



# DEMOGRAPHIC RESEARCH

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*Research Material*

## **APC curvature plots: Displaying nonlinear age-period-cohort patterns on Lexis plots**

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## **APC curvature plots: Displaying nonlinear age-period-cohort patterns on Lexis plots**

**Enrique Acosta<sup>1</sup>**

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### **Abstract**

#### **BACKGROUND**

The analysis of age-period-cohort (APC) patterns of vital rate changes over time is of great importance for understanding demographic phenomena. Given the limitations of statistical modeling, the use of graphical analyses is often regarded as a more transparent approach to identifying APC effects.

#### **OBJECTIVE**

The current paper proposes a Lexis plot for the depiction and analysis of curvature, which is defined as the estimable nonlinear component of age, period, and cohort effects.

#### **METHODS**

In a single visualization, we combine the dynamics of the location, the magnitude, and the spread of nonlinear temporal effects for multiple populations or demographic phenomena. Using vital rates, we provide three examples in which we analyze the APC nonlinear effects of different demographic phenomena.

#### **RESULTS**

We construct several APC curvature plots to display the following patterns: the modal cohort of excess mortality from drug-related causes by racial/ethnic group in the United States among the baby boomer generations; the modal age of excess mortality in young adults; and the modal age of fertility over cohorts and across populations.

#### **CONTRIBUTION**

The use of the APC curvature plot offers more flexibility when analyzing nonlinear APC effects than the use of mathematical models or other Lexis plots.

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## 1. Introduction

It has long been recognized that populations change along the three dimensions of age, period (typically calendar year), and cohort (typically year of birth) (Caselli and Vallin 2005; Keiding 2011). The question of whether it is possible to independently isolate these age, period, and cohort (APC) effects on temporal changes in population phenomena has been vigorously debated since the first half of the 20<sup>th</sup> century (Keyes et al. 2010; Murphy 2010). Recently, discussions of this topic became even more heated in reaction to the set of methods proposed by Yang and colleagues (Bell and Jones 2013; Fosse and Winship 2018, 2019a; Luo 2013; Masters et al. 2016; Reither et al. 2015; Yang and Land 2013). At the center of this latest debate is the identification problem that arises because of the perfect linear dependence between these three dimensions (age = period – cohort), as this dependence makes it impossible to estimate a unique solution without imposing additional constraints.

Given this limitation, the use of graphical analyses is often regarded as a more transparent approach to identify APC effects than the use of statistical modeling (Murphy 2010; Preston and Wang 2006; Willets 2004). Consistent with this idea, the hand-drawn contours indicating cohort mortality improvement patterns presented in the work of Kermack, McKendrick, and McKinlay (1934) have been recognized as a pioneering example of research demonstrating that long-term mortality change tends to follow the birth cohort dimension (Finch and Crimmins 2004; Hobcraft, Menken, and Preston 1982; Preston and Wang 2006). Nevertheless, the question of whether the identification of such visual patterns could also be interpreted as “cohort effects” (i.e., whether improvements in mortality could be the consequence of period-based improvements or the improved performance of newer birth cohorts) has yet to be resolved. This is, for example, the case for APC statistical models; see Murphy (2010) for an interesting review.

The analyses that focus instead on identifying divergence from secular trends (also known as curvature<sup>3</sup>) in each of the three temporal dimensions (Holford 1983; Rodgers 1982; Tango and Kurashina 1987) are more effective and less polemical than those focused on identifying the dominant patterns of change. For example, such analyses might seek to identify a systematic fluctuation that follows a cohort, and that is independent of changes over the age and the period dimensions. Such curvature would

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<sup>3</sup> Note that the term ‘curvature’ used here refers to deviations from linear effects, as originally proposed by Holford and used in the APC literature (Clayton and Schiffers 1987; Holford 1983, 1991, 2005; Tango and Kurashina 1987). Other names used for this component are ‘nonlinear effects’ and ‘nonlinear fluctuations.’ This term should not be confused with ‘local curvature,’ also referred to as ‘contrast’ and ‘second-order difference’ (Acosta et al. 2019b; Clayton and Schiffers 1987; Holford 1991; Pullum 1980; Tango and Kurashina 1987; Tarone and Chu 1996), which are used to measure the change in the slope of the effects.

be indicative of divergence in the behavior of members of a particular set of cohorts from the behavior of the cohorts born before or after them.

Our aim here is to propose a visualization tool that allows for the depiction of such APC curvature and its attributes. In particular, we overcome a key limitation of the existing methods: comparing the change in curvature attributes over time across several demographic phenomena or populations in a single visualization.

The paper is structured as follows. In section 2, we present a short description of some of the existing statistical and graphical methods for the detection and analysis of nonlinear APC effects, while highlighting the advantages and the limitations of each of these methods. For the sake of clarity, we complement the description of each method by applying it to the analysis of excess mortality related to drug abuse among Hispanic baby boomers in the United States. These mortality rates are presented in a Lexis surface in Figure 1. In section 3, we present our proposed visualization technique. We describe its advantages and explain how it can complement existing methodologies. Section 4 consists of a step-by-step description of the construction of the plot. Section 5 provides three empirical applications of the proposed visualization (shown in Figures 5–7). In the final section, we draw some conclusions.

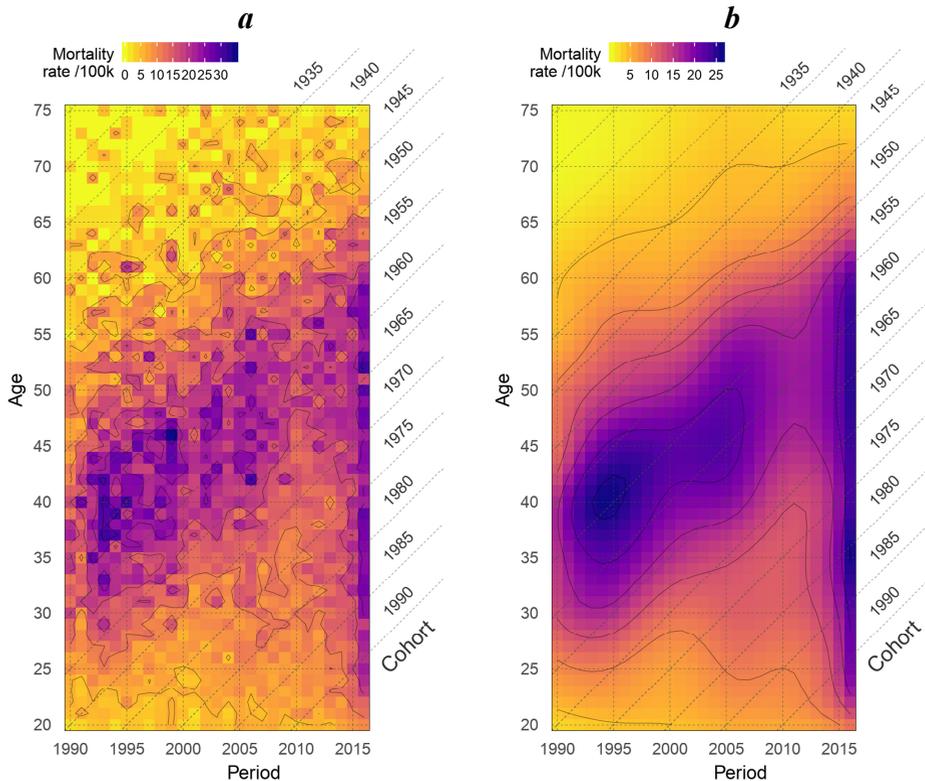
For the examples presented, we used the R programming language (R Core Team 2018) for the analysis and data visualization. In addition, we used the packages *ggplot2* (Wickham 2016), *Epi* (Carstensen et al. 2018), *MortalitySmooth* (Camarda 2012), and *HMDHFDplus* (Riffe 2015). All of the data and code for replicating results are openly available (Acosta 2019).

