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Publication Date

2015-02-10

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Average age at death in infancy and infant mortality level: reconsidering the Coale-Demeny formulas at current levels of low mortality

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Abstract

The longterm historical decline in infant mortality has been accompanied by increasing concentration of infant deaths at the earliest stages of infancy. The influence of prenatal and neonatal conditions has become increasingly dominant relative to postnatal conditions as external causes of death such as infectious disease have been diminished. In the mid-1960s Coale and Demeny developed formulas describing the dependency of the average age of death in infancy on the level of infant mortality from data obtained up to that time.

Almost at the same time as Coale and Demeny's analysis, as shown in this paper, in the more developed countries a steady rise in average age of infant death began. This paper demonstrates this phenomenon with several different data sources, including the linked individual birth and infant death datasets available from the US National Center for Health Statistics and the Human Mortality Database. A possible explanation for the increase in average age of death in infancy is proposed, and modifications of the Coale-Demeny formulas for practical application to contemporary low levels of mortality are offered.

Key words:

infant mortality, Coale-Demeny formulas, average age at death in infancy, concentration of infant deaths, concentration of infant deaths in the neonatal period

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Introduction

During the period from the 1920s to the 1970s infant mortality decline in Europe and other industrialized countries was accompanied by concentration of infant deaths in the neonatal period. The distribution of infant deaths became more and more highly skew. The average length of life for infants who died during the first year in low-mortality countries was less than 0.25, and exhibited systematic decline as the infant mortality rate (IMR) decreased.

The average duration until death for infants who die is an important parameter in life table construction. It is used for calculation of infant mortality rates from infant death rates and other life table functions at age 0, especially if Chiang's simple formula (Chiang 1978) is employed. Where direct data on the average age of death in infancy are not available, a set of formulas known as the "Coale-Demeny formulas"(C-D formula) that describe the relation between the infant mortality rate and the average age of infant death ${}_1a_0$ have been the most widely used to estimate ${}_1a_0$ from ${}_1q_0$. In the 1980s, the C-D formula was included in the UN software package for mortality measurement MORTPAK (1988). In 2001 it was advocated as the basic formula for calculation of the average age of infant death and for calculation of ${}_1q_0$ based on ${}_1M_0$ (Preston, Heuveline, Guillot 2001:47). Finally it became a basic formula employed in the Human Mortality Database (HMD) (Wilmoth et al. 2007) that was launched in May 2002. The C-D formula continues to be recommended for calculation of ${}_1a_0$ in textbooks at the present time.

Preston et al. gave the next explanation of the C-D formulas: "Generally speaking, the lower the level of mortality, the more heavily will infant deaths be concentrated at the earliest stages of infancy; the influence of the prenatal and perinatal environment becomes increasingly dominant relative to the postnatal environment". Nevertheless, as we will show later at the level of the infant mortality rate about 0.017-0.022 this concentration stopped and started to

reverse itself. This conclusion is based on national life tables for countries that do not use the C-D formula (or MORTPAK) and do not accept any conditional estimate of ${}_1a_0$, (For example in all official life tables for France ${}_1a_0$ is equal to $\frac{1}{2}$; hence, we exclude France).

To be specific, the C-D formula indicates that if ${}_1q_0 < 0.011$, the average age of infant death is less than 0.8 month. However in the US life tables for period 2000 and later with ${}_1q_0 < 0.008$ (Arias et al. 2010) the average age of infant death exceeds 1.3 months. In Japan (2007)³, with ${}_1q_0 = 0.00274$ for males, and for females 0.00244, the average age of infant death is more 2 months. We found in the Human Life Table database (HLD)⁴ 277 national level life tables for 23 countries or regions (Australia, Austria, Canada, Chile, Cuba, Germany, Greece, Hong Kong, Hungary, Ireland, Italy, Japan, New Zealand, Poland, Portugal, Singapore, Slovenia, Spain, Taiwan, United States of America) with male infant mortality rates less 0.010 that have been published with all details needed to estimate the exact value of average age of infant death, ${}_1a_0$. In all these life tables except those for Italy, the average age of infant death is more than 1 month. However all countries that use MORTPAK for calculation of national life tables also use C-D formulas. This circumstance does not appreciably distort the table indicators and has almost no effect on life expectancy. Nevertheless, the presence in a life table of an obviously wrong indicator seems to us undesirable.

The history of the C-D formulas is as follows. In the early 1960s Ansley J. Coale and Paul Demeny developed for their influential series of the regional model life tables an algorithm for calculating ${}_1a_0$, the average age of death in age interval (0,1) for infants who died in the interval (1966:(20)). This important parameter is necessary for beginning the life table. The number of person-years lived at in the interval from birth to exact age $1, {}_1L_0$, is related to l_1 ,

³ <http://www.mhlw.go.jp/english/database/db-hw/lifetb07/female.html>

⁴ <http://www.lifetable.de/>

the number of survivors to age 1 and l_0 , the hypothetical number of births, by the actuarial relation ${}_1L_0 = {}_1a_0 \cdot l_0 + (1 - {}_1a_0) \cdot l_1$, which leads in turn to the following formula for deriving the infant mortality rate, ${}_1q_0$, from the death rate observed in the age interval from 0 to 1, ${}_1m_0$:

$${}_1q_0 = \frac{{}_1m_0}{1 + (1 - {}_1a_0) \cdot {}_1m_0}.$$

Table 1. Ansley J. Coale and Paul Demeny formulas for estimation of average age of death in infancy based on the infant mortality rate q_0

Regional family of model life tables	For females	For males
if infant mortality rate ${}_1q_0 \geq 0.1$		
"West," "North," "South"	0.35	0.33
"East"	0.31	0.29
if infant mortality rate ${}_1q_0 < 0.1$		
"West," "North," "South"	$0.050 + 3.000 \cdot {}_1q_0$	$0.0425 + 2.875 \cdot {}_1q_0$
"East"	$0.010 + 3.000 \cdot {}_1q_0$	$0.0025 + 2.875 \cdot {}_1q_0$

Note. Coale and Demeny denoted average age of death in the age interval 0-1 with k_0 but we have adopted a parallel notation.

