The European Journal of Public Health, Vol. 29, No. 3, 575–582 © The Author(s) 2018. Published by Oxford University Press on behalf of the European Public Health Association. This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/4.0/), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited. For commercial re-use, please contact journals.permissions@oup.com doi:10.1093/eurpub/cky143 Advance Access published on 6 August 2018

io.10.10.10.5, culpub, exp115 Advance Access published on 6 August 2016

Systematic Review and Meta-Analyses

The association between social exclusion or inclusion and health in EU and OECD countries: a systematic review

Addi P. L. van Bergen^{1,2,3}, Judith R. L. M. Wolf⁴, Mariam Badou⁵, Kimriek de Wilde-Schutten⁵, Wilhelmina IJzelenberg⁵, Hanneke Schreurs¹, Bouwine Carlier⁶, Stella J. M. Hoff⁷, Albert M. van Hemert²

- 1 Public Health, Municipality of Utrecht, Utrecht, The Netherlands
- 2 Department of Psychiatry, Leiden University Medical Centre, Leiden, The Netherlands
- 3 Department of Epidemiology, Health Promotion and Care Innovation, Public Health Service of Amsterdam, Amsterdam, The Netherlands
- 4 Impuls, The Netherlands Centre for Social Care Research, Radboud University Medical Centre, Nijmegen, The Netherlands
- 5 Department of Health Sciences, Faculty of Sciences, Vrije Universiteit Amsterdam, Amsterdam Public Health Research Institute, Amsterdam, The Netherlands
- 6 Research Group Occupation and Health, HAN University of Applied Sciences, Arnhem and Nijmegen, The Netherlands
- 7 Department of Income and Social Security, The Netherlands Institute of Social ResearchISCP, The Hague, The Netherlands

Correspondence: Addi P. L. van Bergen, Department of Psychiatry, Leiden University Medical Centre, Albinusdreef 2, 2333 ZA Leiden, The Netherlands, Tel:+31 06 28176202, e-mail: 888envanbergen@ziggo.nl

References 41-62 are given in Supplementary file S6.

Background: Social exclusion (SE), or the inability to participate fully in society, is considered one of the driving forces of health inequalities. Systematic evidence on this subject is pertinent but scarce. This review aims to systematically summarise peer reviewed studies examining the association between the multidimensional concepts of SE and social inclusion (SI) and health among adults in EU and OECD countries. Methods: The protocol was registered on Prospero (CRD42017052718). Three major medical databases were searched to identify studies published before January 2018, supplemented by reference and citation tracking. Articles were included if they investigated SE or SI as a multidimensional concept with at least two out of the four dimensions of SE/SI, i.e. economic, social, political and cultural. A qualitative synthesis was conducted. Results: Twenty-two observational studies were included. In the general population, high SE/low SI was associated with adverse mental and general health. For physical health, the evidence was inconclusive. In groups at high risk of SE, support was found for the association between high SE/low SI and adverse mental health but no conclusions could be drawn for physical and general health. Conclusions: This review found evidence for the association between high SE/low SI and adverse health outcomes, particularly mental health outcomes. The evidence is mainly based on crosssectional studies using simple and often ad hoc indicators of SE/SI. The development and use of validated measures of SE/SI and more longitudinal research is needed to further substantiate the evidence base and gain better understanding of the causal pathways.

.....

Introduction

According to the World Health Organization (WHO), social exclusion (SE) is one of the driving forces of health inequalities.^{1–3} SE refers to the inability of people to participate fully in society,⁴ while its antipode, social inclusion (SI) refers to the situation in which individuals are fully involved in the society in which they reside, including the economic, social, cultural and political dimensions of that society.⁵ The pathways linking SE (and lack of SI) to poor health are complex and diverse.¹ The situation of SE encompasses deprivations in areas such as social relations, material resources, access to health services and housing, which are in itself well known determinants of health.^{6,7} In addition, pathways leading to poor health may occur via direct and indirect causation as well as through reverse causation. The experience of exclusion, e.g. low social standing, feelings of alienation and lack of belongingness may directly impact health and well-being via psychoneuroendocrine mechanisms or work indirectly through stressrelated unhealthy behaviours.^{8–10} SE may also give cause to other deprivations, e.g. poor labour conditions or poor nutrition, which also contribute to ill-health.¹ Reverse causation occurs when poor health and disability generate and reinforce exclusionary processes.²

Although SE and SI have considerable public health significance from a theoretical perspective, the empirical evidence-base on this topic is still sparse. Literature reviews on social exclusion or inclusion and health mostly discuss the concepts, operationalisations and instruments used to measure SE or SI^{2,11–13} or describe characteristics of the retrieved studies (research design, country, year of publication etc.).^{14,15} One study systematically reviewed the impact of interventions on SI in adults with intellectual disability.¹⁶ None of these reviews reported systematically on the relationship between SE/ SI and health. The lack of clarity and diversity of meanings associated with SE/SI, the wide variety of SE/SI measures used, the focus on only one dimension of SE/SI and the complexity and sheer magnitude of the literature, severely limited the inferences that could be made from these studies.^{2,11–16}