## 2. Existing methods for analyzing APC curvature

As discussed in the introduction, partitioning temporal variations into linear APC components is controversial due to the identification problem. Systematic divergence from this linearity, also known as curvature, is, however, unambiguously identifiable. Such curvature could have concave (humps) or convex (valleys) shapes, independent of linear trends. When looking at curvature graphically, it becomes clear that if a deviation follows horizontal, vertical, or diagonal trends on a Lexis surface, it is indicative of an age, period, or cohort effect, respectively. For instance, when looking at the Lexis surface of drug-related mortality rates for US Hispanic males depicted in Figure 1, we can see that the diagonal pattern suggests that there is a cohort pattern of increased risk of death among the members of the baby boomer cohorts (i.e., conventionally defined as the cohorts born between 1946 and 1964). In this section, we review some existing statistical (detrended Age–Period–Cohort models) and visual (Lexis surfaces of changes in vital rates, and Lexis surfaces of excess rates) tools for the detection and analysis of a

curvature, and apply each one to the case of drug-related mortality among Hispanic baby boomers.

**Figure 1:** Lexis surface of observed and smoothed drug-related mortality rates for Hispanic males in the United States during 1990–2016, ages 20–75



*Note:* Panel (a) is a Lexis surface of observed drug-related mortality rates for Hispanic males in the United States between 1990 and 2016. Panel (b) is the corresponding Lexis surface of smoothed mortality rates. The smoothing was done using a two-dimensional nonparametric smoothing technique. Specifically, we assume that our events are Poisson-distributed, and smooth the data with P-splines, with the smoothing parameters optimized according to AIC. Smoothing was performed using the *MortalitySmooth* R package developed by Camarda (2012).

## **2.1 Statistical APC models for analyzing curvature**

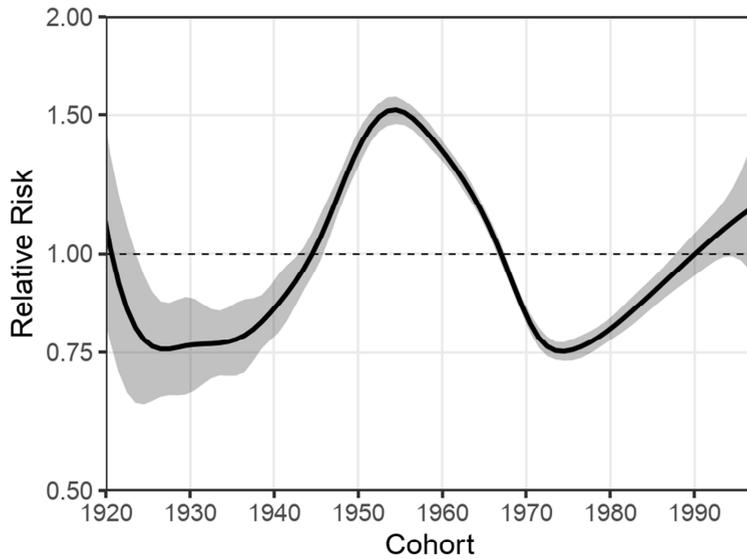
Several arithmetic and statistical models have been proposed to identify deviation from trends, such as the Median Polish technique (Keyes and Li 2010; Selvin 2001; Tukey 1977) and the Detrended Age-Period-Cohort models (dAPC) (Carstensen 2007; Clayton and Schifflers 1987; Holford 1983).

In general terms, the APC effects can be decomposed into linear trend and curvature components (Fosse and Winship 2019b; Holford 1983; Tango and Kurashina 1987). While there are infinitely many ways to partition the linear effects into APC components, the curvature components have the same shape and magnitude regardless of the parameterization used to fit the model (Clayton and Schifflers 1987; Holford 1983, 1991). To obtain these nonlinear components, the effects (log of relative risks) in dAPC models are constrained to be zero on average, with a zero slope (detrended). In this parameterization, the reference category is the overall age, period, or cohort average; centered at its linear trend component. More details about the parameterization of the model can be found in Carstensen (2007).

As an example, Figure 2 shows the estimates of the nonlinear cohort effects in relative risk values that are obtained from a dAPC model applied to the drug-related mortality of Hispanic males. If we focus our attention on the concave shape of the curvature for the boomer cohorts (born between 1940 and 1970), we see that according to these estimates, the most disadvantaged cohort (the curvature peak) is the cohort born in 1954, for whom the risk of dying from a drug overdose is nearly 1.6 times higher than would be expected given the overall linear cohort trend.

Although identifiable, the curvature estimates are average effects that are fixed over the whole length of time (Chauvel 2013). The invariability of these estimates has two particularly undesirable consequences. First, the estimates do not allow the size of the effect to vary over time/age. Second, because of this lack of variability, the estimates attribute the highest or the lowest nonlinear effect to a fixed temporal dimension, permanently labeling it as advantaged or disadvantaged. For instance, from the estimates depicted in Figure 2, is not possible to establish variation in the magnitude of the relative risks over age/time; and, as a consequence, we are unable to determine whether those individuals born in 1954 actually had the highest risk during the entire period under observation.

**Figure 2: Relative risks of drug-related mortality for cohorts of Hispanic males in the United States**



Notes: The cohort relative risk estimates were obtained from a detrended APC model (dAPC) applied to drug-related mortality for Hispanic males aged 20–75 during the 1990–2016 period. Cohort effects (i.e., the log of the relative risks) are constrained to be zero (0) on average, with zero (0) slope. B-splines are used for fitting the APC effects. The reference category is the overall cohort average, indicated in the plot with a horizontal dashed line. The gray area indicates the 95% confidence interval. Estimates were obtained using the R package *Epi* (Carstensen et al. 2018).

An alternative model proposed by Chauvel (2013) allows for the estimation of a hysteresis value that indicates whether the magnitude of the curvature increases or decreases over time. This model is, however, a fixed measure of change that reflects only constant increases or decreases over time, and it does not allow for modifications in the curvature ridge/floor location over time/age.

## 2.2 Graphical tools for analyzing curvature

Several graphical tools have been proposed for uncovering patterns of systematic divergence from linear trends. The focus of most of these tools is the analysis of mortality. Plots of the change in smoothed rates over age/cohort, over period/cohort, and over age/period are effective tools for discerning the dynamics of demographic phenomena over time, and for uncovering patterns of systematic divergence from APC

linear trends. Indicative Lexis diagrams (Willets 2004) and colored Lexis surfaces (Rau et al. 2008, 2018; Richards, Kirkby, and Currie 2006) of these derivatives have been proposed as visualization techniques that could be applied to identify the presence of such patterns in Lexis plots.