These formulas for ${}_1a_0$ were a minor detail within the Coale-Demeny system of model life tables, which has been variously employed to describe relations between levels of fertility and mortality on the one hand, and population structure on the other hand, and which has been widely applied in demographic analysis of Third World countries lacking reliable vital statistics, as well as historical demography of European populations. Somewhat ironically, the

formulas for ${}_1a_0$ seem to have found more widespread use than the model life table system itself, being employed in the construction of life tables for many analyses which have not otherwise involved the model life tables.

In the 1970s related formulas were developed for calculation average age of infant death based on the central death rate at age 0: ${}_1M_0$ (Arriaga, Anderson, and Heligman 1976). They used only the higher (for regions "West," "North," "South") variant of the formula for each sex separately. These formulas were installed under the name of the "Coale Demeny formula" and the widespread use of the C-D formula began.

In the analysis below, we attempt to estimate the actual dynamics of the average age of infant death based on vital statistics data for the United States. These data permit us to demonstrate that the decline of average age of infant death synchronous with infant mortality decline becomes interrupted when the infant mortality rate attains a level of about 10 per 1000 newborn infants. Employing the unique datasets of linked individual birth and infant death records available from the National Center for Health Statistics of the U.S. Centers for Disease Control and Prevention we seek to explain why major infant mortality decline can be combined with relatively high age of infant death.

1. Data and methods

1.1. Methods of calculating average age of death in infancy

For precise computation of the average age of infant death it is necessary to have either the aggregate distribution deaths by age in great detail (e.g. days), or, at a minimum, in days during first month of life and weeks for months 2-12; microdata containing this detail would also be adequate, as in the US case. Regrettably, as a result of infant mortality decrease, national statistical offices as well as international organizations have reduced the amount and

degree of detail in the infant mortality data they publish. Unlike the 1960s and 1970s, in modern publications of infant mortality statistics age in infancy is often provided only in three age groups: 0-6 days, 7-27 days, and 28 or more days.

Fortunately there is an approximation formula for calculation of the average age of death in infancy in an annual birth cohort. This formula uses only numbers of deaths by Lexis triangles. According to this formula, the average age of infant deaths equals the ratio of the number of deaths in the upper Lexis triangle $D(0, t, t+1)$ to the total number of deaths at age 0 in the birth cohort to which these deaths refer:

$$(1) \quad {}_1a_0 = D(0, t, t+1) / (D(0, t, t) + D(0, t, t+1)),$$

where $D(x, y, t)$ is number of death in age x in the cohort of y year of birth during calendar year t . For birth cohort y in the age interval x to $x+1$, the triad $(x, y, t+1)$ corresponds to the upper triangle, while $D(x, y, t)$ represents the lower triangle. The formula is correct under two conditions: 1) the distribution of births throughout the year of birth is uniform; 2) the probabilities of survival to ages less than 1 do not depend on the date of birth within the year.

These assumptions are typical for demographic calculations. In a sense this formula is equivalent to one of the basic formulas of the cohort component method of population projections: the number of newborn infants survived to the end of the calendar year of birth is

equal $b \cdot \frac{{}_1L_0}{l_0}$ (b stands the total number of newborn infants in the given calendar year);

alternatively, if $l_0 = 1$, then this number is $b \cdot {}_1L_0$ (cf. Keyfitz and Flieger 1971, in which 5 should be modified to 1 because we are dealing with single years in this instance). The total number of cohort survivors to age 1 is $b \cdot l_1$. Thus the righthand side of (1) derives from the methodology of population projection: $(b \cdot {}_1L_0 - b \cdot l_1) / (b - b \cdot l_1) = ({}_1L_0 - l_1) / (1 - l_1)$. Inserting into the last expression the formula for ${}_1L_0$ (in the square brackets) we obtain

$$\frac{[{}_1a_0 \cdot 1 + (1 - {}_1a_0) \cdot l_1] - l_1}{1 - l_1} = \frac{{}_1a_0 \cdot (1 - l_1)}{1 - l_1} = {}_1a_0.$$

A more thorough derivation of this equation is given in Appendix 1.

The results obtained through this equation (henceforth termed “the triangle-based formula”) are compared to those obtained by direct calculation based on observed dates of birth and death in infancy through record linkage in the analysis below.

1.2. Data

Our analysis draws upon several datasets, the first being the Human Mortality Database (HMD) available at <http://www.mortality.org>. The HMD is an online database containing detailed data on period and cohort mortality and survival. Currently it includes 37 countries with reliable mortality statistics. All numbers of deaths in the HMD are presented as numbers of deaths in Lexis triangles. However, for the most part these triangles contain the results of splitting numbers of deaths by year and age of death into Lexis triangles (Wilmoth et al. 2007:11-14) in a manner that in many cases predetermines the dependency of ${}_1a_0$ on ${}_1q_0^i$.

The present analysis investigates this relationship and requires data for countries and periods originally received in tabulated form by year of birth, year of death and age at death, or microdata in which these three variables are indicated for each infant death included.