In this study, we conducted a systematic literature review that addressed one of the main obstacles encountered in previous reviews, i.e. lack of clarity and diversity of meanings. SE/SI represents a broad concept that, by its nature, can be defined and operationalised in various ways. Multidimensionality is one of the agreed upon characteristics,^{17'} but the number and nature of the dimensions vary. Burchardt, for example, used four dimensions: consumption, production, political engagement and social interaction;¹⁸ others distinguished six¹⁹ or even seven²⁰ elements or dimensions of SE/SI. The WHO defines social exclusion as 'dynamic multidimensional processes driven by unequal power relationships interacting across four main dimensions-economic, political, social and cultural-and at different levels including individual, household, group, community, country and global levels'.1 These processes may lead to a state of SE characterised by a cumulation of deprivations in multiple dimensions.^{1,10,21} We choose the WHO definition and classification into four societal domains as a template for our study. To further improve homogeneity we made a distinction between often large general population studies and smaller studies in specific groups, mostly at high risk of SE.

The purpose of this study is to systematically summarise the evidence on the association between multidimensional SE and health and to evaluate six hypotheses, i.e. that high SE/low SI is associated with: (i) adverse mental health, (ii) adverse physical health and (iii) adverse general health outcomes in (a) the general population and (b) populations at high risk of SE.

Methods

We followed PRISMA guidelines for reporting this systematic review.^{22,23} The review protocol is registered on the PROSPERO database (registration number CRD42017052718) and is available at https://www.crd.york.ac.uk/PROSPERO.

Electronic search

We developed and executed, with the help of a qualified librarian, a search strategy to identify all studies that reported the association between SE/SI and health. The following three major electronic health databases were searched up to January 2018: PubMed, EMBASE and CINAHL. Searches were conducted in March 2015 and January 2018. The terms 'social exclusion' and 'social inclusion' were searched in title and abstract. Search strategies can be found in Supplementary file S1.

Study selection

Two authors (AvB, plus MB or KS) independently screened all records identified by the electronic search on title and abstract using a sequentially applied algorithm previously introduced by Curran et al.¹⁴ (figure 2). First, records without an abstract and inconclusive title were moved to a separate database to be assessed on the basis of full text. Next, language, study population, country and type of publication/ study design were checked. Studies had to be written in English, Dutch, German, Spanish, or French; involve an adult population; and be set in EU-countries or OECD-countries (http://www.oecd.org/about/ membersandpartners/list-oecd-member-countries.htm), because of their relatively similar welfare regimes. Only research articles published in peer-reviewed journals describing quantitative studies were included. Articles had to test the relationship between SE/SI and a health measure and report statistical results. Next, we excluded studies not using a multidimensional construct of SE/SI (minimum two of four dimensions), studies in which health formed part of the SE/SI measure and studies using an ecological measure of SE/SI. We did not exclude studies on sample size criteria.

We retrieved full-texts of all articles considered potentially eligible by at least one reviewer. Two reviewers (AvB, plus MB, KS or BC) then independently assessed the full texts to ascertain that the inclusion criteria were met. In case of disagreement, one of the other reviewers was consulted to decide. To complement the electronic searches, we hand-searched the reference lists of included studies and other reviews. Citation tracking was performed using Web of Science (WoS) or Google Scholar if studies were unavailable in WoS. Studies identified through reference and citation tracking were screened and assessed by AvB. When uncertain, BC or HS were consulted.

Data extraction

For each included study, the following data were extracted: study design, country, study population, sample size, dimensions and measures of SE/SI, health measure(s), confounding variables, statistical analysis and key results. We classified the health outcomes into three groups: mental health related (MH), physical health related (PH) and general health related (GH). In this, we were guided by the lists of mental and physical adult health measures in the Patient-Reported Outcomes Measurement Information System (PROMIS http://www.healthmeasures.net/explore-measurement-systems/

promis/intro-to-promis/list-of-adult-measures) and UK experience on general health measures.²⁴ Mental illness and its impacts, emotional distress and cognitive functioning were classified under MH, as were intravenous drug use and compulsory hospitalisation on grounds of health and safety risks due to mental illness. Physical functioning, impairments and symptoms such as headache and sleeplessness, were classified under PH. Whereas physical health refers to the physiologic and physical status of the body, general health refers to overall health status. Typical general health measures are self-rated health, presence of chronic diseases (yes/ no) and limitations due to health problems (yes/no).²⁴

Indicators of SE/SI were classified into the four WHO dimensions of SE/SI: social (S), economic (E), political (P) and cultural (C) as operationalised by the Netherlands Institute for Social Research|SCP.^{3,25,26} In the social dimension, we classified SE/SI indicators relating to social isolation, participation in formal and informal social networks and social involvement. In the economic dimension, we classified SE/SI indicators relating to deficits that people experience as shown by debts and the absence of certain basic goods and services; in the political dimension we classified indicators on the ability to exercise the rights people normally have, such as adequate health care, sufficient education, proper housing, a safe living environment and access to public and commercial services; and finally, in the cultural domain, we categorised indicators referring to a lack of normative integration, i.e. non-compliance with core values of society such as low work ethic, low training readiness, not voting, social security abuse or delinquent behaviour. Study populations were classified into two groups: (i) general population and (ii) population groups at high risk of SE. Studies among adults in HIV treatment, problematic drug users and single mothers were classified in the latter category. We classified elderly as general population, viewing them as a demographic group rather than a high-risk group. Data extraction was performed by one reviewer (AvB, MB or KS) and checked by a second (AvB, BC or HS).

