**Title**: Perspectives on differing health outcomes by city: accounting for Glasgow’s excess mortality

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**Abstract**

Several health outcomes (including mortality) and health related behaviours are known to be worse in Scotland than in comparable areas of Europe and the United Kingdom. Within Scotland, Greater Glasgow (in West Central Scotland) experiences disproportionately poorer outcomes independent of measurable variation in socioeconomic status and other important determinants. Many reasons for this have been proposed, particularly related to deprivation, inequalities and variation in health behaviours. The use of models (such as the application of Bradford Hill’s viewpoints on causality to the different hypotheses) has provided useful insights on potentially causal mechanisms, with health behaviours and inequalities likely to represent the strongest individual candidates. This review describes the evolution of our understanding of Glasgow’s excess mortality, summarises some of the key work in this area and provides some suggestions for future areas of exploration. In the context of demographic change, the experience in Glasgow is an important example of the complexity that frequently lies behind observed variations in health outcomes within and between populations. A comprehensive explanation of Glasgow’s excess mortality may continue to remain elusive but is likely to lie in a complex and difficult-to-measure interplay of health determinants acting at different levels in society throughout the life course. Lessons learned from the detailed examination of different potentially causative determinants in Scotland may provide useful methodological insights that may be applied in other settings. Ongoing efforts to unravel the causal mechanisms are needed to inform Public Health efforts to reduce health inequalities and improve outcomes in Scotland.

**Introduction**

From a global perspective, the early part of the twenty first century has seen a significant turning point in population distribution. According to World Health Organisation (WHO) estimates, in 1990, less than 40% of the world’s population were living in cities compared to over half by 2010.1 This rapid transition is having fundamental implications for health and health inequalities. In Europe, unlike other major world regions, the overall population is predicted to fall.2 Low fertility rates, decreasing premature mortality rates, and flattening trends in immigration are leading to an older population profile.2 Along with these important demographic changes, wide variations are seen across Europe in health-related behaviours (such as smoking and alcohol consumption) and health outcomes (such as prevalence of long term conditions and mortality) both between and within countries. Life expectancy at birth and mortality rates, for example, have been shown to vary considerably between different European countries, regions and cities.3

Scotland’s relatively poor health profile, compared with other parts of the United Kingdom and with Europe, has been recognised for some considerable time. In 1989, Carstairs and Morris examined the difference between mortality experience in Scotland and England and Wales and suggested that the excess mortality in Scotland may be explained by more adverse conditions (ie greater deprivation) but were unable to demonstrate a causal relationship.4 Such observations led to the concept of a ‘Scottish effect’ in which Scotland was observed to experience an excess of poor health outcomes greater than that which could be explained by variation in common factors such as differences in age distribution.5 This led Public Health leaders to call for a ‘step change’ in societal efforts to improve health in Scotland.6 However, the optimal targeting of such efforts and their effectiveness in narrowing inequalities may have been somewhat hindered by uncertainty about true causal mechanisms.

Newer techniques, such as multilevel modelling, have allowed for better understanding of the nature of associations (such as that between deprivation and diet), but have also resulted in greater awareness of the lack of simplistic explanations. For example, improved ability to compare the relationship between Scottish health behaviours and associated outcomes has meant that the role of deprivation as the sole explanatory factor has been questioned.7 Similarly, better understanding of factors such as the apparent change in the influence of socioeconomic status over time (when compared with the rest of Great Britain) has revealed a greater underlying degree of complexity than was previously appreciated.8,9

Leading on from the identification of the variation between Scotland and the rest of the UK, the Glasgow Centre for Population Health (GCPH) has been in the forefront of work identifying that several health outcomes and health-related behaviours are worse in particular areas *within* Scotland, particularly West Central Scotland (including Greater Glasgow) and are improving at a slower rate than comparable, post-industrial regions in Europe and the rest of Scotland.8,10,11,12 Yet, despite a growing body of research in this area, a comprehensive explanation for Glasgow’s excess mortality has continued to prove elusive. This effect has sometimes been referred to as ‘the Glasgow effect’, although the term is considered by some to be unhelpful. The excess appears to be increasing over time, is seen for many different causes of death, is seen in comparison of all social classes (although for premature mortality, the excess is greatest in comparisons of those living in the poorest areas), and has been observed in all parts of Scotland when compared to other areas, not just Glasgow.