However, even in this case numbers of deaths presented in the HMD may still be adjusted, for example as a result of prior redistribution of deaths of unknown age. To avoid possibility of artifacts, we have decided to use the initial (in HMD terms “raw”) numbers of deaths. We have collected all raw numbers of deaths tabulated by Lexis triangle presented at HMD website. (Downloaded February 3, 2011).

Some raw data were excluded from our research because they were corrected previously due to some defects in the initial statistical data. The main problem is “false stillbirths”, a situation

where infants that were born alive but died before the birth was registered were reported as stillbirths rather than infant deaths. This has led us to exclude from our dataset observations for France prior to 1975 (Glei et al. 2014) and Netherlands prior to 1950 (Jasilionis 2011). Data for Taiwan were excluded due to “systematic under-registration of infant deaths” (Canudas-Romo et al. 2012). In addition, some East-European countries kept up to the 1960s or later the League of Nations’ definition of live births and stillbirths adopted in 1925, according to which deaths of infants with body mass less than 1000 g. at age less than 7 days were registered as stillbirths: Bulgaria (completely) (Philipov, Jasilionis 2012), the Czech republic (until 1964) (Rychtarikova, Jasilionis, Grigoriev 2012), East Germany (completely) (Scholz, Jdanov, Kibele 2013), Estonia (until 1992) (Jasilionis 2013), Poland (until 1994) (Fihel, Jasilionis 2011), the Slovak Republic (until 1964) (Mészáros, Jasilionis 2011), Ukraine (completely) (Pyrozchkov et al. 2011). Finally, we exclude data for Iceland, where the number of infant deaths is very small and sometimes ${}_1a_0 = 0$. In this manner we have assembled the initial numbers of deaths before age 1 in Lexis triangles for 1001 cohorts that were born in the period 1901-2008 in the following 22 countries: Austria (1971 - 2007), Belgium (1941 - 2008), Canada (1950 - 2006), Czech republic (1965 - 2007), Denmark (1921 - 2007), Estonia (1992 - 2008), Finland (1917 - 2008), France (1975 - 2008), Germany (1991 - 2007), Hungary (1950 - 2005), Italy (1929 - 2005), Japan (1950 - 2008), New Zealand (1980 - 2007), Norway (1993 - 2007), Poland (1995 - 2008), Portugal (1980 - 2008), Slovak Republic (1965 - 2007), Slovenia (1983 - 2008), Spain (1975 - 2005), Sweden (1901 - 2007), USA- (1959 - 2006).

The second source our analysis draws upon a compilation of the cohort linked birth-infant death datasets from the US National Center for Health Statistics, available for years 1983-1991, and 1995-2004, at http://www.cdc.gov/nchs/data_access/Vitalstatsonline.htm. These files match each death record derived from the death certificate of an infant with the corresponding birth certificate, where possible. The linked data allow us to determine exact

ages at death along with other items of interest, such as detailed cause of death and race of mother.

From the linked birth-infant death data files we calculated the total number of deaths and the average exact age at death in infancy using formula (1) for the period under consideration for the total population of the USA from all causes of death and from 5 selected groups of causes: certain conditions originating in the perinatal period (ICD9 codes B45 or ICD10 codes P00 - P96); congenital malformations, deformations and chromosomal abnormalities (codes B44 or Q00 - Q99); diseases of the respiratory system (B31 or B32, J00 - J99); and sudden infant death syndrome (B466 or R95).

These data should help us to tackle the puzzle as to why the decline of exogenous mortality has been associated with a relatively stable average age of infant death.

In 1999, the US NCHS shifted to the ICD 10 classification of cause of deaths in its annual mortality microdatasets. Prior to the implementation of ICD 10, NCHS conducted a comparison study in which a sample of death records filed in 1996 was coded according to both the ICD 9 and ICD 10 classifications. From a crosstabulation of the parallel ICD9/ICD10-coded data from the comparison study, available on the NCHS website, we have confirmed (Appendix 2) that the change in cause of death classifications would be unlikely to result in a major change in the distribution of deaths by cause in terms of the aggregated cause of death groupings we employ.

The triangle-based formula for average age of infant death assumes that the distribution of births during the year in question was uniform. To assess the accuracy of this assumption, we subdivided annual totals of reported births by month of birth in the USA from the Human Fertility Database available at <http://www.humanfertility.org/>.

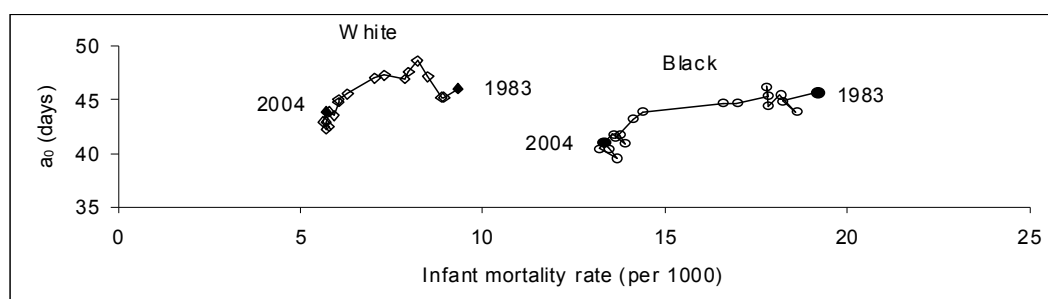
In the present analysis, when making comparisons of US infant mortality by race, we employ categories consistent with the pre-1997 US race definition throughout

In addition, we used data on the distribution of infant deaths by cause for some of the countries mentioned in order to approximate the probability of dying from leading causes of death from the WHO Mortality Database available at <http://www.who.int/whosis/mort/download/en/index.html>.