Risk of bias assessment

As there is currently little consensus on the critical elements for assessing risk of bias in observational studies,²⁷ we opted for a two-track approach. The general methodological quality of each

study was evaluated independently by two reviewers (AvB and MB KS BC or HS) using the Critical Appraisal Skills Programme (CASP) tools for cross-sectional and cohort studies (Supplementary files S2-S3). The respective CASP checklists consist of 10 and 11 questions (e.g. 'Was the outcome accurately measured to minimise bias?' and 'Was the cohort recruited in an acceptable way?'), that can be answered with: 'yes' (1 point), 'can't tell', or 'no' (0 points). The option to answer 'yes moderately' (0.5 points) was added by the reviewers. Disagreements were resolved through consensus and, if necessary, a third reviewer was consulted (BC). A commonly used cut-off point of 60% was used to distinguish between low and acceptable quality studies.²⁸ Only acceptable quality studies were included in the synthesis. As done by De Silva et al.,²⁹ we assessed, in addition to the CASP, a number of specific methodological limitations with a high risk of bias for our research question. We examined whether the definition, operationalisation and measurement of SE/SI were adequately substantiated, whether testing of the association between SE/SI and health was a stated objective of the study and whether adjustment for confounding factors was performed. Details can be found in Supplementary file S4.

Data analysis

Given the variation in health measures and study designs, it was not possible to conduct a meta-analysis. Instead, we used the method of grouping results as originally described by Ramirez et al.³⁰ To examine the six research hypotheses, we grouped the results for each hypothesis into four qualitative patterns. These were: (i) positive, when a significant (P < 0.05) concordant relationship was found for all measured SE/SI dimensions (high-SE/low SI corresponds to low health outcome), (ii) negative, when an inverse association was found, (iii) no association, when the relationships between the SE/SI dimensions and health were not statistically significant and (iv) partly (+/0), when studies reported multiple associations. We classified the result as partly when 30-70% of the tested relations were positive and the remaining 70-30% not significant. If studies reported findings for multiple, non-overlapping, research groups, e.g. men and women, these were included separately in the data analysis and counted as separate instances. When both

unadjusted and adjusted results were presented, only adjusted results were reported. Results were combined by counting the number of instances in each category and weighting by sample size.

Results

Study selection

The digital search yielded 4032 non-duplicated articles: 2038 references in PubMed, 1219 in EMBASE and 775 in CINAHL (figure 1). On the basis of title and abstract screening, 3847 articles were excluded (figure 2). The most common reason for exclusion was publication type (editorials, posters etc.) and study design. Articles that were excluded on language were mostly written in Portuguese. In total 185 articles were selected for full-text screening of which 19 met the inclusion criteria. An important reason for exclusion in this stage was the use of the term SE or SI for a single dimension of SE/SI or for a different concept (e.g. exposure to forms of mistreatment, problems with daily activities or fear for SE). The interrater agreement for the selection of the publications was good (Cohen's $\kappa = 0.77^{31}$). Through reference and citation tracking 1792 more papers were identified of which three met the inclusion criteria. Main reasons for exclusion in this stage were subject (58%) and publication type (22%).

Description of studies

The studies included five cohort studies, four case control studies and 13 studies with a cross-sectional design. The majority of the studies were conducted in Europe (15), mostly in England (6). Eleven studies were conducted in the general population and 11 investigated SE/SI in groups at high risk of SE. The sample sizes ranged from 67 to 25 498 participants. Sixteen papers addressed mental health (MH), six physical health (PH) and six general health (GH). Five papers addressed more than one type of health outcome. The studies are presented in Supplementary file S5, tables S1a and S1b for MH, tables S2a and S2b for PH and tables S3a and S3b for GH.