Variations over APC dimensions are complementary perspectives that can be used to visually detect curvature on Lexis surfaces. When looking at rate changes over age/cohort (i.e., vertical changes along the same period in the Lexis diagram), we can identify within-period divergence in rate changes, although we cannot unambiguously attribute such divergence to age or cohort because of the identification problem. Analogously, when looking at changes over age/period (i.e., diagonal changes along the same cohort in the Lexis diagram) it is possible to determine age/period fluctuations that are independent from variations over cohorts.

Raw derivatives in vital rates are generally noisy, particularly when using high-resolution data or low-frequency events. Thus, it has been suggested that the underlying data should be smoothed over ages and years in order to reduce random fluctuations<sup>4</sup> (Rau et al. 2018); as shown in Figure 1.

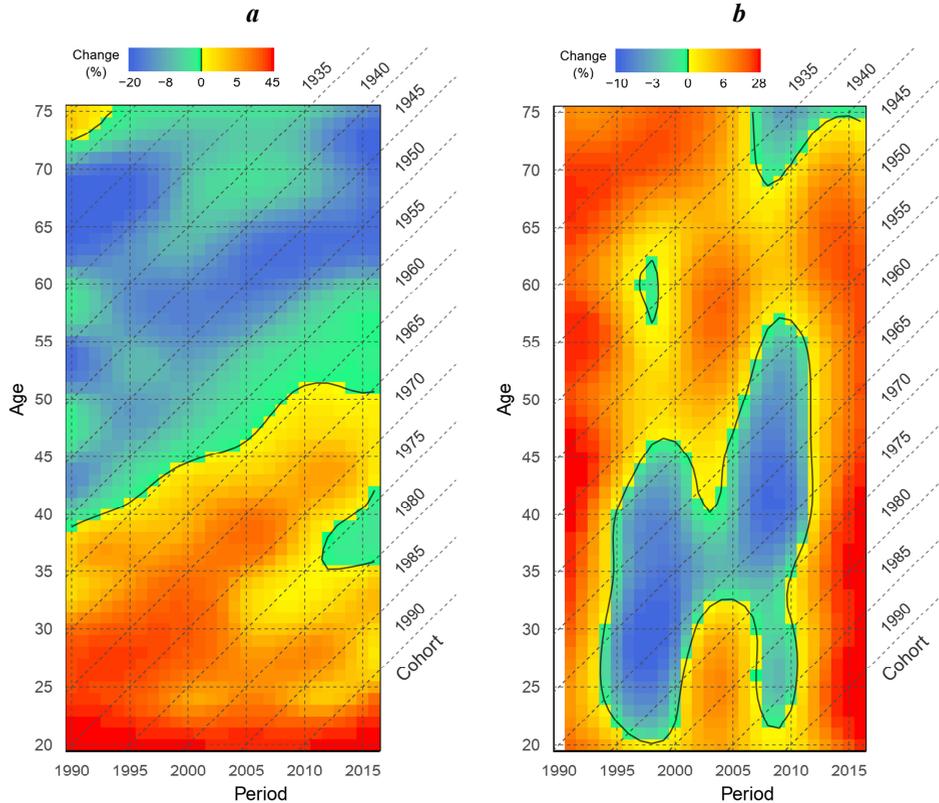
Figure 3 depicts Lexis surfaces of rates of mortality change over age/cohort (Panel a) and over period/cohort (Panel b) from drug-related causes in Hispanic males in the United States. Note that for this empirical example, we do not depict the Lexis surface of rates of mortality change over age/period because our interest is on cohort effects, which are not identifiable in such surfaces.

Given the large variations in drug-related mortality over time, the plot of derivatives over age/cohort (vertical changes along the same period as in the Lexis diagram, Figure 3a) depicts the nonlinear cohort effect of the Hispanic boomers more clearly than the plot of derivatives over period/cohort (horizontal changes along the same age group as in the Lexis diagram, Figure 3b). The diagonal contour line following the cohorts born around 1955 in Figure 3a indicates that the peak in mortality rates followed those cohorts, even if the cohort curvature is camouflaged by the strong period fluctuations visible in Figure 3b.

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<sup>4</sup> Note that smoothed rates are useful for identifying divergence from the trend that occur over large time scales (i.e., effects that appear gradually over several age, period, or cohort groups). However, when divergence is very exceptional and compromises only a few units of time in a given dimension (i.e., effects that appear in only a couple of age, period, or cohort groups) the smoothing process could weaken or remove the divergence of interest. In those cases, the use of observed rates is preferable to the use of smoothed rates.

**Figure 3: Lexis surfaces of changes in drug-related mortality rates over age/cohort and over period/cohort for Hispanic males in the United States**



Notes: Panel (a) is the Lexis surface of changes over age/cohort (read vertically from young to old ages, or from newer to earlier cohorts); with a yellow-to-red scale indicating the relative mortality rate increase for age  $x$  compared to age  $x-1$  (or cohort  $k$  compared to cohort  $k+1$ ) in the same year, and a green-to-blue scale indicating a relative mortality decline between consecutive ages/cohorts. The black contour line depicts zero changes in mortality, which indicates a local maximum or minimum death rate in a given calendar year (i.e., a curvature ridge or floor). For example, if we examine the year 2000, we see that death rates increased with age until hitting their maximum value at age 44 (curvature ridge). From ages 45 to 75, the death rates declined over age. Panel (b) is the corresponding Lexis surface of changes in drug-related mortality rates over period/cohort (read horizontally from earlier to more recent calendar years/cohorts) for Hispanic males in the United States; with a yellow-to-red scale indicating the relative mortality rate increase for year  $t$  compared to year  $t-1$  (or cohort  $k$  compared to cohort  $k-1$ ) in the same age, and a green-to-blue scale indicating a relative mortality decline between consecutive calendar years. Again, the black contour indicates a curvature ridge or floor in a given age. For example, if we examine age 50, we see that the death rates increased with time until hitting a maximum value in 2005 (curvature ridge). From 2006, the death rates declined over time until reaching a local minimum value in 2011 (curvature floor).

Alternatively, the use of Lexis surfaces to depict deviations from trends is an effective way to display variation in the magnitude and the spread of curvature over

time/age. The first step in the extraction of curvature (excesses or depths) is to estimate a baseline, which is a counterfactual scenario of vital rates in the absence of nonlinear age, period, or cohort effects under analysis (for instance, in the absence of nonlinear cohort effects for the Hispanic boomers' example).

Several methods are available to estimate a baseline from which it is possible to obtain curvature in vital rates. These approaches include applying interpolation techniques (Camarda 2012), extracting the irregularity using decomposition techniques (Remund, Camarda, and Riffe 2018), using detrended APC models (Chauvel, Leist, and Smith 2017), or simply detrending the smoothed vital rates over the selected perspective of change (i.e., over age, period, or cohort). There is no ideal generic method that can be applied because each demographic phenomenon and research question has specific underlying hypotheses that should be accounted for.