2. Results

During the period 1983-2004 the US cohort infant mortality rate for both sexes combined declined from 10.9 to 6.8. However, the average age of infant death was relatively stable at a level of 41.9-47.5 days, some 17-21 days more than what is indicated by the C-D formula (Table 2). Remarkably, almost the same average age of infant death was observed also for all groups categorized by race of mother. Very small declines in ${}_1a_0$ were observed for both the white and black subpopulations, but the rather large difference in the infant mortality rates of these two racial groups appears not to be associated with any substantial differences in ${}_1a_0$ (Figure 1).

Figure 1. Dependence of the average age of infant death on the infant mortality rate, by race of mother in birth cohorts 1983-1991, and 1995-2004 in the USA .



Note: Black points correspond to the beginning and the end of the period of observation.

Table 2. Infant mortality rates (IMRs) and three estimates of the average age of infant deaths in the birth cohorts of 1983-1991 and 1995-2004 , by race of mother in the USA.

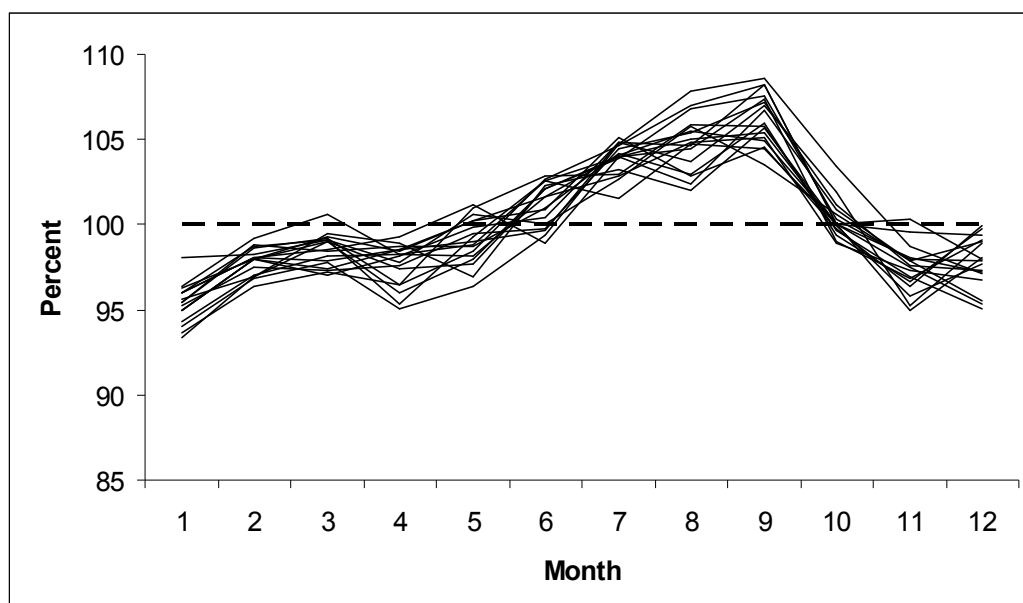
Year of birth	Total				White				Black				Other			
	IMR per 1000	Average age of infant deaths (days)			IMR per 1000	Average age of infant deaths (days)			IMR per 1000	Average age of infant deaths (days)			IMR per 1000	Average age of infant deaths (days)		
		Actual ⁵	Triangle- based	C-D formula		Actual	Triangle- based	C-D formula		Actual	Triangl e-based	C-D formula		Actual	Triangle -based	C-D formula
1983	10.9	46.1	50.3	28.5	9.3	46.0	50.2	26.8	19.2	45.7	49.4	37.4	10.7	50.7	57.1	28.3
1984	10.4	45.3	48.7	28.0	8.9	45.2	48.8	26.4	18.2	44.8	47.4	36.4	10.6	49.4	55.9	28.2
1985	10.4	45.0	47.6	28.0	8.9	45.3	47.5	26.3	18.6	43.9	47.8	36.8	10.2	48.0	49.2	27.7
1986	10.1	46.9	49.4	27.6	8.5	47.2	49.5	25.9	18.2	45.4	48.6	36.4	10.4	51.1	54.5	28.0
1987	9.8	47.5	50.5	27.3	8.2	48.6	52.1	25.6	17.8	44.4	45.9	36.0	9.6	50.7	56.9	27.1
1988	9.6	47.5	50.2	27.1	8.0	47.6	49.1	25.3	17.8	46.2	50.7	35.9	8.7	55.3	64.9	26.1
1989	9.5	46.8	50.1	27.0	7.8	47.0	49.8	25.2	17.8	45.3	49.2	35.9	8.8	55.2	61.0	26.2

⁵ By “actual” we mean calculated from microdata.

