**Lifespan inequality and mortality crises in Scandinavia**

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**Introduction**

Life expectancy is often used as a summary measure to describe the state of a population in terms of mortality as well as in terms of health (van Raalte, Sasson, and Martikainen 2018). In this sense, life expectancy is indeed very convenient, as it expresses the average remaining years of life for a certain cohort, be it real or synthetic (Preston, Heuveline, and Guillot 2001). However, precisely because of this synthesis, life expectancy can camouflage other important characteristics of a population. One of such characteristics is the variation in the age at death, also known as lifespan variation. Lifespan variation, which at the individual level describes the uncertainty of the timing of death, has been decreasing as life expectancy and the modal age at death have increased (Colchero et al. 2016; Kannisto 2001; Smits and Monden 2009; Vaupel, Zhang, and Raalte 2011). Yet, life expectancy and lifespan variation have been shown to follow different historical and contemporary trends, so that an increase in life expectancy does not necessarily imply a decline in lifespan variation (Aburto and Raalte 2018; Brønnum-Hansen 2017; Sasson 2016; Wilmoth and Horiuchi 1999). Therefore, by only focusing on life expectancy, we miss a fundamental inequality in age at death.

Studies on lifespan variation have mostly focused on populations with continued improvements in mortality or analysed differences by social determinants, such as SES or educational level (Edwards and Tuljapurkar 2005; Lariscy et al. 2016; Permanyer and Scholl 2019; van Raalte 2011). Yet, to the best of our knowledge, lifespan variation has never been studied in circumstances where mortality increases sharply. We fill this gap by focusing precisely on populations that have experienced a mortality crises.. Although such mortality crises happened in the past and are not a current event in contemporary Europe, . Recent evidence suggests that they will become an ever more pressing question as extreme weather events increase in frequency with climate change, potentially bringing natural catastrophes and food shortages in their wake (Cynthia et al. 2001). By analysing and comparing the evolution of different populations, we aim to understand whether a regular pattern emerges which could precede or be a consequence of mortality crises. In this way, studying the patterns in lifespan variation of past populations could help us better understand the impact of mortality crises today.

Mortality patterns during times of crises have already been the object of research. Although these studies do not tackle the issue of lifespan variation, they do shed light on what we can expect from a broader point of view. First of all, no single pattern can be expected, as age and gender specific mortality rates vary, depending on the nature of the crisis itself. For example, male adults are more at risk during wars or some epidemics such as those caused by HIV/AIDS (Gaylin and Kates 1997; Hosegood, Vanneste, and Timæus 2004), while natural disasters seem to affect more women, children and the elderly (Bern et al. 1993; Frankenberg et al. 2011; Neumayer and Plümper 2007). In this paper, however, we focus on two specific types of mortality crises: famines and epidemics.

Bongaarts and Cain (1982, in Kane 1987) hypothesised that mortality would increase during a famine to reach a peak at its end. Afterwards, mortality rates would gradually decrease as the long-term consequences of food deprivation took their toll. Age and gender specific mortality patterns vary depending on the cultural and social environment. Some information about Scandinavian trends can be found in Bengtsson, Campbell and Lee (2009), who analysed historical data linked with increased food prices. They found that infants were generally less affected by increased food prices, as they mostly depend on breast-feeding, while older children are much more sensitive to external conditions. Because of breastfeeding and pregnancy, women are more vulnerable to food deprivation, which may also affect them more in case of an unequal distribution of food in the household, which often favours males (although Zarulli *et al.* (2018) find that the life expectancy gender gap advantages females at almost all ages even during crises). Finally, the elderly are also affected, but show little differences in terms of gender and socio-economic status, possibly as a result of the selection of the most robust individuals into old age.

When looking at epidemics, trends become even more complicated, as each disease is characterised by a specific age pattern, dictated by biological and social determinants. Of the three epidemics we consider, two were caused by measles, which traditionally affects children, but also non-immunised adults, common in isolated communities previously spared by the virus. The last crises we analyse was a typhus and dysentery epidemic, diseases which kill especially weakened individuals, such as children and the elderly (Castenbrandt 2014).

**Data and Methods**

We use data from the Human Mortality Database (HMD). The HMD supplies data covering multiple population with nearly complete information, assuring a high level of quality. For this reason as well, only few countries, mostly European, are included in the database and fewer yet contain data series from the XVIII or XIX centuries, where mortality crises were more common in Europe. For this reason, we have chosen to focus on Scandinavian countries, for which data were collected by parishes at a reasonably precise level during this time. More precisely, we use these data to study four mortality crises, two in Sweden and two in Iceland.