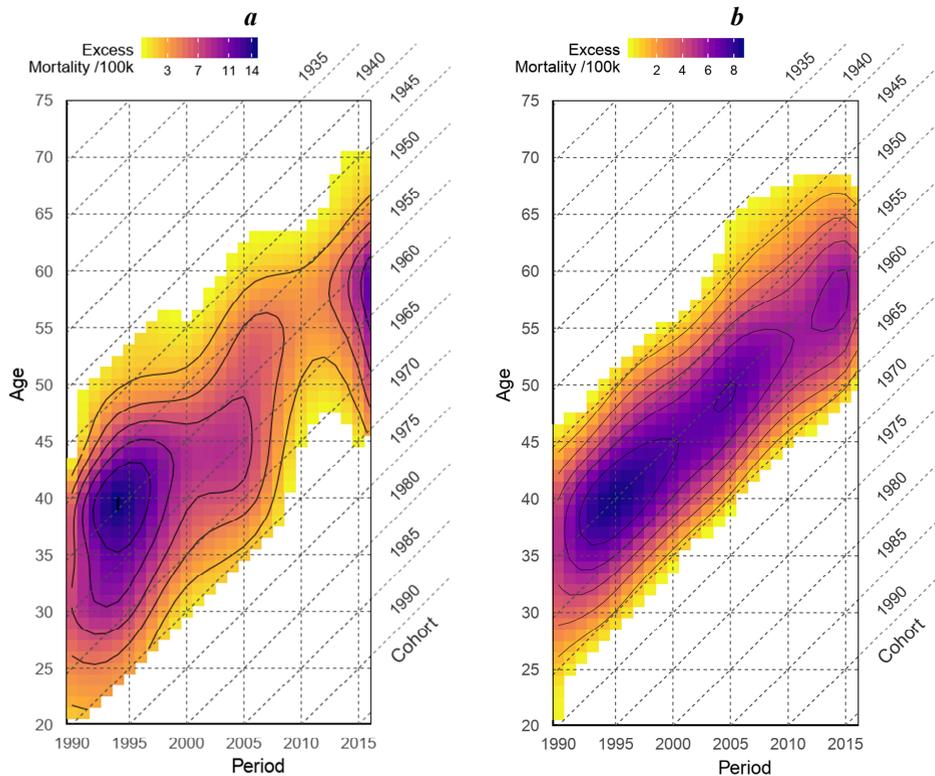
For the case of cohorts of Hispanic males in the United States who are disadvantaged in terms of drug-related mortality, we present two examples using the interpolation and dAPC approaches for the estimation of the mortality baseline. For the interpolation process, we excluded deaths pertaining to disadvantaged cohorts, who were previously estimated to be born between the advantaged 1940 and 1970 cohorts (see Figure 2), and interpolated the surface with these 31 cohorts removed. The estimation of the mortality baseline was done through the two-dimensional interpolation option available in the R package *MortalitySmooth*. Figure 4a depicts the excess mortality rates (per 100,000 population) among these cohorts over the interpolated baseline; i.e., the difference between the smoothed observed mortality rates and the hypothetical mortality that is predicted if mortality had developed according to a linear trend from the 1940 to the 1970 birth cohort.

Alternatively, to obtain a mortality baseline that accounts for all cohorts present in the observed data, we fitted a dAPC model with all linear trends attributed to age and period variations (cohort-detrended), and set all cohort terms to zero, while removing all cohort curvature components from the predicted baseline. Figure 4b presents the excess mortality among cohorts over a baseline that excludes cohort curvature; i.e., the difference between the smoothed observed mortality rates and the hypothetical mortality that is predicted if mortality had developed according to the overall linear trend for all of the cohorts.

Although they are similar in shape, the excess mortality estimates in Figures 4a and 4b are considerably different in magnitude and must be interpreted differently. As the magnitude and the shape of the excess mortality obtained from the interpolated baseline depend exclusively on the two cohorts selected for the interpolation, they are sensitive to the arbitrariness of the selection of these cohorts. In contrast, the excess mortality that is obtained from the baseline without cohort curvature components is

relative to the performance of all of the observed cohorts included in the observation window.

**Figure 4: Lexis surfaces of the excess drug-related mortality rates for male Hispanic boomers in the United States**



*Note:* Panel (a) is the excess mortality rates (/100k) estimated as the difference between the smoothed mortality rates and an interpolated baseline that omits the 1940–1970 cohorts from the Lexis mortality surface. Panel (b) is the excess mortality rates (/100k) estimated as the difference between the smoothed mortality rates and a baseline obtained from a dAPC model with the cohort terms set at zero; i.e., centered at the linear trend component of the cohort effects. In this case, the excess mortality is relative to the overall mortality trend across cohorts.

Compared to the curvature effects obtainable from statistical dAPC models, the plots of rate derivatives (Figure 3) and of trend divergence (Figure 4) are much more flexible in depicting the temporal dynamic of nonlinear fluctuations, as they allow the shape of curvature to move freely through the Lexis diagram, and depict patterns with a higher degree of fidelity to the observed data. Another important advantage of these

plots is that they depict general patterns that modulate the changing phenomena over a wide age and time frame. In a single image, it is possible to identify several irregularities potentially indicative of APC effects on the dynamics of a specific phenomenon of a population.

These plots are useful for the analysis of temporal patterns in a single phenomenon in a single population. However, when comparisons across several phenomena within a single population or of a single phenomenon across several populations are desired, it is necessary to construct a surface for each phenomenon/population analyzed. In addition to requiring considerable space, an important limitation of such comparisons is that displaying contrasting patterns across surfaces is visually difficult, even when surfaces are faceted.

### **3. The proposed visualization**

In the current paper, we follow in the tradition of the aforementioned literature by attempting to visually depict and contrast the nonlinear changes in demographic phenomena over the APC dimensions. Specifically, we demonstrate the value of making a broad-picture comparison of how the location of a curvature feature of a demographic phenomenon is changing over time. This curvature feature could be anything from the location of the cohort with maximum excess mortality by cause of death to the mode of age-specific fertility in different populations. In the former case, the visualization enables the comparison of the temporal patterns for many phenomena (causes of death) in one population on one plot. In the latter case, the aim is to compare the temporal patterning of a single phenomenon (age-specific fertility schedules) across a variety of populations or subgroups. While the simplest visualization depicts the changing location of these demographic phenomena (i.e., a curvature ridge or floor), the use of visual attributes such as color, size, and opacity allows for more information about the densities to be depicted; including, for instance, the change in the magnitude and the spread of curvature.

It should be noted that if the emphasis is placed on the maximum/minimum point of the curvature over time (ridge/floor), other local maximums/minimums would be neglected in the case of multimodal distributions. This information could be of great importance, and alternative plots could depict the dynamics of multiple modes over time. Moreover, the depiction of several features in the same plot also offers the possibility of analyzing the interrelation among them, such as the correlation between magnitude and spread. Nevertheless, the choice of the amount of information to display in the plots depends on the context of the research question, and should be made while

strategically considering the tradeoff between visual complexity and the clarity of the visualization (Munzner 2014).

Although the APC curvature plots proposed here contribute additional information that is not available in APC statistical models, they should be seen as complementary. Some estimable functions from statistical APC models are of great value, and are not obtainable by graphical analysis, such as the statistical significance of a particular pattern (Holford 1991). In Table 1, we summarize some of the advantages and limitations of the statistical and visual methods we have discussed.