1990	8.9	46.8	49.8	26.3	7.3	47.3	50.3	24.6	16.9	44.6	46.5	35.0	7.9	57.1	66.2	25.3
1991	8.6	46.6	49.6	26.0	7.1	47.0	50.6	24.3	16.6	44.6	46.7	34.6	7.0	56.7	57.0	24.3
1995	7.5	45.4	47.1	24.8	6.3	45.6	47.8	23.5	14.4	43.9	44.0	32.2	5.9	54.8	58.8	23.1
1996	7.3	44.7	46.5	24.6	6.0	45.0	46.9	23.2	14.1	43.2	45.1	32.0	6.0	50.7	51.4	23.2
1997	7.2	44.2	46.6	24.5	6.0	44.8	46.8	23.2	13.6	41.5	45.2	31.4	5.7	53.2	54.3	22.8
1998	7.2	43.3	44.9	24.4	5.9	43.5	44.9	23.1	13.8	41.8	43.8	31.6	6.3	48.6	51.9	23.5
1999	7.0	43.4	45.1	24.3	5.8	43.9	45.6	23.0	13.9	40.9	42.8	31.7	5.6	52.5	53.1	22.8
2000	6.9	43.2	46.1	24.2	5.7	43.0	45.4	22.9	13.6	41.8	45.1	31.4	5.7	54.0	61.2	22.8
2001	6.8	42.2	45.8	24.1	5.7	42.3	46.0	22.9	13.2	40.5	43.7	31.0	5.5	50.7	55.1	22.6
2002	6.9	41.9	42.7	24.2	5.8	42.5	43.3	22.9	13.7	39.5	39.4	31.5	5.5	47.9	54.4	22.6
2003	6.9	42.1	43.8	24.1	5.6	43.0	44.1	22.8	13.4	40.4	42.4	31.2	7.3	41.7	46.6	24.6
2004	6.8	43.2	46.1	24.1	5.7	43.8	46.7	22.9	13.3	41.1	43.7	31.1	5.3	47.7	51.6	22.4

The results of our calculations by the “triangle-based” approximation formula in all years and for all groups are a little bit higher than the exact values calculated from the microdata. There appears to be a systematic difference averaging 2.5 days for the whole population, as well as for Whites and Blacks separately. This can be explained by departures from the conditions required for accuracy, including, for example, that the distribution of births during the year of birth is not uniform (Figure 2). The number of births during the second half of the each year is on average 6% greater than the number in the first half-year. For this reason, the number of person-years lived in the upper Lexis triangle is on average 1.4% greater than the corresponding number in the lower triangle. This increases the weight of the upper Lexis and leads to overstatement of average age of death in infancy. Note that the differences are almost the same for the whole population, both races, both sexes combined, and for male and female infants separately. However, we consider an error of 2.5 days, or less than 0.7% of a year, to be acceptable.

Figure 2. Average daily number births by month as a percent of the annual average daily number births in 1983-1991, 1995-2004 in the USA



We computed ${}_1a_0$ from the HMD input data files by the triangle-based formula and the results of each calculation are plotted as “×” markers in the Figure 3. Simultaneously, we calculated average age of infant death using the C-D formula. The thin black line corresponds to these ${}_1a_0$ estimates. The second heavy line in the Figure 3 is our alternative to C-D formula, described later.

The estimates from the C-D formula tend to be lower than the triangle-based values for 79% of male cohorts and 90% of female cohorts. If we look only at cohorts with ${}_1q_0 < 10$ per 1000, then the C-D formulas underestimate ${}_1a_0$ for 98% of our observations. In the example for the USA, ${}_1a_0$ estimated with the triangle-based formula is on average 2.5 days greater than the actual values calculated from the linked birth and death records. Perhaps analogous circumstances account for the bias in relation to estimates from the C-D formulas. The difference between ${}_1a_0$ estimated with the C-D formula and that based on the Lexis triangle formula is more than 5 days in 71% of the male and 80% of the female cohorts. If ${}_1q_0 < 10$ per 1000, then the difference between ${}_1a_0$ estimated with the C-D formula and on the basis of the Lexis triangle formula is more than 5 days in 98% of the male and 95% of the female cohorts, and this difference exceeds 10 days in 92 and 89% of male and female cohorts respectively. Detailed analysis of each of the 24 countries listed above (not presented here) has shown that in all countries secular decrease in ${}_1a_0$ has stopped or has slowed down greatly between the 1960s and 1980s. In one of the countries it was characterized by stagnation and in all the others the trend has reversed itself and a_0 has risen. Slow decline in ${}_1a_0$ synchronous with decline in the IMR is observed only in the Netherlands and Slovenia. Unfortunately data by Lexis triangles for the Netherlands are available only for the cohorts born in the years 1979-1998. The average IMR for male infants in the cohorts born in 1995-1998 is 5.9 per 1000 and

average ${}_1a_0$ is 40 days, as opposed to 24.7 days according to the C-D formulas. Major fluctuations in ${}_1a_0$ in Slovenia complicate our analysis. However, ${}_1a_0$ is on average 33.8 days in the 1999-2008 male birth cohorts and ${}_1q_0$ is 4 per 1000, which by the C-D formulas corresponds to ${}_1a_0 = 22.7$ days.

Figure 3. Relationship between average age of infant death and the infant mortality rate in the HMD database by sex.