Figure 2 Exclusion algorithm title and abstract screening

	Mental health													Physical health							General health							
	general population							high-risk groups									general population					high-risk	ge	general population			on	high-risk
	Sacker et al. (41)	Bayram et al. (42)	Halleröd & Larsson (43)	Honey et al. (44)	Richter & Hoffmann (45)	Van de Beek et al. (46)	Flores et al. (47)	Todd et al. (48)	Webber & Huxley (49)	Choi et al. (50)	Cole et al. (51)	Fakhoury & Priebe (53)	Killaspy et al. (53)	Maia et al. (54)	March et al. (55)	0' Brien et al. (56)	Saito et al. (57)	Waterstone et al. (58)	Bayram et al. (42)	Halleröd & Larsson (43)	Richter & Hoffmann (45)	O' Brien et al. (56)	Bryngelson (59)	Gannon & Nolan (60)	Sacker et al. (41)	Halleröd & Larsson (43)	Urbanos-Garrido (61)	Johner et al. (62)
1, Theoretical motivation SE/SI	٠	٠	٠	•	٠	•	٠	•	•	•	٠	•	٠	•	•	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠
2, Data set originally designed to measure SE/SI	•	٠	•	•	•	٠	٠	٠	•	٠	٠	•	٠	٠	٠	•	•	•	٠	•	•	•	•	٠	•	•	٠	٠
3, All dimensions of SE/SI measured	•	٠	٠	•	•	•	٠	٠	•	•	•	•	٠	•	•	•	•	٠	٠	٠	•	٠	•	•	•	٠	•	٠
4, Composite measure SE/SI	٠	•	•	•	•	•	٠	•	٠	٠	•	•	•	<	•	٠	٠	٠	•	•	•	٠	<	•	٠	•	•	
5, Existing SE/SI measure	•	٠	•	<	•	<	•	•	C	<	•	•	٠	•	•	•	•	•	٠	•	•	•	•	•	•	•	•	•
6, Testing of association SE/SI - health was a stated objective	٠	٠	•	•	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	٠	•	٠	٠	٠	٠	٠	•	٠	٠
7, Adjustment for confounding	٠	<	•	٠	<	<	•	٠	٠	٠	٠	٠	٠	٠	٠	<	٠	٠	<	•	<	•	<	٠	٠	•	٠	٠
Key: • Limitation not present;	•	Lim	itatic	on pa	rtly i	orese	nt;	Li	nitat	ion p	orese	nt																

Figure 3 Summary of study specific limitations with a high risk of bias

Risk of bias assessment

The methodological quality of each study is summarised in Supplementary file S5. Details on the measurement of SE/SI and confounding variables can be found there as well. All studies had CASP scores of 6 or more, indicating acceptable quality. Figure 3 shows that the most frequent methodological limitations were incomplete measurement of SE/SI (<4 dimensions) and lack of an existing SE/SI measurement instrument. Most studies used self-chosen indicators without testing the psychometric properties. Only two studies used an existing instrument for SE/SI, that is, the Social Inclusion Questionnaire User Experience (SInQUE)^{32,53} and Social exclusion index.^{33,42} Three studies used existing scales to measure dimensions of SE/SI.^{44,46,51} In two studies an index of social exclusion was constructed and partly validated.^{49,50} The majority of studies did not use a composite measure for SE/SI, and those that did, mostly calculated simple sum scores.^{41,47,49,54,57–59} In 10 studies, the data were not originally designed to measure SE/SI (e.g. case files, registration or monitoring data). One in three studies lacked a theoretical underpinning of SE/SI. Control for confounding factors was missing or incomplete in 7 of the 22 studies.

Mental health in the general population

Figure 4 shows evidence in favour of our hypothesis that high SE/ low SI is associated with adverse health outcomes for MH in the general population. Our hypothesis is supported by 92% of the combined sample (27881 persons, six instances, five studies)^{41,43-46} and partly supported by 8% of the sample (2493 persons; one instance).⁴² All but one study were cross-sectional in design. A retrospective cohort study showed an association between high psychological distress in elderly persons and later SE. High levels of SE, in turn, were found to be predictive of high psychological distress.⁴¹ Three cross-sectional studies found positive associations between a



Figure 4 Significance and direction of the relationship between SE/SI and health: total sample size (X-axis) and number of instances (between brackets)

large number of SE indicators and self-reported anxiety and anguish,⁴³ common mental illness and severe mental illness;⁴⁵ depressive symptoms and psychotic experiences.⁴⁶

Another supportive study⁴⁴ found that the relationship between disability and MH was moderated by the social and economic dimensions of SE (operationalised as low social support and financial hardship, respectively); and that the combination of the two dimensions strengthened the effect. The study with partial evidence⁴² found a significant relation between low MH and the social dimension of SE but not with the cultural and economic dimension. Within the political dimension one indicator (adequate housing and safe neighbourhood) showed a concordant relation with MH whereas the other did not (access to institutions).

Mental health in high-risk groups

Figure 4 shows that the association between SE/SI and MH was tested in 13 high-risk study populations. Due to the typically small samples, the total sample size is modest compared to the general population sample (figure 4; Supplementary file S5 tables S1a and S1b). This does not indicate less evidence per se. Our hypothesis was supported by 80% of the combined sample (4646 persons; 8 out of 13 instances) and partly supported by 12% of the sample (692 persons; three instances). Supporting evidence was derived from two case control studies^{47,48} and five cross-sectional studies.^{50,51,54–} ⁵⁶ The case control studies showed an elevated prevalence of DSM III personality features associated with SE in men with AIDS and/or drug addiction;47 and an elevated prevalence of substance use disorders in clients of mental health services with SE characteristics.48 The cross-sectional studies found significant associations between SE/SI and, respectively, perceived stress in patients in substance abuse treatment;⁵¹ elevated intravenous drug use in drug users in public places;⁵⁵ symptoms of depression⁵⁴ and mental symptoms and impairments in HIV patients⁵⁶ and higher levels of complex post-traumatic symptoms in torture survivors.⁵⁰

Partial evidence was found in a study among patients of Assertive Outreach teams.⁵² In this population, alcohol abuse and dependency was associated with the social and cultural dimensions of SE, but not with the political dimension. Drug abuse and dependency was associated with the political and part of the cultural dimension of SE and not with the social dimension. Partial evidence was also found by Killaspy et al.⁵³ Patients interviewed after developing a psychotic illness showed a significant deterioration in two of the four SI dimensions measured, i.e. the social and economic dimensions. Older age at onset of illness and longer duration of illness were associated with greater changes in the economic dimension. Higher current quality of life was associated with less decline in the social dimension.