**Table 1: Comparison of the properties of different methods for the analysis of nonlinear APC patterns**

Type	Method	Detection of curvature	Temporal dynamic of curvature features			Comparability across populations / phenomena	Statistical measures <sup>a</sup>
			Location	Magnitude	Spread		
Statistical models	dAPC models	+	-	-	-	+	+
	dAPC hysteresis model	+	-	+/-	-	+	+
Graphical tools	Surface of derivatives	+	+	-	-	-	-
	Surface of excess	-	+/-	+	+	-	-
	APC curvature plots	-	+	+	+	+	-

Note: (+) indicates that the method is able to perform the property; (+/-) that it does so to some degree, albeit imprecisely; and (-) that it is not able to perform the property.

<sup>a</sup>Some examples of these statistical measures are tests of statistical significance and confidence intervals.

In the following, we describe the procedure for constructing APC curvature plots for demographic phenomena, and apply the procedure to three empirical examples: (1) the excess mortality of baby boomer cohorts in the United States from drug-related causes across racial/ethnic populations, (2) the stability of the age location of the young adult mortality hump, and (3) the temporal patterning of age-specific cohort fertility peaks across countries.

## 4. Construction of the plot

For this kind of analysis, we suggest the use of the finest possible resolution for the data. The smaller the grid in the Lexis surface, the clearer and more precise the depiction of the temporal dynamic of demographic curvature will be. The construction of the APC curvature plot involves three main steps:

#### **4.1 Detection of curvature and the temporal section frame of interest**

As mentioned in the introduction, there are statistical methods (e.g., dAPC models in Figure 2) and pre-existing visualization tools (e.g., Lexis surfaces of changes in vital rates over age/cohort and over period/cohort in Figure 3) that allow for the detection of curvature patterns.

After some features of curvature (e.g., curvature ridge and floor) have been identified, their temporal positions in the Lexis diagram determine the temporal frame of interest. This framing is important because other “irregularities” located outside of this temporal section represent a potential source of noise for the analysis.

#### **4.2 Estimation of curvature features**

Remund and colleagues (2017) proposed three attributes of interest that could be used in describing the young adult mortality hump and provided examples of summary indices that could be used to measure them. These attributes are location (e.g., mode, mean, and median), magnitude (e.g., loss in life expectancy, years of life lost, and death counts), and spread (e.g., standard deviation and quantile). These dimensions and indices could be enriched so that they can be used to analyze other demographic phenomena.

The minimum requirement of a comparative APC curvature plot is the location measure (i.e., mode, mean, etc.). Other dimensions, such as magnitude and spread, are optional and complementary measures that could be used to enrich the comparative analysis, but that are not strictly necessary for the construction of a basic version of the plot.

In cases in which the deviance from the trend is also a local or an absolute maximum or minimum, the mode of the ridges and the valleys may be obtainable by extracting the age/period coordinates of the maximum or the minimum smoothed values within the temporal section frame. However, for cases in which these irregularities are not a local maximum or minimum, the estimation of their respective location, magnitude, and spread requires additional information pertaining to the divergence of vital rate estimates.

As discussed in section 2.2, several methods are available for estimating divergence in vital rates, and which method is appropriate depends on the underlying hypotheses regarding the demographic phenomenon and the research question to be addressed. The location, magnitude, and spread of curvature can be estimated from this divergence in vital rates. It is worth noting that all three features could be estimated

using age, period, or cohort perspectives depending on the temporal dimension that is of interest.

### **4.3 Translation of curvature attributes into visual properties of the plot**

The population or the demographic phenomenon to be compared in the Lexis surface is a categorical value that should be translated into color; preferably using a color-blind safe palette (as is done here), or into different point shapes for a black-and-white printout<sup>5</sup>. The location measures should be translated into the age and period coordinates in the Lexis diagram. To prevent unrelated curvatures from being fused, unifying these point coordinates by lines should be avoided.

The magnitude and the spread measures should be standardized and can be translated into opacity level and shape size.

## **5. Empirical application**

For illustrative purposes, we applied the procedure described above to construct APC curvature plots for three demographic outcomes with different temporal perspectives: (a) comparing excess mortality from drug-related causes across several racial/ethnic groups of boomers in the United States; (b) examining excess mortality among young adults; and (c) comparing cohort fertility rates across several countries. In this section, we describe how we estimated the location, the magnitude, and the spread features of curvature; and how we translated these features into an APC curvature plot that displays the attributes for each case. Although our main objective is to propose a visualization tool, we briefly comment on some of the determinants that might be driving the nonlinear APC patterns that are easily identifiable and comparable using this visualization technique.

### **5.1 Excess mortality from drug-related causes among boomers**

Previous exploratory and descriptive analyses have suggested that baby boomers in the United States have a disproportionate susceptibility to drug-related mortality throughout

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<sup>5</sup> When using different point shapes, equal areas must be given equal values of spread across the different point shapes employed. As this is usually not done by default in the plotting systems (e.g., *ggplot2*), a correction factor must be applied to adjust the proportionality of areas across the shapes.

their life course (Acosta et al. 2019a; Miech, Koester, and Dorsey-Holliman 2011; Zang et al. 2019). Thus, for this specific case, we are interested in comparing the attributes (location, magnitude, and spread) of the cohort mortality curvature of drug-related mortality for several racial/ethnic groups of US males in a single Lexis plot.

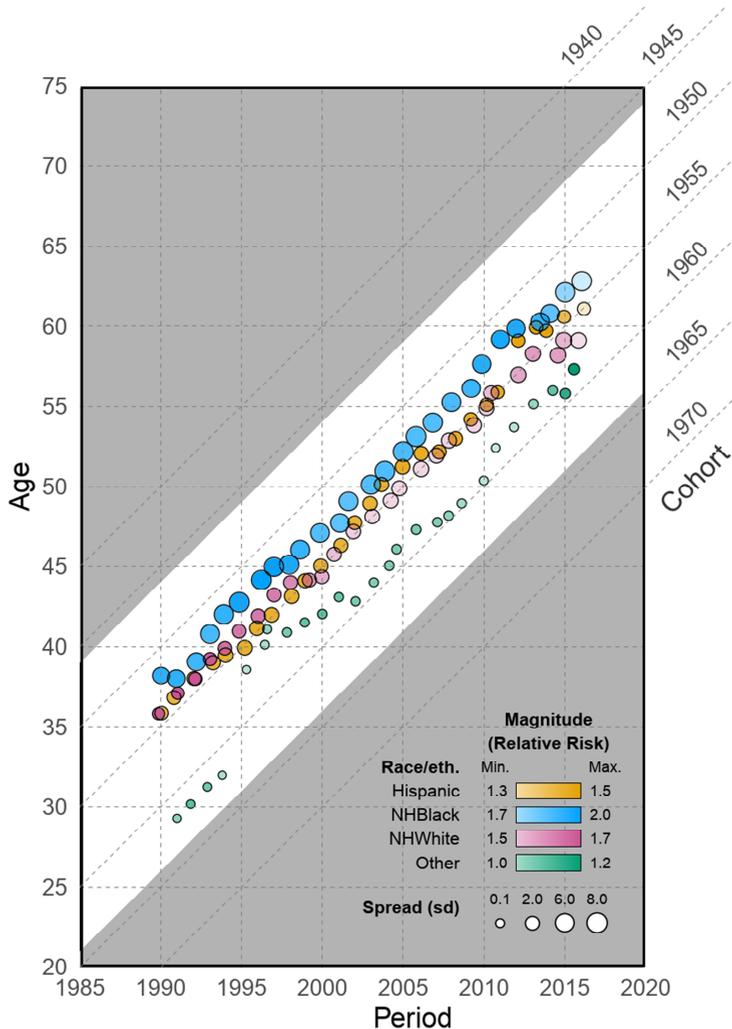
We use cause-specific mortality data and population estimates by calendar year, age, sex, race, ethnic group, and cause of death from 1990 to 2016 drawn from the US National Vital Statistics System (NVSS 2019a, 2019b). This dataset allows us to use single-year resolution in the estimation and visualization of nonlinear cohort effects. Here, we define drug-related mortality as deaths involving drug use registered in the categories of accidental and undetermined intent overdoses, or in the categories of mental or behavioral causes (i.e., ICD 10 codes F11-19, F55, X40-44, Y10-14).