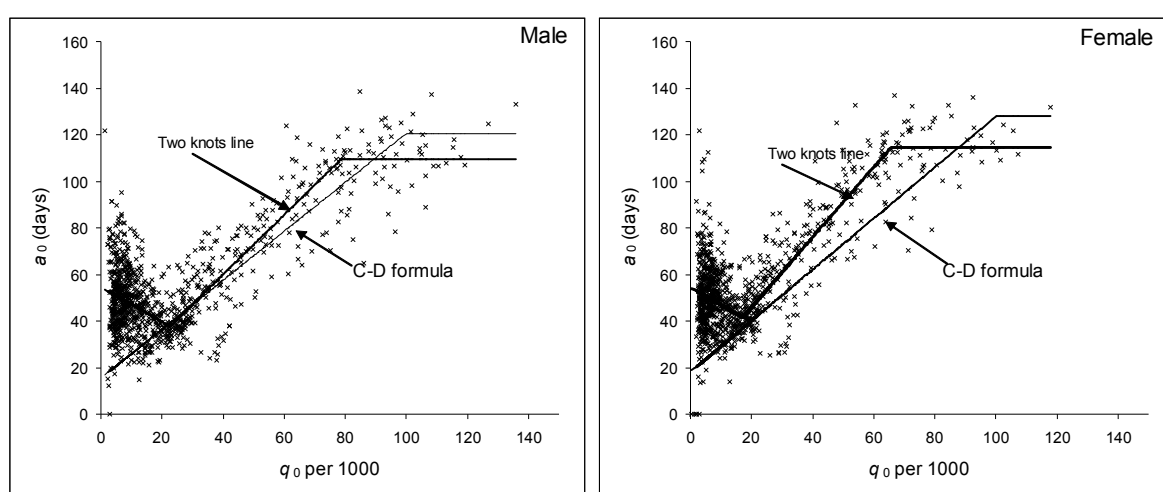
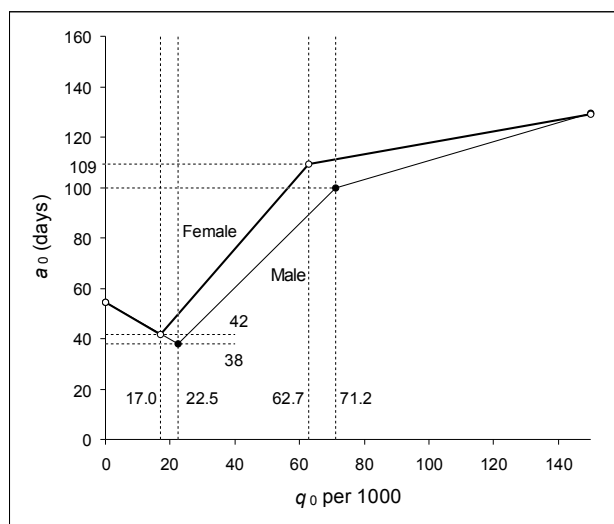


Figure 3 also shows that the relationship between ${}_1a_0$ and the infant mortality rate is complex and the data are noisy, making it difficult to find a simple functional relation of ${}_1a_0$ from the infant mortality rate by methods such as regression. If the probability of death is less than 10 per 1000 then the Pearson correlation coefficient between infant mortality rates and average ages of observed cohort deaths in infancy is less than 0.03 in magnitude. However it is possible to find some average relation between these variables that can be used where other information concerning the average age of infant death is unavailable.

Taking Coale and Demeny as a precedent, we looked for a piecewise linear function that best approximates the empirical data, but in contrast to the former authors, we sought 3-segment

piecewise linear splines on criteria of best fit. We fit the splines using the **R** routine *curfit.free.knot* in package *Dierckxspline* (Doray-Raj and Graves 2009). The package is an extension of the fitting procedures developed by Paul Dierckx and incorporated in the FITPACK package of FORTRAN subprograms (Dierckx 1987). The fundamental nonlinear least squares algorithm is described in Dierckx (1993). (Simplifying deletion) The resultant fitted linear splines for males and females are presented in Figure 4.

Figure 4. Three-segment piecewise linear spline relating ${}_1a_0$ to ${}_1q_0$ fit to the HMD database by sex.



It is important to note that although the calculations for males and females were performed independently, the abscissas of the left internal knots for males and females agree well with each other: the average female IMR in countries with male IMRs in the interval 22-23 per 1000 is 17.3. This does not hold for the right knots: the male IMR in the interval 70-73 per 1000 corresponds to female IMRs in the interval 52-61, averaging 57 per 1000. As to the rightmost segment, it is clear that ${}_1a_0$ cannot increase indefinitely with increasing IMR. We assume, as did Coale and Demeny, that at some high level of IMR, ${}_1a_0$ is constant; in other

words the third segment should be a flat line. Unfortunately, there seem to be too few observations in our dataset to estimate the appropriate third knot and the corresponding horizontal segment: our HMD-based dataset includes only 83 observations with male IMRs of more than 71 and 57 observations with female IMRs of more than 64 per 1000. There are only 21 observations of male IMRs and 8 of female IMRs at levels of more 100 per thousand. In our opinion, these data are not enough for substantively reliable definition of one more knot. Therefore, we adopted a 2-step procedure to cover the entire range of ${}_1q_0$ values. In the first step we estimated a 2-segment piecewise linear approximation for ${}_1a_0$ if ${}_1q_0 < 71$ for males and ${}_1q_0 < 63$ per 1000 for females using the *curfit.free.knot* procedure constrained to a single interior knot. In the second step we continued the 2-segment piecewise line with a horizontal segment that best approximates the leftover points (Figure 3, Table 3). We cannot display a very good fit to the empirical data. Nevertheless exactly half of the empirical observations are situated above our line.

We offer these formulas as an alternative to the Coale-Demeny equations for use in circumstances where more direct calculation of ${}_1a_0$ is not a viable option due to lack of reliable data or for other reasons. Practical recommendations for calculation of the average age at death in infancy based on the central death rate at age 0 are presented in Appendix 3. The last results we present pertain to changes in average age of infant death by cause of death. For each race we present results for the first four and last four available cohorts in the US NCHS data, namely the 1983-1987 and 2000-2004 birth cohorts (Table 4).