Dribe, Olsson and Svensson (2015) describe the mortality response to the 1772-1773 famine and the 1808-1809 epidemic in Sweden. Crop failures in large regions of Sweden caused by unusual weather in 1772 exacerbated already high food prices and led to a famine which peaked the following year. In 1773 mortality rates were 86% higher in the most affected counties, compared to the others and that CDR doubled in central Sweden. Although all age-groups were affected, children between 1 and 14 years of age were suffered the most, while infants witnessed a relatively small increase in mortality. Mortality was mostly driven by nutrition-related diseases, specifically typhus and dysentery (which alone accounted for 50% of the excess mortality that year). Typhus and dysentery are also the diseases involved in the 1808-1809 epidemics, which followed troop movements involved in the Finnish War. However, the increase in mortality is thought to have resulted from epidemics than from war itself (Glei et al. 2019). As a consequence, mortality follows the same age-pattern as in 1773, although the difference between children over 1 year and the other age groups is even greater.

In Iceland, we consider two measles epidemics, in 1846 and 1882. In both years, particularly cold spring and summer forced fishermen to concentrate in shore villages, facilitating the spread of the disease, brought by Danish sailors. In 1846, even the oldest Icelanders had never been in contact with measles, which spread rapidly through the unimmunised population. Although mortality increased for all ages, children and the elderly were affected more severely, because of their physiological weakness. The epidemic lasted from July to December and caused the death of around 3% of the whole population. The individuals that survived were better prepared to face the following epidemic in 1882, which mostly affected ages under 50. The immunisation of the population also meant that this epidemic lasted only from June to August and led to the death of around 2% of the population (Cliff, Haggett, and Graham 1983; Shanks et al. 2015).

*Lifespan variation indicators*

Lifespan variation can be measured using a wide range of techniques and all of them have been found to be highly correlated in empirical datasets when measured from young ages (Wilmoth and Horiuchi 1999). They are not, however, completely interchangeable. As van Raalte and Caswell (2011) point out, they differ in their formal properties and in the underlying concept they gauge. Following the authors’ analysis, we have decided to include three measures of lifespan variation in this work: the standard deviation *S*, life disparity *e†* (add reference to Vaupel & Canudas-Romo 2002) and the Gini coefficient *G (Hanada or Scholkinov et al 2003)*.These represent a mix between absolute and relative measures of variation. It is useful to consider relative variation for comparisons, especially when considering dramatic changes in life expectancy, which could mask significant differences. At the same time, absolute variation is a more directly understandable measure and can better inform us on the concrete changes experienced by our populations. Moreover, using various methods with different sensitivities to changes in age-specific mortality rates will allow us to come to more robust conclusions. We also prefer these to other measures, because they are easily interpretable.

*Decomposition methods*

Although studying the variation in the mortality distribution will already inform us on its general trend, we also plan to analyse these changes more in detail, as the information included in the HMD makes it possible to identify gender and age at death. We will first compare lifespan variation differences between males and females, to study the behaviour of the gender gap under crisis conditions. Then, we will decompose the change in variation by age, to understand whether certain age groups contributed more to the increase or decrease in lifespan variation. We will do so by using the life table response experiments (LTRE) method, which expresses the observed change in the value of a function (in our case one of the measures of lifespan variation to its parameters and as a combination of the sensitivity of this function to its parameters and of the changes in the parameters themselves (in our case, age-specific mortality rates) (Caswell 2019). Although other methods of decomposition by age have been developed (see for example Appendix B of Wilmoth and Horiuchi 1999), LTRE has been shown to be applicable to the measures we consider through relatively straightforward calculations (van Raalte 2011).

We hope that studying the patterns in lifespan variation across age and gender of past populations will help us better understand the impact of mortality crises today. These results could be particularly helpful for organising the response to mortality crises in the future. Indeed, if some specific age groups are revealed to be especially vulnerable in such cases, the organisations responsible for the response will be able to better prepare and organise it.

**Research questions**

In this work, we will answer three main research questions.