Detecting the mortality curvature and measuring its features (location, magnitude, and spread) involved three distinct steps. First, we smoothed the entire drug-related mortality surface over period and age using two-dimensional P-splines, with the smoothing parameters optimized by AIC, for all male racial/ethnic groups; as presented in Figure 1b for the Hispanic ethnic group. Second, from these smoothed rates, we calculated the change in mortality from age  $x$  to age  $x+1$  (from cohort  $k$  to cohort  $k-1$ ) for each year (Figure 3a), and the change in mortality from period  $t$  to period  $t+1$  (from cohort  $k$  to cohort  $k+1$ ) for each age (Figure 3b). The ridges and floors of curvature were identified with black lines used to indicate the ages at which the mortality change was zero. In the derivative over age/cohort (plotted in Figure 3a), the curvature ridge indicating the maximum level of drug-related mortality is mostly diagonal, following the cohorts born between 1950 and 1960 throughout their life course. Third, using a dAPC model, we estimated the excess mortality relative to the overall cohort average risk for each racial/ethnic group; as the example presented in Figure 4b for US Hispanic males shows. From these excess mortality estimates, we were able to extract information about the temporal dynamic of the location (age/cohort with the largest mortality relative risk), the magnitude (largest mortality relative risk compared at the curvature), and the spread (steepness of the excess mortality compared to the mortality of the surrounding ages) of the curvature during the 1990–2016 period. We can see that the curvature trajectory is not completely straight, and that its magnitude and spread are not constant over time. As discussed in the introduction, these changes in the location, magnitude, and spread over age/time cannot be identified from the dAPC estimates, such as those depicted in Figure 2. To compare the curvature ridge pattern differences across racial/ethnic groups, we plotted in Figure 5 the following attributes of the excess drug-related mortality among boomers on an APC curvature plot:

1. *Category*: Each observed racial/ethnic group was identified by a distinct color, or by a different point shape in the case of a black-and-white printout (see Figure A-1).
2. *Location*: For each calendar year, we plotted the age/cohort location of the maximum relative risk of drug-related mortality.
3. *Magnitude*: The relative risk compared to the baseline value for each point was translated into opacity. Note that the opacity scale reflects the difference in relative risk within each racial/ethnic group. We use a relative scale because it allows us to better identify the variation in magnitude over time and for each group. The absolute differences in magnitude across racial/ethnic groups can be seen by comparing the minimum and maximum values reached by group. This range in magnitude is indicated in the legend for each group.
4. *Spread*: The standard deviation of the hump in relative risk for each period was translated into point size.

In this case, we were not only able to extract and summarize the differences in location from four different plots in a single visualization; we were able to compare additional outputs to the rate of mortality change plots, such as the magnitude (relative risk) of the mortality curvature ridge compared to the expected values and the spread (standard deviation) of the curvature. Thus, we were able to clearly identify the disadvantaged cohorts for all of the racial/ethnic groups by observing the alignment of the points along a diagonal line. Our finding that the degree of cohort disadvantage, relative to the overall cohort average, was greater among non-Hispanic black boomers than among other race groups is reflected in the larger values of relative risk among the former group (i.e., 1.7 to 2.0). Note that we picked a cause of death with particularly strong cohort patterning. For other causes of death, such as cerebrovascular diseases, nonlinear excess mortality risks would not follow cohort patterns; thus, the depicted points would not fall along a diagonal line.

**Figure 5: APC curvature plot of the features of excess drug-related mortality among four racial/ethnic groups of boomer males in the United States**



Notes: The coordinates of the points indicate the location of the curvature ridge over time (i.e., the modal age/cohort with the excess mortality in each single-year period). The magnitude, indicated by the color opacity, is measured as the relative risk of the death rate in the modal age/cohort to the corresponding death rate in the baseline. The minimum and maximum levels of relative risk that the curvature ridge of each racial/ethnic group reached during the period under observation are indicated in the legend. The spread, indicated by the point size, is estimated as the standard deviation of the curvature in each single-year period. The white band indicates the baby boomer cohorts (i.e., born between 1946 and 1964).

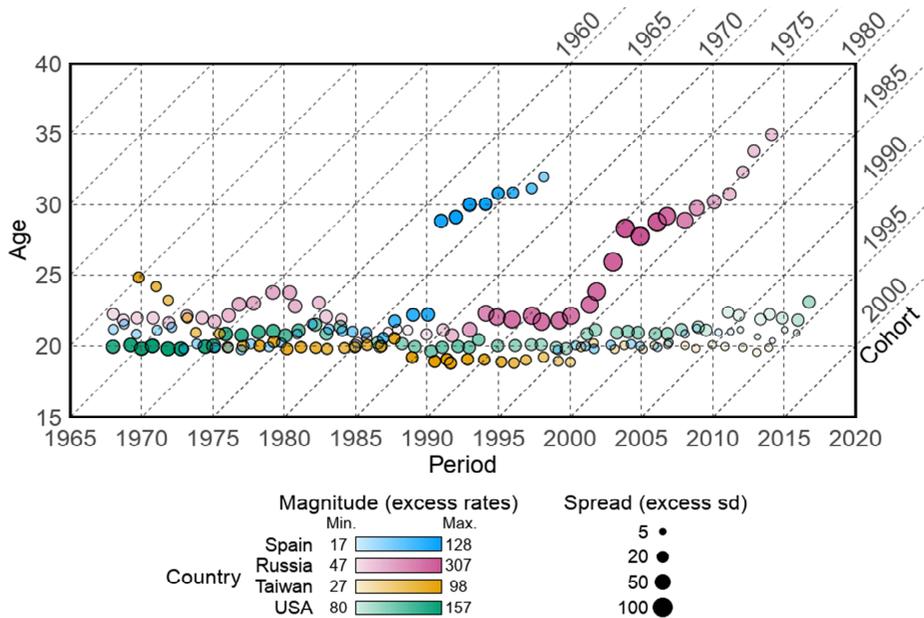
## 5.2 Young adult mortality hump

Excess mortality in young adults – also known as the accident or young adult mortality hump – is a well-known feature of the age structure of most mortality regimes, particularly among males. Goldstein (2011) presented a plot of the hump peaks over time for several countries, and Remund and colleagues (2018) decomposed and plotted the contributions by cause of death in order to analyze several features of the hump in the United States. Here, we used mortality data for certain countries drawn from the Human Mortality Database (HMD 2019) to compare not only the curvature ridges of the hump, but the additional features of their changes in magnitude and spread over time and across different countries. As in the method employed by Goldstein, we defined the hump as the difference between the cross-sectional period smoothed mortality rates and an interpolated mortality baseline between ages 10 and 40. We performed the smoothing and the baseline interpolation using the methods we previously applied to obtain Figure 4a. However, unlike the example presented in Figure 4a, we interpolated the hypothetical baseline with excess age-related mortality rather than cohort-related mortality.