Across these periods the infant mortality rate declined from by 6 to 10 points per thousand. The main contributions to this decline (70-76 percent) came from two groups of causes of death: 1) certain conditions originating in the perinatal period and congenital malformations; and 2) deformations and chromosomal abnormalities.

Table 3. Parameters of 3-segment piecewise linear splines recommended for estimation of average age of death in infancy based on the infant mortality rate ${}_1q_0$.

Lower limits ${}_1q_0$	Upper limits ${}_1q_0$	Equation
Male		
0	0.0226	$0.1493 - 2.0367 \cdot {}_1q_0$
0.0226	0.0785	$0.0244 + 3.4994 \cdot {}_1q_0$
0.0785	+	0.2991
Female		
0	0.0170	$0.1490 - 2.0867 \cdot {}_1q_0$
0.0170	0.0658	$0.0438 + 4.1075 \cdot {}_1q_0$
0.0658	+	0.3141

Changes in average age were less significant, but it declined for all race categories by some 2-4 days. (Table 4).

Table 4. Infant mortality rate and average age of infant death by race of mother and major causes of death in infancy in the birth cohorts of 1983-1987 and 2000-2004, USA.

Race	Period of birth	Total	Certain conditions originating in the perinatal period	Congenital malformations, deformations and chromosomal abnormalities	Diseases of the respiratory system	Sudden infant death syndrome	All other and unknown causes
Infant mortality rate per 1000							
All Races	1983-1987	10.3	4.9	2.2	0.3	1.4	1.5
	2000-2004	6.9	3.4	1.4	0.2	0.6	1.3
White	1983-1987	8.8	3.9	2.2	0.3	1.2	1.2
	2000-2004	5.7	2.7	1.3	0.1	0.5	1.1
Black	1983-1987	18.4	10.2	2.4	0.7	2.3	2.8
	2000-2004	13.4	7.8	1.7	0.4	1.1	2.5
Other	1983-1987	10.3	4.3	2.4	0.4	1.5	1.7
	2000-2004	5.8	2.7	1.3	0.2	0.4	1.2
Average age of infant death (day)							
All Races	1983-1987	46.1	9.9	37.1	105.4	94.0	118.9
	2000-2004	42.5	8.2	37.9	120.0	95.0	106.0
White	1983-1987	46.4	10.1	36.2	104.4	94.0	119.3
	2000-2004	42.9	7.9	35.8	118.4	95.4	106.5
Black	1983-1987	44.9	9.5	41.5	106.7	94.0	118.6
	2000-2004	40.7	8.9	45.1	121.8	92.7	103.0
Other	1983-1987	50.0	10.8	36.1	109.4	93.3	114.3
	2000-2004	47.9	7.8	42.8	126.4	102.3	114.6

4. Discussion

Using perfectly exact methods for data on the USA, and for other countries an approximate formula for calculation of average age of death in infancy verified against the US data, we showed that in the process of infant mortality decline, the negative relationship between the probability of death and the average age of death disappears after infant mortality reaches some low level. How does this relate to our ideas about the long-term evolution of infant mortality?

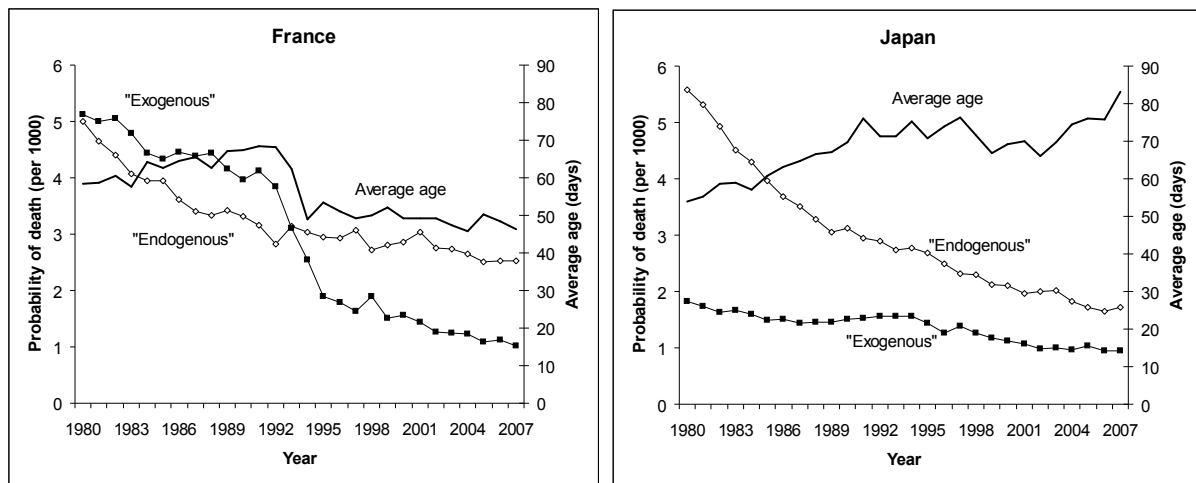
In the early 1950s, the eminent French demographer Jean Bourgeois-Pichat created the model of infant mortality that explained why infant mortality decline leads to concentration of infant deaths in the neonatal period (1951a,b). He maintained that there are two types of infant mortality: exogenous mortality due to the influence of postnatal conditions as infants become exposed to the external environment, and endogenous mortality due to conditions of the prenatal period, including congenital diseases. Endogenous mortality is concentrated in the first month of life and its level is relatively stable through time. In general, historical mortality decline has been connected with declining exogenous mortality, including in infancy. Thus, rapid infant mortality decline was observed at ages 1-11 months. Bourgeois-Pichat included under the heading of “endogenous mortality” the following 4 major causes of (infant) death: “congenital defects, prematurity, congenital anomalies, and diseases of earliest childhood” (1951b). These categories coincide with two first items on our list of causes of death in infancy. These two groups alone account for the recent infant mortality decline in low mortality countries, starting from the 1970s. It seems that at present we witness a new stage of mortality decline in the USA, which is largely the result of endogenous mortality decline in Bourgeois-Pichat’s terminology. The impact of exogenous mortality is smaller. The rapid decline of the endogenous component is what leads to the rise in the average age of infant deaths in the US.