Our hypothesis was not supported by two case control studies (490 persons, two instances).^{48,49} One study found that in clients

with substance use disorder, the co-occurrence of mental health problems was not associated with higher levels of SE.⁴⁸ The authors suggest that the association between substance abuse and SE is stronger than between mental health and SE. The second study⁴⁹ showed that SE increased the likelihood of compulsory admission among people assessed under the Mental Health Act, but, when other factors such as diagnosis, life-threatening self-neglect and physical aggression towards others, were taken into account, the association became non-significant. It is plausible that these factors might act as mediators in the relation between SE and compulsory admission.

Physical health in the general population

Figure 4 shows a more mixed picture for PH in the general population. Two studies support the hypothesis that high SE is associated with adverse PH (56% of the combined sample, 21 058 persons), two studies partly support the hypothesis (33%, 7879 persons) and two studies do not (21%, 9001 persons). Findings from a prospective cohort study57 showed that elderly Japanese women who were excluded both in the social and in the economic dimension were 1.7 times more likely to die prematurely than those who were not socially excluded. In elderly men, the association between SE and mortality was not significant. The results were adjusted for age, marital status, education, municipality, disease and impairment. Supporting evidence was also found from crosssectional studies on severe obstetric complications in general, on severe pre-eclamptic conditions and severe haemorrhage specifically,⁵⁸ on headache and sleeplessness⁴³ and severe physical illness or disability.45 No significant associations were found with severe haemorrhage and uterine rupture,⁵⁸ with obesity,⁴³ and with the PH domain of the WHOQOL-BREF.⁴² This domain covers among others pain, physical problems, sleep and energy.

General health in the general population

Evidence was found for the association between high SE/low SI and adverse GH in the general population. Our hypothesis was supported by 80% of the combined sample (32 537 persons, three out of six instances),^{41,60,61} partly supported by 16% of the sample (6481 persons; two instances),^{43,59} and not supported by 4% of the sample (1604 persons; one instance).⁵⁹ The results were heavily influenced by one large cross-sectional study in 25 498 adults in Spain which found significant relationships between SE factors and socioeconomic inequalities in self-assessed health, presence of any chronic disease and limitations in daily activity due to health problems.⁶¹ Two cohort studies showed positive associations between SE and disability onset and persistent disability,^{41,60} and onset and persistence of low self-assessed health. A third cohort study showed partial evidence.⁵⁹ In women, long-term sickness

absence adjusted for age and previous SE increased the risk of the combination of economic and social exclusion, but not of the combination of economic and cultural exclusion. In men, no significant associations were found between dimensions of SE and long-term sickness.⁵⁹ Partial evidence was also found in a cross-sectional study among 4941 adults demonstrating a positive association between the presence of any chronic disease and the social, economic and part of the political dimensions of SE/SI, but not with the cultural dimension.⁴³

Physical health and general health in high-risk groups

The literature did provide little evidence on the association between SE/SI and PH or GH in high-risk groups. The number of studies was low, with one study on PH⁵⁶ and one study (2 instances) on GH.⁶² In HIV patients, physical symptoms and impairments and difficulties in day-to-day activities due to illness were associated with low SI.⁵⁶ In single mothers, self- assessed health was not associated with SE. This was true for both single mothers on social assistance as for single mothers without social assistance.⁶²

Discussion

We set out to systematically summarise existing evidence on the association between SE/SI and health and evaluate the hypotheses that high SE/low SI is associated with adverse MH, PH and GH outcomes, in the general population and in groups at high risk of SE. The evidence base is currently strongest for the association between SE/SI and MH. The hypotheses that high SE/low SI is associated with adverse MH outcomes are supported by studies with various designs, sample sizes and settings, in both the general population and high-risk groups. Conflicting evidence was only found in two studies,^{48,49} in which the relation between SE/SI and MH appeared to be mediated by other factors.

This review also found support for the association between SE/SI and GH in the general population. The outcomes included some that are widely used in public health monitoring such as self-assessed health, presence of any chronic disease and limitations due to health problems. Two aspects deserve closer attention. First, the results are confined to the social and economic dimensions of SE/SI. The cultural and social rights dimensions were not well presented and little or no significant relations with these dimensions were found. Second, none of the studies used a composite measure for SE/SI, and only one study provided insight into the cumulative impact of the underlying dimensions.⁵⁹

Our review failed to confirm or refute a direct association between high SE/low SI and adverse PH in the general population. The wider literature provides ample evidence for associations between aspects of SE/SI and PH outcomes, for example, between social relations and mortality⁶ and between neighbourhood characteristics and cardiovascular health.⁷ We expected that a cumulation of these aspects would also be associated with adverse PH outcomes. One reason for the absence of association may be the much broader spectrum of PH outcomes included in this study, ranging from headache and obesity to severe obstetric complications. Another reason may be that these studies use other terms such as deprivation or precariousness and did not get included in this review.