This review does not aim to represent a complete synopsis of all work in this area, but to describe some of the studies that have been key in shaping our understanding of the excess mortality in Scotland and, more specifically, in Glasgow, to summarize current thinking in the light of more recent work and to identify where uncertainties remain that may require future investigation.

**Deprivation and health in Scotland**

In 2005, Hanlon and colleagues published the findings of a cross sectional analysis of the British population using data from the 1981,1991 and 2001 censuses, demonstrating that, between 1981 and 2001, Scotland became less deprived relative to the rest of Great Britain but that age and sex standardised all-cause mortality rates were 12% higher in Scotland in 1981 and 15% higher in 2001.8 From their findings, they also suggested that measures of deprivation (using the Carstairs score – a measure of area-based deprivation based on four variables – adult male unemployment, lack of car ownership, low social class and overcrowding)13 did not explain most of the higher mortality in Scotland (Table 1).8

Subsequently, in 2007, analyses of combined data from Scottish Health Surveys in 1995, 1998 and 2003 (nationally representative population surveys, total n=25,127) showed that, by comparison with the rest of Scotland, men in Greater Glasgow had higher mortality rates from cancer, chronic liver disease and drug-related mental health disorders even after adjustment for area deprivation (also using the Carstairs index). In considering these findings, it is important to remember that identifying excess mortality in Glasgow relative to the rest of Scotland may be challenging, given that Scotland already exhibits an excess in relation to other areas. Higher rates of acute sickness, potential psychiatric morbidity and long standing illness (in men) also remained after adjustment for socioeconomic factors in this study as did some behavioural factors, such as poor diet (low green vegetable consumption) in men. Higher rates of excess alcohol consumption and binge drinking were found in both deprived and non-deprived areas of Glasgow compared with the rest of Scotland even after adjustment for age, survey year and socioeconomic status. Interestingly, differences in some other behavioural factors, such as smoking and using additional salt (in men) did appear to be explained by socioeconomic status.11

In 2008 Gray et al reported a similar study, but with comparison of Glasgow broadened to include a variety of European countries and regions (total n=101,923, of whom 1267 were participants of the 2003 Scottish Health Survey resident in Greater Glasgow). Again, the research question of interest was the degree to which socioeconomic factors (this time using occupational status and educational attainment as more internationally-comparable measures) explained differences in health behaviours and outcomes. The authors found that Greater Glasgow had a comparable socioeconomic profile in terms of occupation-based social class to many of its European counterparts, though conceded there were some differences in profile in terms of education status (with a higher proportion with no qualifications in Glasgow). After adjustment for age, social class and education qualification, binge drinking, smoking (in both men and women), obesity, diabetes, self-reported poor health, acute sickness, long standing illness and psychological morbidity tended to all be higher in Glasgow than in other regions, though the picture was slightly mixed (with some similarities with other areas for certain behaviours and outcomes). In this study they also concluded that the variations they had observed could not be explained solely by socio-economic deprivation.12

In 2009, Gray and Leyland then published a study looking specifically at variations in smoking behaviour. Using data from three Scottish Health Surveys again (1995, 1998 and 2003, total n=25,127), they compared smoking behaviour between Greater Glasgow and the rest of Scotland. For both men and women, the likelihood of smoking was considerably higher in Glasgow than in the rest of Scotland (odds ratios 1.30, (95% CI = 1.08-1.56) and 1.43 (95% CI = 1.22-1.68) respectively). Crucially, the detailed nature of these surveys had allowed for four measures of socioeconomic status to be considered (including both individual and area measures): the Carstairs index, occupation-based social class, educational attainment and economic activity. With this richer definition of social class than could be attained in other studies using fewer measures, they were able to demonstrate that higher rates of smoking in Greater Glasgow were attributable to the social pattern of smoking and lower socioeconomic status, with associations attenuated to statistical non-significance after adjustment (Table 1).14 This is an important finding, perhaps demonstrating that other measures of deprivation suffer a flooring effect i.e. that these deprivation measures have a lower limit to the data values they can specify. At a very similar time, they reported on a further analysis (also using the same three Scottish Health Surveys) that compared dietary habits in Glasgow with those in the rest of Scotland. They found that certain dietary habits (such as lower consumption of high-fibre bread and potatoes/pasta/rice in Glasgow) were also explained by socio-economic factors. Other factors (such as higher butter and salt consumption in women), however, were not, which hints once again at the complexity of such relationships and their measurement. They concluded that much of the tendency of people in Glasgow to have a poorer diet was explained by socioeconomic factors, although acknowledged the limitations of the study, particularly the use of self-completed questionnaires to assess diet and differential non-response rates by social class (Table 1).15