First of all, we want to study whether lifespan variation changes during and after a mortality crisis. In order to provide an answer, we will compare the mean variation of the years before the crisis to the one measured during the crisis year(s) and in the following years. One could expect that a severe enough crisis would cross social lines, affecting the whole population equally, as happened for the European Black Plague (Livi Bacci 2012), so that lifespan variation would decrease in such situations. However, more recent episodes have shown clear inequalities in the mortality during extreme events, famously hurricane Katrina (Zoraster 2010). In fact, most of our data is connected to famine episodes, from which the wealthy are protected to a certain degree. Even if a crisis were to cross social lines, it would likely affect individuals differently depending on their age, disproportionately increasing the mortality of the extreme and more vulnerable ages and thus variation. Finally, a decrease in life expectancy, which is inevitable during a mortality crisis, gives mechanically more space for variation in age at death, as the modal age at death shifts to the left. All of these considerations lead us to believe that variation will increase rather than decrease during a mortality crisis. After the end of the episode, we expect that variation will continue to be higher than pre-crisis level, but that it will gradually decrease as the parts of the population most affected by the crisis recover. This trend could be balanced by a reduction in lifespan variation due to a selection, during the crisis, of the more robust individuals, who would die later on average.

Our second research question looks at gender differences. Zarulli *et al.* (2018) have found that the gender gap in life expectancy remains during high-mortality regimes. In the same way, we expect that mortality crises will affect both subpopulations similarly, so that the gender gap in lifespan variation, which generally favours females (van Raalte 2011), will not change in high mortality situations.

Our final question asks whether some ages especially contributed to the change in variation witnessed during and after the crisis, and if so which ones. We predict that a mortality crisis will particularly affect children and the elderly, as these sub-groups are physically less equipped to deal with extreme conditions and because their survival might become less of a priority in situations where resources are scarce. Moreover, deaths at the extremes of a distribution will more heavily affect variation. Therefore, we expect that these age groups will largely contribute to the expected increase in lifespan variation.

**Preliminary Results**

A couple of graphs would be good as preliminary results for the abstract. You could plot the age-at death distribution (function dx of the lifetable), trends in lifespan variation for males and females. We should open a github repository to have all the code to reproduce the results.

**Discussion**

**References**

Aburto, José Manuel and Alyson van Raalte. 2018. ‘Lifespan Dispersion in Times of Life Expectancy Fluctuation: The Case of Central and Eastern Europe’. *Demography* 55(6):2071–96.

Bengtsson, Tommy, Cameron Campbell, and James Z. Lee, eds. 2009. *Life under Pressure: Mortality and Living Standards in Europe and Asia, 1700 - 1900*. 1. MIT press paperback ed. Cambridge, Mass.: MIT Press.

Bern, C., J. Sniezek, G. M. Mathbor, M. S. Siddiqi, C. Ronsmans, A. M. Chowdhury, A. E. Choudhury, K. Islam, M. Bennish, and E. Noji. 1993. ‘Risk Factors for Mortality in the Bangladesh Cyclone of 1991.’ *Bulletin of the World Health Organization* 71(1):73–78.

Castenbrandt, Helene. 2014. ‘A Forgotten Plague’. *Scandinavian Journal of History* 39(5):612–39.

Caswell, Hal. 2019. *Sensitivity Analysis: Matrix Methods in Demography and Ecology*. Cham: Springer International Publishing.

Cliff, Andrew D., Peter Haggett, and Rosemary Graham. 1983. ‘Reconstruction of Diffusion Processes at Local Scales: The 1846, 1882 and 1904 Measles Epidemics in Northwest Iceland’. *Journal of Historical Geography* 9(4):347–68.

Colchero, Fernando, Roland Rau, Owen R. Jones, Julia A. Barthold, Dalia A. Conde, Adam Lenart, Laszlo Nemeth, Alexander Scheuerlein, Jonas Schoeley, Catalina Torres, Virginia Zarulli, Jeanne Altmann, Diane K. Brockman, Anne M. Bronikowski, Linda M. Fedigan, Anne E. Pusey, Tara S. Stoinski, Karen B. Strier, Annette Baudisch, Susan C. Alberts, and James W. Vaupel. 2016. ‘The Emergence of Longevous Populations’. *Proceedings of the National Academy of Sciences* 201612191.

Cynthia, Rosenzweig, Anna Iglesias, Xiao-Bing Yang, Paul R. Epstein, and Eric Chivian. 2001. ‘Climate Change and Extreme Weather Events; Implications for Food Production, Plant Diseases, and Pests’. *Global Change & Human Health* 2(2).

