CVDs (IRRs: 1.3–1.6) and respiratory diseases (IRRs: 1.3–2.3). Estimates were less consistent for cancers (IRRs: 1.2–1.4). In general, associations are stronger for the placed group than for those investigated. Though IRRs for investigated women had similar directionality as for the

placed women, this group was small with larger CIs that often include one (online supplemental table 4).

Table 3 shows the HRs of all-cause mortality after hospitalisation across ages 20–66. Experiences of ICWS were associated with increased risks of death after hospitalisation due to external causes (HRs: 1.7–2.6) and NCDs (HRs: 1.6–2.3), and, particularly, after hospitalisation due to CVDs (HRs: 1.8–2.5). Except for investigated women, ICWS was also associated with higher death hazards after hospitalisation due to mental and behavioural disorders (HRs: 1.5–2.0).

DISCUSSION

Based on a 1953 birth cohort from Stockholm, followed over almost seven decades, our findings could largely confirm the initial hypotheses. Individuals with experiences of childhood

Table 3 The association between ICWS and all-cause mortality after hospitalisation, stratified by sex and cause group

	Observations/deaths	Men (n=7204)				Women (n=6930)				
		Investigated		Placed		Investigated		Placed		
		HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI	
External causes	2459/533	1.65	1.34 to 2.02	2.35	1.87 to 2.96	1989/291	2.21	1.54 to 3.16	2.55	1.90 to 3.44
Self-harm	247/116	1.59	1.00 to 2.51	1.74	1.09 to 2.78	257/77	1.10	0.54 to 2.23	1.58	0.91 to 2.73
NCDs	5033/871	1.69	1.43 to 1.99	2.33	1.95 to 2.79	5064/567	1.61	1.21 to 2.15	2.08	1.66 to 2.62
Mental and behavioural disorders	1146/435	1.48	1.17 to 1.88	1.56	1.23 to 1.99	867/208	1.34	0.87 to 2.06	2.00	1.46 to 2.75
Anxiety	251/81	1.20	0.66 to 2.18	1.23	0.70 to 2.16	258/51	0.44	0.10 to 1.90	1.83	0.94 to 3.56
Depression	283/89	1.70	0.96 to 3.02	1.80	1.08 to 2.99	313/70	0.85	0.33 to 2.16	1.64	0.94 to 2.87
Substance use disorders	757/342	1.09	0.83 to 1.42	1.29	0.97 to 1.70	419/133	1.32	0.79 to 2.18	2.03	1.37 to 3.02
CVDs	2195/376	1.75	1.37 to 2.23	2.49	1.88 to 3.30	1530/225	2.09	1.39 to 3.14	2.27	1.59 to 3.25
Cancers	848/295	1.56	1.17 to 2.08	1.52	1.08 to 2.15	1094/309	1.38	0.90 to 2.11	1.47	1.01 to 2.13
Respiratory	816/198	1.22	0.86 to 1.73	2.30	1.58 to 3.35	814/151	1.13	0.61 to 2.11	1.63	1.09 to 2.43
Diabetes	381/100	1.40	0.86 to 2.29	3.00	1.74 to 5.17	243/57	1.97	0.86 to 4.52	1.65	0.80 to 3.38

Results from Cox proportional hazards models.

Reference group is 'Without ICWS'. (HR=1.00). The models are centred around the first admission date and uses a calendar time scale. All models are adjusted for born out of wedlock, mother's age at birth, occupational status of the father at birth, and family education at age 10. Bold numbers signify a p value ≤ 0.004 .

CI, CI interval; CVDs, Cardiovascular diseases; HR, HR ratio; ICWS, involvement with child welfare services; NCDs, Non-communicable diseases.

adversity, by proxy of ICWS, had higher risks of being hospitalised in adulthood due to most of the major causes included here (differential risk of hospitalisation). Overall, they were also more likely to die after being hospitalised (differential survival). The results generally followed a gradient reflecting the severity of ICWS, which suggests that the results are driven by the underlying exposure to adversity rather than solely being due to placement in OHC itself. Moreover, there were no clear differences between men and women.

Extending the follow-up to mid-adulthood, our study corroborates findings from earlier studies showing that individuals with experiences of OHC have increased risks for hospitalisation due to external causes and mental and behavioural disorders—especially substance use disorders and self-harm.^{14 16} We further add evidence for elevated hospitalisation risks due to NCDs, and the subcategories of CVDs, respiratory diseases and diabetes. This is remarkable as such diseases typically have a later onset, and selective mortality among those with ICWS would be expected to bias these estimates towards the null. The investigation into subcategories also shows that the conditions most strongly related to health risk behaviours are particularly increased among those with experiences of OHC placement. This potentially indicates that such behaviours are perhaps used as strategy to cope with the increased burden of mental health problems in those with experiences of ICWS.