In 2009, Walsh et al compared mortality trends in Scotland with ten other post-industrial regions of Europe (using a variety of different data sources depending on the region). This was an important study given the proposal that deprivation, driven by the underlying effects of deindustrialisation, may have been the main drivers of poor health in Scotland. They found that mortality trends in West Central Scotland compared badly with other, similar, post-industrial regions of Europe (including some Eastern European regions with high levels of poverty). They therefore challenged the notion that post-industrial decline alone could be implicated (Table 1).10 Subsequent to this, in a study comparing mortality (using standardised mortality ratios (SMRs)) by levels of income deprivation at small area level in Glasgow (350 merged data zones with average 750 people per data zone) with the same data for Liverpool and Manchester (291 and 259 Lower Super Output Areas (LSOA) respectively with average 1500 people per LSOA) , they identified that Glasgow’s mortality rates, particularly premature mortality, were significantly higher than those of Liverpool and Manchester even after adjusting for income deprivation. They identified that a high proportion of the excess premature mortality in Glasgow was related to alcohol and drugs, and concluded that, if deprivation alone does not completely explain the increased mortality experience of Glasgow, other explanations such as the possibility of an ‘extreme’ behavioural risk profile in Glasgow should be explored (Table 1).9

Taken together, these studies created a mixed picture of the relationship between deprivation, health behaviours and health outcomes in Glasgow compared to other areas and began to reveal that the underlying explanation was unlikely to be straightforward. The later studies seemed to suggest that more complete measures of socio-economic status went further in being able to explain some of the observed variation, raising the possibility that simpler measures lacked the breadth or granularity to be able to discriminate true variation, particularly when also assessing the influence of complex variables such as diet. A review of the 2010 Walsh paper written by the second author (SG) picked up on this point, suggesting that existing measures of deprivation may not be comprehensive enough to pick up variation at extremes, such as might be present in Glasgow. It also suggested that other, potentially more ‘hidden’ (and certainly challenging to measure) factors, such as sectarianism, might play a hidden part in explaining some of the differences.16

To investigate this further, a study using 2008 and 2009 Scottish Health Survey data (n=13,996 adults of whom 3242 were resident in Greater Glasgow) was conducted for the Scottish Government to update and extend the studies that had used the 1995, 1998 and 2003 Scottish Health Surveys. Among its extensive findings, this showed that, when area and individual-level measures of socio-economic status were combined (area socio-economic status using the Scottish Index of Multiple Deprivation, individual socio-economic status using income-related benefits, National Statistics Socio-economic Classification (a measure of occupational status), economic activity (eg in paid employment), educational attainment, housing tenure and marital status) socio-economic status explained most of the differences in health behaviours and outcomes. However, there were a couple of important exceptions: heart attack (doctor diagnosed) and anxiety that were not explained by socio-economic differences. 17

Although this study seems to have clarified a great deal by using even more detailed definitions of socio-economic status, questions still remained. In the meantime, other groups had been studying a variety of other potential explanatory factors such as ethnicity and migration effects.

**Ethnicity and migration**

Fischbacher et al used mortality data from 362,029 deaths in Scotland to examine whether country of birth was linked to differences in mortality experience. They identified only small excess cardiovascular mortality among people born in India, Pakistan, Bangladesh and Ireland compared to people born in Scotland, but significantly greater mortality differences among South Asians when compared with residents of England and Wales (Table 1).18 Their findings highlighted the differences in mortality between Scotland and England and Wales for people of non-Scottish birth, but did not suggest that ethnicity alone would explain these differences..