Dribe, Martin, Mats Olsson, and Patrick Svensson. 2015. ‘Famines in the Nordic Countries, AD 536–1875’. *Lund Papers in Economic History* 138.

Edwards, Ryan D. and Shripad Tuljapurkar. 2005. ‘Inequality in Life Spans and a New Perspective on Mortality Convergence Across Industrialized Countries’. *Population and Development Review* 31(4):645–74.

Frankenberg, Elizabeth, Thomas Gillespie, Samuel Preston, Bondan Sikoki, and Duncan Thomas. 2011. ‘Mortality, The Family and the Indian Ocean Tsunami’. *The Economic Journal* 121(554):F162–82.

Gaylin, Daniel S. and Jennifer Kates. 1997. ‘Refocusing the Lens: Epidemiologic Transition Theory, Mortality Differentials, and the AIDS Pandemic’. *Social Science & Medicine* 44(5):609–21.

Glei, Dana, Hans Lundström, John Wilmoth, Gabriel Borges, Mia Zhong, and Magali Barbieri. 2019. ‘Sweden - Background and Documentation’.

Hosegood, Victoria, Anna-Maria Vanneste, and Ian M. Timæus. 2004. ‘Levels and Causes of Adult Mortality in Rural South Africa: The Impact of AIDS’. *AIDS* 18(4):663.

Kane, Penny. 1987. ‘The Demography of Famine’. *Genus* 43(1/2):43–58.

Kannisto, Vaino. 2001. ‘Mode and Dispersion of the Length of Life’. *Population: An English Selection* 13(1):159–71.

Lariscy, Joseph T., Claudia Nau, Glenn Firebaugh, and Robert A. Hummer. 2016. ‘Hispanic-White Differences in Lifespan Variability in the United States’. *Demography* 53(1):215–39.

Livi Bacci, Massimo. 2012. *A concise history of world population*. Chichester, West Sussex, UK: Wiley-Blackwell.

Neumayer, Eric and Thomas Plümper. 2007. ‘The Gendered Nature of Natural Disasters: The Impact of Catastrophic Events on the Gender Gap in Life Expectancy, 1981–2002’. *Annals of the Association of American Geographers* 97(3):551–66.

Permanyer, Iñaki and Nathalie Scholl. 2019. ‘Global Trends in Lifespan Inequality: 1950-2015’. *PLOS ONE* 14(5):1–19.

Preston, Samuel, Patrick Heuveline, and Michel Guillot. 2001. *Demography: Measuring and Modeling Population Processes*. Oxford: Blackwell Publishing.

Raalte, Alyson A. van, Isaac Sasson, and Pekka Martikainen. 2018. ‘The Case for Monitoring Life-Span Inequality’. *Science* 362(6418):1002–4.

van Raalte, Alyson. 2011. ‘Lifespan Variation: Methods, Trends and the Role of Socioeconomic Inequality’. Erasmus University, Rotterdam.

Sasson, Isaac. 2016. ‘Trends in Life Expectancy and Lifespan Variation by Educational Attainment: United States, 1990–2010’. *Demography* 53(2):269–93.

Shanks, G. D., M. Waller, H. Briem, and M. Gottfredsson. 2015. ‘Age-Specific Measles Mortality during the Late 19th–Early 20th Centuries’. *Epidemiology & Infection* 143(16):3434–41.

Smits, Jeroen and Christiaan Monden. 2009. ‘Length of Life Inequality around the Globe’. *Social Science & Medicine* 68(6):1114–23.

Vaupel, James W., Zhen Zhang, and Alyson A. van Raalte. 2011. ‘Life Expectancy and Disparity: An International Comparison of Life Table Data’. *BMJ Open* 1(1):e000128.

Wilmoth, John R. and Shiro Horiuchi. 1999. ‘Rectangularization Revisited: Variability of Age at Death within Human Populations\*’. *Demography* 36(4):475–95.

Zarulli, Virginia, Julia A. Barthold Jones, Anna Oksuzyan, Rune Lindahl-Jacobsen, Kaare Christensen, and James W. Vaupel. 2018. ‘Women Live Longer than Men Even during Severe Famines and Epidemics’. *Proceedings of the National Academy of Sciences of the United States of America* 115(4):E832–40.

Zoraster, Richard M. 2010. ‘Vulnerable Populations: Hurricane Katrina as a Case Study’. *Prehospital and Disaster Medicine* 25(1):74–78.