We compared the young adult mortality hump in Spain, Russia, Taiwan, and the United States during the 1965–2016 period. Figure 6 shows the variation over time of the modal age of the hump, its magnitude measured by excess death rates (per 100,000 population), and the spread of the hump measured by standard deviation. A black-and-white printout of this plot is presented in Figure A-2.

In contrast to the more regular location patterns we see for Taiwan and the United States, we observe in this plot an age-period interaction effect for Spain around age 30 during the 1990s and a cohort effect for Russia among the cohorts born between 1975 and 1980. In the case of Spain, this finding of an age-period interaction effect is consistent with evidence indicating that in this period, Spain was the western European country with the highest incidence of HIV infection, which was mainly driven by the sharing of contaminated needles during the “heroin boom” of the 1980s (Valdes and George 2013). In the case of Russia, the cohort pattern could be indicative of a long-term mortality disadvantage for those cohorts who entered adulthood in the early 1990s, when the country was undergoing a profound sociohistorical transformation, as well as a severe alcohol abuse epidemic (Keenan et al. 2015). This pattern is consistent with a cohort pattern found in Belarus (not shown here), a country that experienced similarly far-reaching contextual changes during the same period. It is noteworthy that part of the age-period interaction effect depicted in the APC curvature plot for Spain would be attributed to a cohort effect if a statistical approach would have been used. This is because, unlike the visual tools, the existing APC statistical methods are unable to make the distinction between age–period interaction effects and sustained cohort effects.

**Figure 6: APC curvature plot of the features of excess mortality in young adult males in four countries**



Notes: The coordinates of the points indicate the location of the curvature ridge over time (i.e., the modal age/cohort of the excess mortality by single-year period). The magnitude, indicated by the color opacity, is the excess death rates (/100k), calculated as the difference between the death rate in the modal age/cohort and the corresponding death rate in the baseline. The minimum and maximum excess mortality rates reached by the curvature ridge of each country during the period under observation are indicated in the legend. Finally, the curvature spread, indicated by the point size, is estimated as the standard deviation of the curvature in each period.

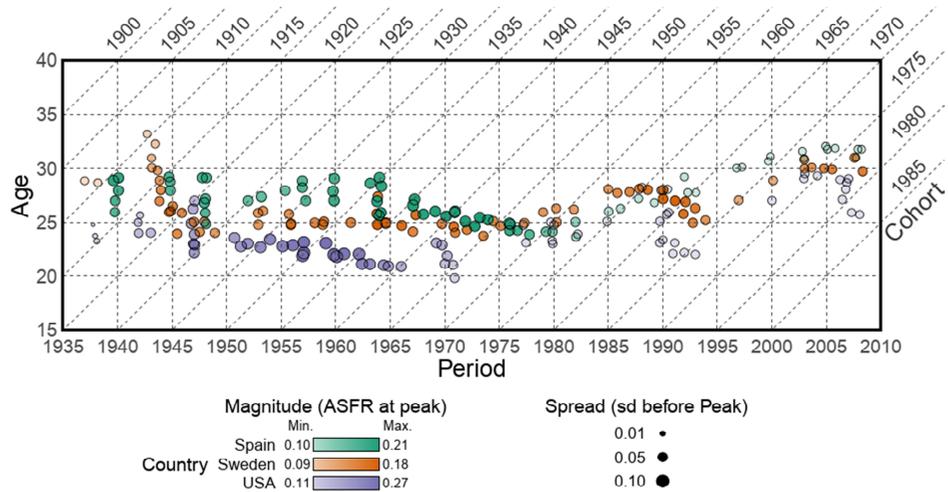
### 5.3 Cohort fertility rate

As a final example to show the applications beyond mortality, we plotted some aspects of fertility behavior along cohorts. Figure 7 depicts the age-specific fertility outcomes of the cohorts born between 1905 and 1985 in Spain, Sweden, and the United States. Figure A-3 displays a black-and-white version of this plot. Cohort age-specific fertility rates (ASFR) were obtained from the Human Fertility Database (HFD 2019) and extracted using the R package *HMDHFDplus* (Riffe 2015).

In this case, we plotted the modal age/period of fertility for each cohort (location), the ASFR pertaining to this mode (magnitude), and the standard deviation before that age/period (spread). The standard deviation before the mode was used to enable us to

consider more recent cohorts who are likely to have reached the modal age in fertility but have not yet completed their childbearing. As mentioned previously, any number of indices could be used to capture the three dimensions of location, magnitude, and spread.

**Figure 7: APC curvature plot of cohort fertility rate peaks in three countries**



*Notes:* The coordinates of the points indicate the location of the curvature ridge along the cohort (i.e., the modal age/period of the ASFR by single-year cohort). The magnitude, indicated by the color opacity, is measured as the ASFR mode by cohort. The minimum and maximum ASFR reached by the modal age/period of each country are indicated in the legend. The curvature spread, indicated by the point size, is estimated as the standard deviation of the curvature before the modal age/period.

This plot highlights a number of interesting features in the fertility dynamics of these three countries. We see strong evidence of period effects for many of the countries in which the fertility booms were synchronous for several cohorts (e.g., Sweden in 1943–1945; the United States in 1947, 1990, and 2007; and Spain in 1940, 1948, and 1964). The plot also illustrates the sustained differences in modal ages of fertility over time (e.g., the ages are consistently youngest for the United States and oldest for Spain; with an exceptional shift between Sweden and Spain during the 1980s). The overall decline in fertility in Spain and the United States is also illustrated by the fading peak ASFR over time, whereas this pattern is irregular in Sweden. Lastly, the concentration of fertility at more advanced ages is reflected in the decline in the standard deviation before the modal age at fertility in all three countries.

## **6. Conclusions**

In this paper, we discussed the advantages of using visualization tools on Lexis diagrams for analyses of age-period-cohort nonlinear effects of vital rates rather than mathematical models, as well as some of the limitations of these tools. We proposed the APC curvature plot to enrich the analysis of irregularities in vital rates that are indicative of nonlinear age, period, or cohort effects. We argue that compared to mathematical models and other Lexis plots, this visual display provides a higher level of flexibility because it allows us not only to depict the dynamics of the location, the magnitude, and the spread of temporal effects over time together; but to contrast different populations or subtypes of demographic phenomena in a single visualization. We outlined the process that can be used to construct APC curvature plots for the analysis of nonlinear APC effects. Using vital rates, we provided some examples of how this approach can be applied by analyzing cohort effects on drug-related mortality by racial/ethnic groups within the United States; age effects on young adult mortality in Russia, Taiwan, Spain, and the United States; and age/period effects on fertility in Spain, Sweden, and the United States. Finally, we have provided R-code that can be used to reproduce these examples.