Lastly, as our review identified only a few studies focusing on the relation between SE/SI and PH or GH in high-risk populations, no conclusions can be drawn about the hypotheses on PH and GH in high-risk groups.

Causality and directionality

The studies we found employ different assumptions about the relationship between SE/SI and health. Some authors consider SE as a cause of adverse health^{47,57,58} while others regard SE as a consequence of adverse health^{53,56,60} or as a mediator.⁴⁴ The observational design of these studies does, however, preclude firm causal inference. The few longitudinal studies give us some insight in directionality. One longitudinal study showed that SE preceded negative health outcomes, i.e. mortality in Japanese elderly women.⁵⁷ A second longitudinal study⁵⁹ points to a reverse directionality; long-term sickness absence was associated with a deterioration of the economic and social dimensions of SE in women, independent of their earlier situation. A reciprocal relation was found in two longitudinal studies.^{41,60} Further longitudinal studies may contribute to unravel the dynamic relation between SE/SI and health.

Risk of bias within studies

In line with previous reviews we found almost no study using a valid measure for SE/SI. Most studies used self-chosen indicators and in nearly half of the studies the data were not originally designed to measure SE/SI. The lack of valid measures for SE/SI prevents very firm conclusions being drawn from this review. We agree with previous reviews that the development and use of validated multi-dimensional measures^{12,13,16,34} is warranted in future research. As SE is a multi-interpretable concept that can be operationalised in various ways depending on one's theoretical perspective, political position and purpose, it is not obvious that one agreed upon measure for SE/SI will surface. This need not be problematic as long as choices are explicated and substantiated.

Finally, seven of the 22 studies did not adjust for demographic and other potential confounding factors. As confounding may affect the results of our review through over-estimation, the evidence was also analysed without these seven studies and the inferences remained unchanged. It is important to note that in all observational studies, residual confounding may account for part of the associations observed.

Strengths and limitations of this review

The principal strengths of this review are its systematic approach, tactical search strategy and clear conceptual framework. These made it possible, despite the great diversity of studies, to take a step further than previous reviews, which did not report on the relationship between SE/SI and health, or did not do this systematically, but merely as exemplary descriptions.^{2,12–16} Another strong points is the inclusion of papers in languages other than English.

There are limitations too. The method we used to summarise the evidence is based on P-values. P-values give an indication of the compatibility of the data with the null-hypothesis of each paper, and not of the effect size or the importance of the results.³⁵ To enable interpretation of the results we reported for each paper effect sizes and/or other statistics in the tables and provided some qualitative context in the main text. Another limitation arises from the classification of health outcomes, which was not always straightforward. In a number of studies no clear distinction could be made between MH and PH components, for instance, when researchers considered other, non-congruent, classes of diseases. As these results were classified as GH, this category may have become somewhat ambiguous. Yet another limitation is that our review is not exhaustive. The downside of applying strict selection criteria is that, e.g. studies not using a multidimensional construct of SE/SI were left out. As a previous review using a comprehensive search strategy yielded unmanageable amounts of 100 000 plus titles,¹⁴ we choose a narrow search strategy to identify papers that focus specifically on SE/SI and not on related subjects such as income, housing, social cohesion etc. This way we may have missed relevant papers not using the specific terms SE or SI in title or abstract, but we do not expect there to be many, for two reasons. First, studies on only one dimension of SE/SI or on a constituent element, such as social isolation, housing or access to health care, are beyond the scope of this review as our focus was solely on papers employing a multidimensional interpretation of the concept SE/SI. Second, more comprehensive concepts such as social cohesion, social capital, citizenship or (multiple) poverty, were deliberately kept out of the

Implications for future research

Our paper revealed a great number of weaknesses in research methodology and provides ideas and directions for future research. A research agenda required to have a better understanding of potential mechanisms and putative pathways should include longitudinal studies, studies into mediating and modifying factors such as gender and previous disadvantage; and into the accumulation and interaction of SE/SI dimensions. Equally important for enhancing the knowledge base on SE/SI and health, is a more systematic and standardised terminology of SE/SI domains and the development and validation of composite measures of SE/SI. The WHO/SCP model used in this paper may serve here as a useful template.^{3,25,26}

Policy implications

The association between high SE and poor MH came most clearly to the fore in people with severe mental illness and substance use disorder. Through the implementation of recovery-orientated services, the mental health sector can contribute to the SI of their clients.³⁶ but more may be needed. As several studies in our review show an association between the economic and political dimensions of SE/SI and MH, e.g. with income, economic deprivation, employment, education and housing, we expect that there may be little chance of improving the situation of those with mental health problems without attention being given to these other problems. These need to be addressed by social and economic policies,³⁶ involving not just the health sector but a range of sectors and services such as housing, employment, education, income support, debt counselling and community building.³⁷⁻⁴⁰ The evidence on the association between SE/SI and poor mental and general health in the general population also calls for more macro level policies and interventions, targeting the general population and not only those at highest risk.