Following this, in 2010, using mortality data from the England and Wales Office for National Statistics and the General Register Office for Scotland along with census data, Popham et al compared mortality differences between Scotland and England and Wales according to country of birth and country of residence. They found that, for people living in Scotland (n=3.3 million), risk of death was higher for those born in Scotland (n=3.0 million) than those born in England and Wales (n=332,255) . For people living in England and Wales (n=32 million), those born in Scotland (n=719,199) had a higher mortality rate than those born in England and Wales (n=31.2 milion). They concluded that country of birth may be a more important determinant of Scotland’s adverse outcomes than country of residence and suggest that early life factors may therefore be important in determining excess Scottish mortality (Table 1).19

In 2011, focusing more on Glasgow’s excess mortality, Popham et al conducted another interesting analysis of data from the Scottish Longitudinal Study (an anonymous representative sample of the Scottish population (n=137,073) linking 1991 and 2001 census records with health records and other events data), in which they explored whether the excess mortality rate in Glasgow compared with the rest of Scotland (not fully explained by socio-economic status) could be explained by selective migration. Selective migration is a net movement of people who are better off and healthier away from areas that are relatively more deprived. They compared migration from one area to another (from 1991 to 2001) between three areas: Glasgow, the next three largest Scottish cities (Edinburgh, Aberdeen and Dundee) and the rest of Scotland. Their outcome of interest was age and sex standardised mortality rates. They found that, although Glasgow lost a significant proportion of its population between 1991 and 2001 (which was more than the other three cities), this migration difference did not seem to account for differences in mortality rates between the areas (Table 1).20

**Current thinking**

Given the complexity of the problem, and the seeming lack of a single simple solution, in 2011 the Glasgow Centre for Population Health (GCPH) published an extensive literature review that tried to capture all of the different hypotheses that had been put forward to date to explain Scotland’s (and Glasgow’s) excess mortality. The review used Bradford Hill viewpoints on causality to try and evaluate how well each theory explained the mortality trends (see Box 1).21 They synthesised the theories into seventeen separate potential mechanisms (see Box 2) and identified outstanding research questions for each. 22 This review provided a comprehensive summary of the literature and a critical review of the strength of evidence for each hypothesis. Its findings and recommendations for future research form a very useful starting point for investigators interested in this area. The authors suggested that each of the hypotheses identified may form part of the explanation for Scotland’s mortality pattern but that some, such as inequalities and health behaviours, were likely to constitute a greater component of the causal pathway than others (see Figure 1 and Figure 2). They concluded that no single ‘cause’ was likely to fully explain the phenomenon but that the strongest individual candidates were negative health behaviours linked to cultural context.22 They commented that the divergence in mortality experience from European counterparts predominantly occurred from 1950 onwards. They also grouped possible explanatory factors into ‘downstream’ (such as greater exposure to negative health behaviours), ‘midstream’ (such as some of the cultural factors) and ‘upstream’ (such as poverty and unemployment) with greater or lesser degrees of likelihood (according to Bradford-Hill).22

In considering the GCPH review, it is worth remembering that, though widely used in this way, the use of Bradford-Hill ‘criteria’ as a checklist for evaluating whether a reported association might be causal, has been criticised in the past, particularly by Rothman, who felt that the Bradford - Hill ‘criteria’ do not clearly distinguish causal from non-causal relations (hence the term ‘viewpoints’ used here, rather than ‘criteria’).23 In the GCPH review, they did, however, provide a useful framework within which to consider the different hypotheses and may help to tease out where ongoing uncertainties remain and the authors recognised that their conclusions were hindered by a lack of empirical data for many of the hypotheses that were considered.

**Box 1. Bradford Hill viewpoints on causality**

1. Strength of association
2. Consistency
3. Specificity
4. Temporality
5. Biological gradient
6. Plausibility
7. Coherence
8. Experiment
9. Analogy

**Box 2: Hypotheses with the potential to explain / partially explain the Scottish effect identified in the GCPH review**22

1. Deprivation

2. Migration

3. Genetic differences

4. Health behaviours

5. Individual values

6. Different culture of substance misuse

7. Culture of boundlessness and alienation

8. Family, gender relations and parenting differences

9. Lower ‘social capital’

10. Sectarianism

11. Culture of limited social mobility

12. Health service supply and demand

13. Deprivation concentration ‘area effects’

14. Greater inequalities

15. Deindustrialisation

16. Political attack

17. Climatic differences

**Figure 1**: **The number of Bradford-Hill viewpoints met by each hypothesis for the divergence of the Scottish mortality pattern from the rest of Europe in the middle of the 20th Century** (reproduced with kind permission of the authors).22

**Figure 2.** **The number of Bradford-Hill viewpoints met by each hypothesis as explanatory factors in Scotland and Glasgow’s excess mortality** (reproduced with kind permission of the authors).22