The findings also highlight that cancers are an exception, potentially because we could not differentiate the types of cancers with stronger behavioural risk factors. Our results are nonetheless in line with two of the most comprehensive earlier prospective inquiries into hospitalisation risks, which found a higher incidence across many conditions, but not neoplasms among individuals with a history of foster care.^{28 30} This is in contrast to other retrospective studies on cumulative adverse childhood experiences,² and to two prospective studies finding evidence for increased cancer risks among women and lung cancer risks among those with childhood adversity.^{34 35}

This study uniquely investigates the association between childhood adversity and survival prospects following hospitalisation. Individuals with experiences of ICWS generally had higher mortality risks and died sooner after hospitalisation due to external causes or NCDs than individuals without such experiences. These results were consistent in the subcategory of CVDs but not in the other subcategories. The worse survival prospects after a CVD-related hospitalisation could mean that individuals with experiences of ICWS are more vulnerable to the adverse consequences of such diseases, perhaps due to comorbidities, worse mental health or other coinciding risk factors for worse prognoses. Alternatively, it could reflect differences in disease severity when they present to the hospital, or differences in quality, access or response to treatment. These worse survival prospects might point to unmet care needs among individuals with experiences of childhood adversity; something which has been shown in OHC-experienced populations already during their childhoods.³⁶

This study has several strengths. Most importantly, the population-based birth cohort data that could be followed up to age 66. Based on national registers, there was minimal attrition and selection. Some limitations should, however, be considered. We cannot exclude the possibility of measurement error. Our indicators of ICWS were based on child welfare records from the Social Registers kept locally by municipalities in Stockholm; not all cohort members lived in Stockholm before age 10 and we might miss some cases that were solely recorded outside the metropolitan area.³⁷ In terms of hospitalisations, the Swedish

National Patient Register did not have full national coverage between 1973 and 1987.³⁸ However, the coverage in Stockholm was more or less complete (for a more extensive discussion, see Ludvigsson *et al*).

There are possible unmeasured confounders, such as baseline health, and other factors related to genetic predisposition to disease or risk-taking behaviours. There are also a number of potential mediators that have been not included in this study, for instance, life style behaviours, mental health, cognitive ability and educational attainment. Our results, nonetheless, lend some support to the hypothesis that mental and behavioural problems might be important explanatory factors. How much, for example, of the worse survival prospects, might be explained by mental health warrants further investigation: in patients with several comorbidities who experienced ICWS, mental health problems are very common (online supplemental table 5). Nevertheless, the cause groups used here were too broad and numerous to consider disease aetiology.

It remains a question whether our findings may explain previously reported inequalities in premature all-cause mortality between those with and without experiences of ICWS.⁷ Differential risk of hospitalisation and differential survival after hospitalisation have been considered important for producing socioeconomic inequalities in mortality more generally.³⁹ Further investigating the mechanisms driving the inequalities in mortality following childhood adversity is a task for future research to consider.⁴⁰

As this study only reports associations for one cohort born in one city, future research would need to verify the results in other contexts, and use designs and methods better equipped for making causal inferences. Regardless, the impact of adversity seems to persist across the life course, making it a relevant issue for public health and the inquiry into health inequalities. Our results further have implications for healthcare policy, as they strongly suggest that populations with experiences of childhood adversity might not be optimally served by the healthcare system, whether due to intrinsic susceptibility, help-seeking behaviours, comorbidity or other barriers. With their higher risks of disease and death following diseases, much could be gained from improving access to and quality of care in this vulnerable population.

Funding This study was supported by grants from the Swedish Research Council for Health, Working Life and Welfare (Forte), grant numbers 2016-07148 and 2019-00058. The funders had no role in the analyses interpretation of results or the writing of this manuscript.

Competing interests None declared.

Patient consent for publication Not applicable.

Ethics approval This study was approved by the Stockholm Regional Ethical Review Board (number 2017/34-31/5; 2017/684-32; 2011/1907-31/5).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Under Swedish law and ethical approval, patient level data cannot be made publicly available. The statistical code is available from the corresponding author. Data access can be granted upon request to YBA and conditional on approval of the review board.

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ORCID iDs

Josephine Jackisch <http://orcid.org/0000-0002-5958-2303>
Ylva B Almquist <http://orcid.org/0000-0001-7576-9410>

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