Conclusions

To our knowledge, this is the first paper that succeeded in systematically synthesizing evidence on the association between the multidimensional concept of SE/SI and health. Preliminary evidence was promising. Most studies confirmed the expected relationship between high SE/low SI and adverse health outcomes, particularly for mental health. We recommend a greater focus on the valid measurement of SE/SI in future research.

Supplementary data

Supplementary data are available at EURPUB online.

Conflicts of interest: None declared.

Key points

- Social exclusion is generally regarded as an important social determinant of health, yet, its evidence base is still weak.
- In this systematic review, we operationalised social exclusion as the cumulation of deprivations in four dimensions, i.e.

social, economic, political and cultural and social inclusion as full involvement in these dimensions.

- Evidence was found for the interconnectedness of social exclusion and inclusion and health. Available evidence is stronger for mental and general health than for physical health.
- There is need for the development and use of validated multidimensional, and preferably composite, measures for social exclusion and inclusion.

References

- Popay J, Escorel S, Hernández M, et al. Understanding and Tackling Social Exclusion. Final Report to the WHO Commission on Social Determinants of Health from the Social Exclusion Knowledge Network. Geneva: World Health Organization, 2008.
- 2 Mathieson J, Popay J, Enoch E, et al. Social Exclusion. Meaning, Measurement and Experience and Links to Health Inequalities. A Review of Literature. Geneva: WHO Social Exclusion Knowledge Network, 2008.
- 3 Poverty, Social Exclusion and Health Systems in the WHO European Region. Copenhagen: WHO Regional Office for Europe, 2010.
- 4 Millar J. Social exclusion and social policy research: defining exclusion. In: Abrams D, Christian J, Gordon D, editors. *Multidisciplinary Handbook of Social Exclusion Research*. Chichester: John Wiley & Sons Ltd., 2007: 1–16.
- 5 Standing Senate Committee on Social Affairs Science and Technology. In from the Margins, Part II: Reducing Barriers to Social Inclusion and Social Cohesion. Ottawa, Ontario, Canada: Senate, 2013.
- 6 Courtin E, Knapp M. Social isolation, loneliness and health in old age: a scoping review. Health Soc Care Community 2017;25:799–812.
- 7 Gomez SL, Shariff-Marco S, DeRouen M, et al. The impact of neighborhood social and built environment factors across the cancer continuum: current research, methodological considerations, and future directions. *Cancer* 2015;121:2314
- 8 Marmot M. Status Syndrome: How Your Social Standing Directly Affects Your Health. London: Bloomsbury Publishing Plc., 2004.
- 9 Wilkinson R, Pickett K. The Spirit Level: Why More Equal Societies Almost Always Do Better. London: Penguin Books Ltd., 2010.
- 10 Sen A. Social Exclusion: Concept, Application, and Scrutiny. Manila, Philippines: Asian Development Bank, 2000.
- 11 Huxley P, Evans S, Munroe M, et al. Development of a Social Inclusion Index to Capture Subjective and Objective Domains (Phase I). Birmingham, UK: National Coordinating Centre for Research and Methodology, 2006.
- 12 Morgan C, Burns T, Fitzpatrick R, et al. Social exclusion and mental health: conceptual and methodological review. *Br J Psychiatry* 2007;191:477–83.
- 13 Wright N, Stickley T. Concepts of social inclusion, exclusion and mental health: a review of the international literature. J Psychiatr Ment Health Nurs 2013;20:71.
- 14 Curran C, Burchardt T, Knapp M, et al. Challenges in multidisciplinary systematic reviewing: a study of social exclusion and mental health policy. *Soc Policy Admin* 2007;41:289–312.
- 15 Evans-Lacko S, Courtin E, Fiorillo A, et al. The state of the art in European research on reducing social exclusion and stigma related to mental health: a systematic mapping of the literature. *Eur Psychiatry* 2014;29:381–9.
- 16 Bigby C. Social inclusion and people with intellectual disability and challenging behaviour: a systematic review. J Intellect Dev Disabil 2012;37:360–74.
- 17 Tsakloglou P, Papadopoulos F. Aggregate level and determining factors of social exclusion in twelve European countries. *J Eur Soc Policy* 2002;12:211–25.
- 18 Burchardt T, Le Grand J, Piachaud D. Degrees of exclusion: developing a dynamic, multi-dimensional measure. In: Hills J, Le Grand J, Piachaud D, editors. Understanding Social Exclusion. Oxford: Oxford University Press, 2002.
- 19 Secker J, Hacking S, Kent L, et al. Development of a measure of social inclusion for arts and mental health project participants. J Ment Health 2009;18:65–72.
- 20 Levitas R, Pantazis C, Fahmy E, et al. *The Multi-Dimensional Analysis of Social Exclusion*. Bristol: University of Bristol, 2007.
- 21 Vrooman J, Hoff S. The disadvantaged among the Dutch: a survey approach to the multidimensional measurement of social exclusion. Soc Indic Res 2013;113:1261–87.
- 22 Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med* 2009;6:e1000097.