Of the hypotheses shown in Figure 2, the most striking is perhaps ‘Political attack’. This has been described as the adverse neo-liberal influence on the organised working class implemented by the post-1979 UK Conservative governments.24 While difficult to immediately understand the connection with health outcomes, it has been argued that the effects of political forces on social inequalities, for example, are often underestimated and have a significant effect on health. 25 There is no doubt that political influence was closely linked to the profound degree of deindustrialisation occurring in the region in the late 1970s and early 1980s; the same time that Glasgow’s excess mortality was starting to emerge.26 Rapid loss of employment with associated increase in poverty and reduction in council housing was particularly intense in Glasgow and such change may well have had profound effects on health, both in the short term and reaching far into the future as successive generations are affected by the legacy of that period across their lifetime. 24

Since the GCPH review, other theories of interest have emerged in this field such as the importance of neighbourhood context. Research questions identified in the GCPH review (such as ‘Is there a difference in the geographical patterning of deprivation between Glasgow, Scotland and other populations – either now, or in the past, and is this associated with mortality patterns?) have led to some innovative research in subsequent years.22 Work assimilated by Chandola from a symposium in 2011 included examination of the effects of variation in the spatial distribution of urban populations and their link to health outcomes.27 Livingston presented a study at this symposium comparing Glasgow with Liverpool and Manchester, looked at contextual data at the neighbourhood level (as well as deprivation) to show differences in the way in which deprived neighbourhoods were arranged; either clustered around the city centre as in Liverpool and Manchester, or spread, as in Glasgow. He concluded that the ‘surrounding deprivation of a neighbourhood’ has an impact on mortality within a neighbourhood.27 However, in a subsequent paper with more detailed analyses of this potential effect, despite the more dispersed pattern of deprivation in Glasgow, Livingston and Lee conclude that patterning was not a major contributor to mortality in Glasgow as a similar effect is seen in Liverpool which has a different spatial distribution of deprivation.28 This does, however, still suggest a greater degree of complexity to the impact of deprivation than would be detected by, for example, measuring area deprivation alone. It is an example of how, in trying to fully explain Glasgow’s excess mortality, consideration may need to be given to much more difficult-to-measure issues, such as the way in which cities are constructed and change over time, and the relationship between this and health outcomes. A more recent study by Taulbut et al, for example, examined sub-regional spatial inequalities in 160 districts within selected similarly deindustrialised European regions (of which 22 were in West Central Scotland) and showed different patterns of life expectancy, with larger intraregional difference in life expectancy seen in West Central Scotland, suggesting that there may yet be more to explore in this area.29

Other more recent studies arising from the research questions identified in the GCPH review has included work on vitamin D deficiency, sectarianism, childhood and early years influences, and sense of cohesion (and there is ongoing work on the role of social capital). Among these are a systematic review and meta-analysis of studies investigating the link between low vitamin D and premature mortality, in which Rush et al identified that low vitamin D was associated with an increased all-cause mortality risk (hazard ratio 1.19, 95%CI 1.12-1.27) in adjusted models and recommended the need for further research in this area.30 Graham and colleagues compared Glasgow with Belfast – a similar post-industrial city with a stronger history of sectarianism. They concluded that, with this stronger history, if sectarianism were an important factor in explaining excess mortality in Glasgow, it is likely that similar findings would be seen in Belfast.31 Taulbut et al studied two British birth cohorts looking for evidence of a link between adverse childhood experience and mortality. They concluded that there was no evidence for an increased risk of adverse childhood events in Scotland compared with England (and specifically between Glasgow and the Clyde Valley and Merseyside and Greater Manchester), this reducing the likelihood of such adverse events being linked to excess mortality.32 Walsh et al applied a ‘sense of cohesion’ measure to a representative sample of adults in Glasgow, Liverpool and Manchester and identified that sense of cohesion was higher in Glasgow, making it an unlikely candidate to explain Glasgow’s excess mortality.33