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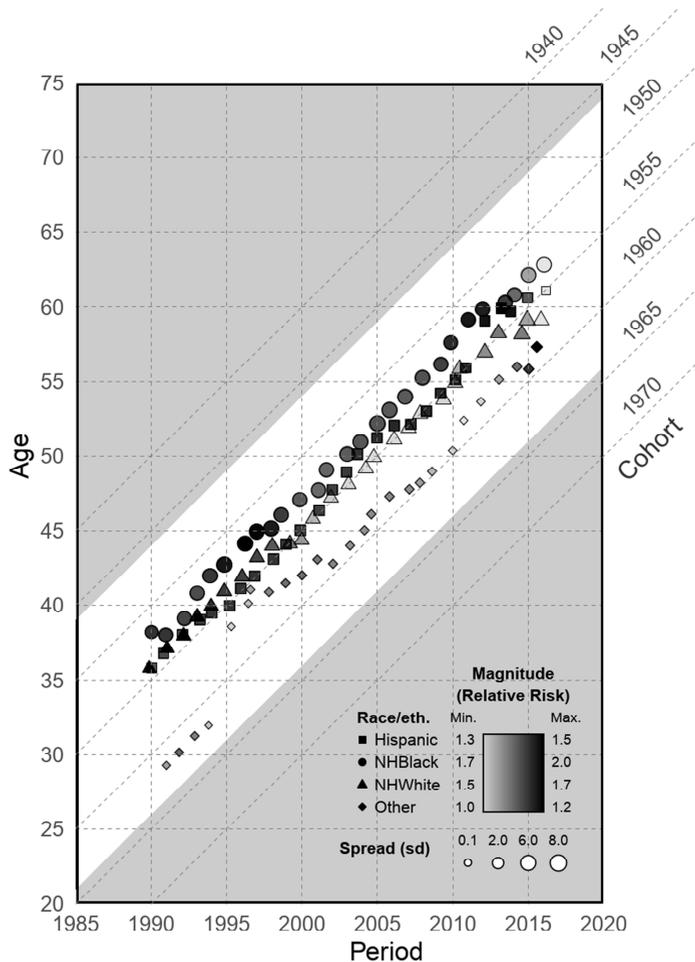
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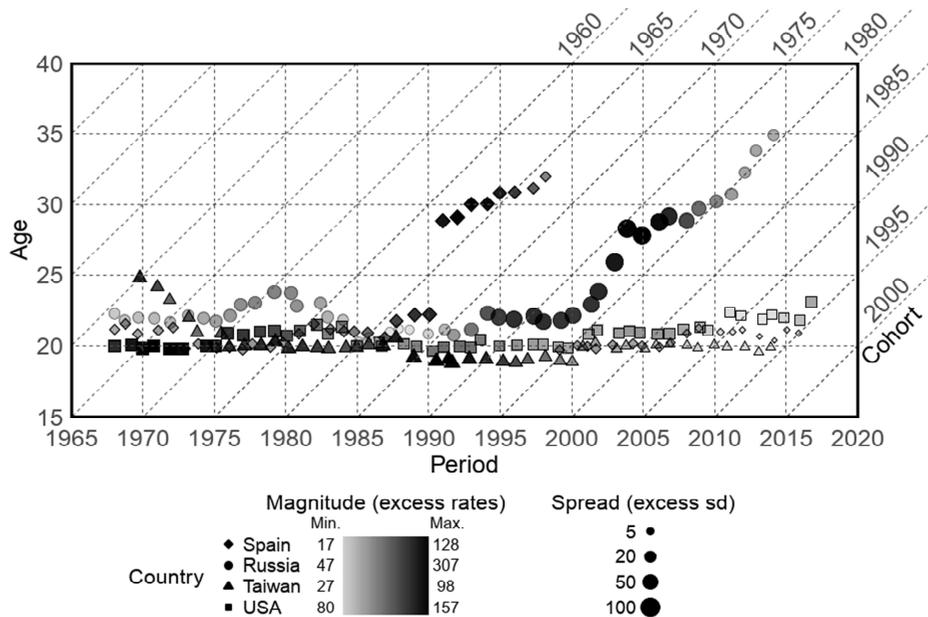
## Appendix: Black-and-white printout of the Lexis plots

**Figure A-1: B&W APC curvature plot of the features of excess mortality from drug-related causes in four racial/ethnic groups of boomer males in the United States**



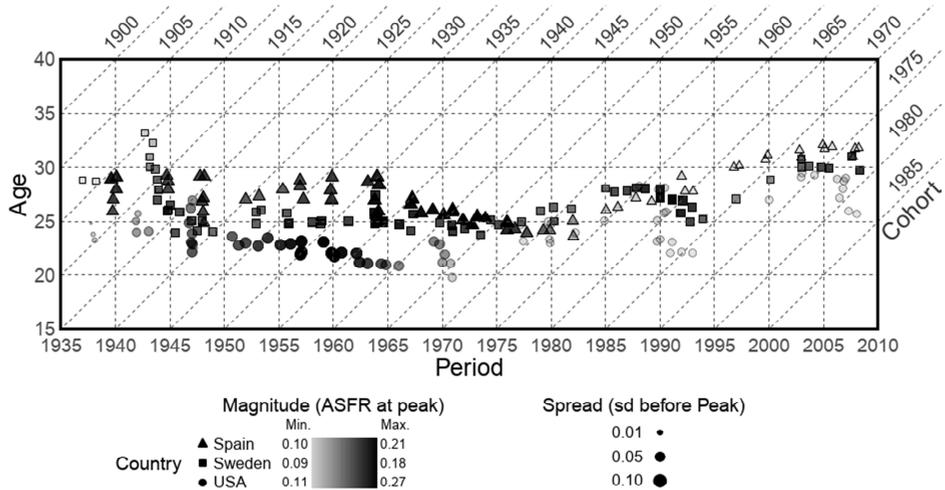
*Notes:* The coordinates of the points indicate the location of the curvature ridge over time (i.e., the modal age/cohort with the excess mortality in each single-year period). The magnitude, indicated by the opacity, is measured as the relative risk of the death rate in the modal cohort to the corresponding death rate in the baseline. The minimum and maximum levels of relative risk that each racial/ethnic group reached during the period under observation are indicated in the legend. The spread, indicated by the point size, is estimated as the standard deviation of the curvature in each single-year period. The white band indicates the baby boomer cohorts (i.e., born between 1946 and 1964). A correction factor was applied to adjust the proportionality of areas across the shapes.

**Figure A-2: B&W APC curvature plot of the features of excess mortality in young adult males in four countries**



Notes: The coordinates of the points indicate the location of the curvature ridge over time (i.e., the modal age/cohort of the excess mortality by single-year period/cohort). The magnitude, indicated by the opacity, is the excess death rates (/100k), calculated as the difference between the death rate in the modal age/cohort and the corresponding death rate in the baseline. The minimum and maximum excess mortality rates reached by country during the period under observation are indicated in the legend. Finally, the curvature spread, indicated by the point size, is estimated as the standard deviation of the curvature in each period. A correction factor was applied to adjust the proportionality of areas across the shapes.

**Figure A-3: B&W APC curvature plot of cohort fertility rate peaks in three countries**



Notes: The coordinates of the points indicate the location of the curvature ridge along the cohort (i.e., the modal age/period of the ASFR by single-year cohort). The magnitude, indicated by the opacity, is measured as the ASFR mode by cohort. The minimum and maximum ASFR reached by the cohorts of each country are indicated in the legend. The curvature spread, indicated by the point size, is estimated as the standard deviation of the curvature before the modal age/period. A correction factor was applied to adjust the proportionality of areas across the shapes.