- 23 Welch V, Petticrew M, Petkovic J, et al. Extending the PRISMA statement to equityfocused systematic reviews (PRISMA-E 2012): explanation and elaboration. Int J Equity Health 2015;14:92.
- 24 Meltzer H. General measures of health for use in health interview surveys and censuses: the UK experience. In: Ottawa, Canada: Proceedings of the Washington City Group meeting, 2003.
- 25 Hoff S, Vrooman C. Dimensies Van Sociale Uitsluiting: Naar Een Verbeterd Meetinstrument [Dimensions of Social Exclusion: Towards an Improved Measurement Instrument]. The Hague: The Netherlands Institute for Social Research|SCP, 2011.
- 26 Jehoel-Gijsbers G. Sociale Uitsluiting in Nederland [Social Exclusion in The Netherlands]. the Hague. The Hague: The Netherlands Institute for Social Research|SCP, 2004.
- 27 Jarde A, Losilla J-M, Vives J. Methodological quality assessment tools of non-experimental studies: a systematic review. Ann Psychol 2012;28:617-28.
- 28 van der Noordt M, IJzelenberg H, Droomers M, Proper KI. Health effects of employment: a systematic review of prospective studies. Occup Environ Med 2014;71:730-36.
- 29 De Silva MJ, McKenzie K, Harpham T, Huttly SR. Social capital and mental illness: a systematic review. J Epidemiol Community Health 2005;59:619-27.
- 30 Ramirez AJ, Westcombe AM, Burgess CC, et al. Factors predicting delayed presentation of symptomatic breast cancer: a systematic review. Lancet 1999;353:1127-31.
- 31 Altman D. Inter-rater agreement. Pract Stat Med Res 1991;5:403-9.

- 32 Mezey G, White S, Thachil A, et al. Development and preliminary validation of a measure of social inclusion for use in people with mental health problems: the SInQUE. Int J Soc Psychiatry 2013;59:501-7.
- 33 Jehoel-Gijsbers G, Vrooman C. Explaining Social Exclusion: A Theoretical Model Tested in The Netherlands. The Hague: The Netherlands Institute for Social Research|SCP, 2007.
- 34 Baumgartner JN, Burns JK. Measuring social inclusion-a key outcome in global mental health. Int J Epidemiol 2013;43:354-64.
- 35 Wasserstein RL, Lazar NA. The ASA's statement on p-values: context, process, and purpose. The American Statistician 2016;70:129-33.
- 36 Boardman J. Social exclusion and mental health-how people with mental health problems are disadvantaged: an overview. Ment Health Soc Inclus 2011:15:112-21
- 37 Braveman PA, Cubbin C, Egerter S, et al. Socioeconomic disparities in health in the United States: what the patterns tell us. Am J Public Health 2010;100:S186-96.
- 38 Marmot M, Allen JA. Social determinants of health equity. Am J Public Health 2014:104:S517-9
- 39 Montez JK, Zajacova A. Trends in mortality risk by education level and cause of death among US white women from 1986 to 2006. Am J Public Health 2013:103:473-9
- 40 Neumayer E, Plümper T. Inequalities of income and inequalities of longevity: a cross-country study. Am J Public Health 2016;106:160-5.

The European Journal of Public Health, Vol. 29, No. 3, 582-587

© The Author(s) 2018. Published by Oxford University Press on behalf of the European Public Health Association. This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http:// creativecommons.org/licenses/by-nc/4.0/), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited. For commercial re-use, please contact journals.permissions@oup.com doi:10.1093/eurpub/cky189 Advance Access published on 10 September 2018

Mortality after the death of a parent in adulthood: a register-based comparison of two ethno-linguistic groups

Jan Saarela¹, Mikael Rostila²

- 1 Demography Unit, Åbo Akademi University, FI-65100 Vaasa, Finland
- 2 Department of Public Health Sciences, Stockholm University and Centre for Health Equity Studies, Stockholm University/ Karolinska Institutet, SE-106 91 Stockholm, Sweden

Correspondence: Jan Saarela, Demography Unit, Åbo Akademi University, Strandgatan 2, FI-65100 Vaasa, Finland, Tel: +358 (0) 46 921 9438, e-mail jan.saarela@abo.fi

Background: Most research on parental bereavement and health have analysed health consequences of parental loss in childhood, while collateral health in adulthood has been less studied. Methods: Using register-based population data from Finland, we analyse adult offspring aged 18–50 years with discrete-time hazard models that adjust for offspring and parental socioeconomic and demographic characteristics. In focus are adult children whose parents were alive and lived together at the beginning of the observation period. We compare two culturally distinct but otherwise similar ethno-linguistic groups, Finnish speakers and Swedish speakers. Results: The results suggest that bereaved men have an approximately 30% higher death risk than nonbereaved men, while there is practically no difference in women. Associations between parental and child deaths are, as expected, stronger for concordant causes of death than for discordant causes of death. However, some associations for discordant causes of death remain, which may indicate causality. Among Swedish speakers, who have notably higher family stability than Finnish speakers, the death of one or both parents shows a stronger association with own mortality. Conclusions: The estimated associations found are generally larger than in the neighbouring country Sweden, which may be due to a stronger obedience to traditional family values and patriarchal family roles in Finland. These findings suggest that the association between parental death and mortality in adult offspring may depend on the societal context as well as on cultural practices. These factors should be increasingly acknowledged in future studies on collateral health.