**Glasgow’s excess mortality in the context of determinants of health**

Seminal work by Dahlgren and Whitehead in 1991 described a model of determinants of health that included ‘general socio-economic, cultural and environmental conditions’ as an outer layer influencing the material and social conditions in which people live (education, work environment, health services and so on), which in turn influences social and community networks and individual lifestyle factors.34 They recognised that ‘the age, sex and genetic make-up of each individual also plays a part, of course, but these are fixed factors over which we have little control’.34 As with the ‘downstream’, ‘midstream’ and ‘upstream’ factors described in the GCPH review, the hypotheses that have been explored in attempts to explain Glasgow’s excess mortality in Box 2 may be usefully considered in the context of the Dahlgren and Whitehead model. Moreover, the hypotheses summarised in the GCPH review could be thought of as applying at various levels. For example, ‘a different culture of substance abuse’ may be considered to have influence on ‘general socio-economic, cultural and environmental conditions’, perhaps affecting societal opinions and actions at the macro level (and thereby impacting a wide range of attitudes, policy and funding decisions for example), but also at the level of ‘individual lifestyle factors’ affecting personal behaviour (see Table 2). Considering both concepts in tandem may help to identify unexplored areas in the investigation of cause, as shown in the table. Although beyond the remit of this paper to describe in detail, an area that, to our knowledge, is underexplored as a potential explanatory factor in Scotland is epigenetics. Rapid developments in the field are identifying potentially important links between smoking, DNA methylation and cardiovascular disease, for example, and merit further investigation given the degree of unexplained excess mortality.35 Findings from the 239 people in the ‘Psycho- logical, social and biological determinants of ill health (pSoBid)’ cohort in Glasgow identified associations between DNA methylation and socioeconomic status and lifestyle factors.36 An important recent study is that by McCartney et al (2015) which examined mortality differences between Scotland and England by pooling 18 nationally representative observational studies (total n=222,829). They showed that only a quarter of the excess mortality in Scotland could be explained by factors such as socio-economic status and differences in health behaviours.37 This may tempt some to adopt a somewhat fatalistic view of Scotland and Glasgow’s excess mortality if it gives the impression that ‘little can be done about it’. We would argue, however, that it strengthens the argument, not only for further exploration of causality, but for adopting a proactive approach in Public Health terms, as it underlines that there is still much that can be done. Their finding, for example, that alcohol-related mortality is much higher in Scotland, even after adjustment for age, sex and socio-economic factors (HR 4.64 (95%CI 3.55-6.05) demonstrates the vital, ongoing role for Public Health in Scotland in addressing cultural and behavioural change around substance.37

Key to the further exploration of this idea, however, is the need to understand, as recognised by Dahlgren and Whitehead, that no ‘level’ of determinant exists in isolation, but that there is interplay between the levels. It may be that differences in the extent and nature of the interactions between these levels in Glasgow for many determinants of health, when compared to other cities, is as important as the impact of the individual determinants themselves.34

**The challenges of measurement**

Any review of this topic would be incomplete without some consideration of the difficulties encountered in measuring many of the factors under consideration. Measurement of almost any determinant in the Dahlgren and Whitehead model in order to derive its attributable risk, for example, presents significant challenges. If consideration is then given to the measurement of the interactions between determinants, the problem is even greater. To take deprivation as an example; Grundy and Holt described the complexities around accurate measurement of socio-economic status, particularly among older people.38 They raised issues such as the logistic difficulties of collecting information on household income, the relevance of past occupation (among retired people) to current socioeconomic status, and the lack of applicability of household measures to older people living in institutions.38 Measuring the impact of an ‘individual’ determinant such as socioeconomic status is therefore fraught with challenges. To fully understand socioeconomic status, a lifecourse approach may be more applicable (though even more complex to measure).39

In Glasgow, it is also of vital importance to consider historical events in terms of both societal-level events (such as deindustrialisation) as well as individual-level / personal history (such as drug use in childhood) and their impact not only on society, but on individuals across their lifecourse.10 This approach is in keeping with opinion expressed by Donnelly that it is important to consider adverse childhood experience and its health-damaging sequelae.40 Measuring such things and their effect in order to incorporate them into a causal pathway may not be achievable, and is certainly very difficult to do in the context of the kind of cross-sectional survey from which many of the associations with Glasgow’s excess mortality have been described.

However, their measurement may not be the most important consideration in Public Health terms. In contrast to the other studies discussed here, a recent study by Levin using data from the cross sectional 2006 Health Behaviour in School-Aged Children survey has shown that mental wellbeing among adolescents was better in Glasgow compared with the rest of Scotland.41 This suggests that the future lifecourse experience of current young Glaswegians has the potential to be very different from that of their forebears. It is also therefore true, that Public Health efforts to change the Glasgow effect will take time to become evident (and to measure), though there is evidence of progress seen in interventions such as those reducing the levels of dental caries.42 Observable changes in later outcomes, including mortality, are likely to take considerably longer.43 A particular focus is likely to be needed for younger people. There is good evidence from a study of the 31,648 suicides in England and Scotland between 2001 and 2006, for example, that the rate of suicide, particularly among young people, has been considerably higher in Scotland than in England (rate ratio 1.79 (95%Ci 1.62-1.98)).44 Similarly, although numbers may be small in absolute terms, the contribution of drug use to Scotland’s excess mortality among young people, is important because of its high mortality risk. In a cohort study of 1033 drug users across Scotland, Bloor et al estimated that, among people aged 15-54, drug use may account for 32% (95% CI 22.3-43.0%) of Scotland’s excess mortality.45 This reinforces the point made above about the on-going need for Public Health (and societal) action in these areas.

Similarly, the measurement of other individual determinants such as ‘ethnicity’ will include layers of complexity that are yet unexplored. Heterogeneity within ethnic groups, for example, may lead to differential impact on health-related behaviour and outcomes such as cardiovascular mortality.46

Further unravelling the web of complexity around Glasgow’s excess mortality may therefore include not only the need to consider the interplay of determinants in the Dahlgren and Whitehead model, but also address the methodological issues around their measurement, assess their relative influence, and make adequate allowance for a lifecourse perspective. The effort required in this challenging task is, however, of great importance. An updated synthesis report is due to be produced by the GCPH in 2015 and will no doubt address many of these complex issues.47 As Gavine and colleagues point out, continuing to understand the factors contributing to the ‘Glasgow effect’ is necessary to accurately inform Public Health endeavours to improve the health outcomes and reduce health inequalities in Scotland; it should not slip into common parlance as an explanation in its own right.48

**Conclusion**

A comprehensive explanation of Glasgow’s excess mortality remains elusive but is likely to lie in a complex and difficult-to-measure interplay of health determinants acting through the life course. It is an important example of the complexity that frequently lies behind health variations within and between populations. The research to date serves as an exemplar to usefully inform the critical evaluation of studies investigating morbidity and mortality variations in other countries and regions and the methodology of future study design in this area. Ongoing efforts to unravel its complexity are needed.

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The authors declare that they have no conflicts of interest

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**New Table 1:**

**Table 1. Summary of selected studies of health outcomes in Scotland and their main findings**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Study** | **Year** | **Geographical area of interest** | **Comparator area** | **Characteristic / exposure** | **Outcome** | **Main finding and effect size (where available)** |
| Carstairs4 | 1989 | Scotland | England and Wales | Socioeconomic status | Mortality | Scotland is more deprived than England and Wales and part of Scotland’s excess mortality may be explained by deprivation. Excess mortality is seen in all social classes, but greatest in people living in the poorest areas. |
| Hanlon8 | 2005 | Scotland | England and Wales | Socioeconomic status | Mortality | Scotland’s excess mortality is increasing over time and deprivation explains less of it over time.  Scotland to England and Wales ratio of mortality rates, directly standardised for age, sex and deprivation, increased from 104.7% in 1981 to 108.0% in 2001. |
| Fischbacher18 | 2007 | Scotland | England and Wales | Country of birth | Mortality | In Scotland, compared to people born in Scotland, those born in other parts of the UK had lower all-cause mortality (SMR 80.1 (95% CI 77-83) compared to SMR 100 in Scotland). In England and Wales, those born in Scotland had higher all-cause mortality (SMR 127.8 (95% CI 126-129) compared to SMR 100 in E&W). |
| Bloor45 | 2008 | Scotland | England | Drugs | Mortality | The contribution of deaths in drug users to national death rates was estimated as an attributable risk fraction for Scotland of 17.3% (95%CI 12.3% to 22.8%) and England of 11.1% (95%CI 7.8% to 14.8%). Estimated that 32.0% (95%CI 22.3% to 43.0%) of the excess mortality in people aged 15-54 in Scotland is due to drug use. |
| Gray11,15 | 2009 | Glasgow | Rest of Scotland | Socioeconomic status | Diet | Greater Glasgow area had unfavourable dietary profile, but not all explained by differences in socioeconomic status. E.g. lower consumption of high fibre bread in men (OR 0.74 (95%CI 0.62-0.88) age and survey year adjusted) in Glasgow compared with the rest of Scotland attenuated to (OR 0.88 (95%CI 0.74-1.05) with the addition of adjustment for socioeconomic status |
| Gray14 | 2009 | Glasgow | Rest of Scotland | Socioeconomic status | Smoking | Men living in Glasgow more likely to smoke than men living in the rest of Scotland (OR 1.3 (95%CI 1.08-1.57) age and survey year adjusted). This was attenuated to non-significance after additional adjustment for socioeconomic status (OR 0.92 (95%Ci 0.78-1.09)) |
| Walsh10 | 2009 | West Central Scotland | Ten comparable post-industrialised regions of Europe | De-industrialisation | Life expectancy  Mortality | Life expectancy improving at slower rate in West Central Scotland than most of the ten comparable European regions.  All cause mortality rate is higher in West Central Scotland and improving at a slower rate than comparable European areas.  Chronic liver disease and cirrhosis mortality rising much faster in West Central Scotland |
| Walsh9 | 2010 | Glasgow | Manchester and Liverpool | Income deprivation | Mortality | All-cause mortality higher in Glasgow than Manchester and Liverpool despite very similar deprivation profile (SMR 114.4 (95%Ci 113.2=115.5)). Excess mortality seen in all deciles of deprivation. |
| Popham19 | 2010 | Scotland | England and Wales | Country of birth | Mortality | Age and sex adjusted relative risk of death 1.2 (95% CI 1.12-1.29) for people born in Scotland and living in Scotland, and 1.15 (95% CI1.07-1.22) for people born in Scotland and living in England and Wales compared with people born in England and Wales and living in England and Wales. |
| Popham20 | 2011 | Glasgow | Three other Scottish cities (Edinburgh, Aberdeen, Dundee) and the rest of Scotland | Internal migration | Mortality | Relative age sex standardised mortality rate between Glasgow and the other three cities was 1.53 and between Glasgow and the rest of Scotland was 1.42 (no CIs given). Adjusting for area of migration did not affect these mortality differences. |
| McGuinness36 | 2012 | Glasgow (more deprived areas) | Glasgow (less deprived areas) | Epigenetics | DNA methylation | Global DNA methylation content was 17% (95%CI 4-29%) lower in most deprived vs. least deprived group. |
| Mok44 | 2012 | Scotland | England | Suicide | Mortality | Suicide rate ratio for Scotland vs. England between 2001 and 2006 was 1.79 (95%Ci 1.62-1.98). In men aged 15-44, the ratio was 2.06 (95%CI1.81-2.35) |
| McCartney37 | 2015 | Scotland | England | Socioeconomic and behavioural characteristics | Mortality | Hazard ratio for all-cause mortality Scotland vs. England was 1.40 (95%Ci 1.34-1.47), attenuated to 1.29 (95%CI1.23-1.36) after adjustment for socioeconomic and behavioural characteristics. Excess mortality was greatest among young adults and observed across all social classes (though greater in the unskilled). The excess was observed for a wide variety of causes of death including cancer, ischaemic heart disease and stroke. |

Footnote: CI = confidence interval

**Table 2. Potential links between Dahlgren and Whitehead determinants and GCPH review hypotheses with some suggested areas of future potential exploration.**

|  |  |  |
| --- | --- | --- |
| **Dahlgren and Whitehead determinants** | **GCPH review hypotheses** | **Potentially under-explored areas that may impact health outcomes** |
| General socio-economic, cultural and environmental conditions | * Culture of boundlessness and alienation * Culture of limited social mobility * Sectarianism * Greater inequalities * Deindustrialisation * Political attack * Climatic differences * Different culture of substance misuse | * Education provision, structure and opportunity * Culture of nihilism and cynicism * Conflict * Epigenetics |
| Living and working conditions | * Deprivation * Health service supply and demand * Deprivation concentration ‘area effects’ * Deindustrialisation | * Community safety * Access to opportunities for physical activity * Employment |
| Social and community networks | * Family, gender relations and parenting differences * Migration * Lower ‘social capital’ | * Perceived community safety * Perceived value to society |
| Individual lifestyle factors | * Health behaviours * Different culture of substance misuse * Individual values | * Health literacy * Psychological resilience and mental health * Spiritual wellbeing * Lifecourse effects * Epigenetics |
| Age, sex and constitutional factors | * Genetic differences | * Ethnicity * Lifecourse effects * Epigenetics |