Table 5. Infant mortality rates from endogenous and exogenous causes and average ages of infant deaths by race of mother in the birth cohorts of 1983-1987 and 2000-2004, USA.

		Infant mortality rate			Average age of infant death		
		Total	Endogenous	Exogenous	Total	Endogenous	Exogenous
All Races	1983-1987	10.3	7.1	3.2	46.1	18.3	106.7
	2000-2004	6.9	4.8	2.1	42.5	16.9	104.2
White	1983-1987	8.8	6.1	2.7	46.4	19.5	106.4
	2000-2004	5.7	4.0	1.7	42.9	17.0	103.9
Black	1983-1987	18.4	12.6	5.8	44.9	15.6	107.4
	2000-2004	13.4	9.5	4.0	40.7	15.4	102.0
Other	1983-1987	10.3	6.7	3.6	50.0	19.9	105.0
	2000-2004	5.8	4.0	1.8	47.9	19.2	113.2

Using data from the HMD combined with the WHO Mortality Database, we calculated average ages of death and probabilities of infant death from endogenous and exogenous causes for France and Japan after 1980 (Figure 5). In Japan, where the initial infant mortality rate was 7.4 per 1000 in 1980, it declined by 4.7 per thousand. About 81 percent of the decrease was due to endogenous mortality; thus, the average age of infant deaths grew by 29 days. Another situation was observed in France. The French infant mortality rate also declined between 1980 and 2007 more than twofold, from 10.1 to 3.6 per 1000 (dates ?), but the role of exogenous causes was more important and the average age of infant deaths declined by 12 days.

Figure 5. Probability on infant death from endogenous and exogenous causes and average age of infant death in France and Japan after 1980.



The infant mortality rate in Japan in 1988 was about the same as in France in 1995 but the probability of death from exogenous causes was lower than in France by 0.6 per 1000 and the average age in France was lower by 11 days.

In the US birth cohorts of 2000-2004 the proportions of total infant deaths due to endogenous and exogenous causes for the black population are about the same as for whites (Table 5) and average ages of death in infancy are quite similar, even though the infant mortality rate for Blacks is 2.4 times greater than that for whites.

Thus, in the mid-1970s, the decline of infant mortality from causes that Bourgeois-Pichat referred to as endogenous, and which in the 1960s seemed unassailable, commenced. In some populations it occurred even more rapidly than the decline of exogenous mortality. This fact explains the observed abatement and even disappearance of the connection between the level and average age of infant mortality. Starting from the 1980s, the average age of infant death became almost independent of the infant mortality rate. However, on average in countries with ${}_1q_0$ less than 0.017, further decrease of infant mortality is associated with increase in the

average age of death in infancy. Our analysis suggests that this is due to the influence of the decline in mortality from endogenous causes of death, which tends to raise the average age of death in infancy.

5. Conclusion

The Coale-Demeny model life tables and the formulas which underlie them have proven to be exceptionally resilient, remaining in use in demographic analysis for 3-4 decades. Throughout this timespan, the formulas relating ${}_1a_0$ (k_0 in their notation) to ${}_1q_0$ have remained unchanged. This is most remarkable. Ansley Coale and Paul Demeny had 326 life tables, of which 212 pertained to the period from 1945 from 1960 onwards (1966:(7)) according to Coale and Guang, (1989) data after 1960 were practically not present in their collection. In this time period the minimum infant mortality rate for both sexes was more than 12 per 1000. It would be miraculous if the empirical formulas based on this dataset remained accurate up to the present moment.

Our assembled data demonstrate that the Coale-Demeny formulas for estimation of the average age of infant death no longer hold for countries with low infant mortality by current standards. In most of these countries, starting from some moment in the process of infant mortality decline, the decrease in the average age of death in infancy has given way to increase. However, the relation between these two indicators is characterized by a reversal in the main effect as well as considerable uncertainty, making it difficult to describe with a “traditional” parametric mathematical model. Our two-knot spline is preferable for mortality modeling.

Inaccuracy in average age estimation does not influence either the magnitude of infant mortality rates or life expectancy. But this average age of death in infancy is an important

characteristic of infant mortality and is regularly used in various demographic calculations such as life table construction. We recommend calculating the infant mortality rate directly whenever possible, and estimating it by the triangle based formula when not. If estimates of ${}_1a_0$ are needed and no other data are available than infant mortality rates by sex, our formulas may be employed. Our analysis shows that these approaches are preferable to the Coale-Demeny formulas at low levels of mortality.

Acknowledgements

We are grateful to Dr. Vladimir Shkolnikov from the Max Planck Institute for Demographic Research for useful discussions on some methodological problems and preliminary results.

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Appendix 1. Proof triangle-based formula for average age of deaths.

Consider an annual birth cohort born within year $Y = [t_0, t_0 + 1)$ that satisfies the following two conditions:

- the distribution of births is uniform within Y , (the density function of the birth distribution, $\beta(t)$, is constant within Y);
- the cohort survival function $l(x, t)$, $t \in [t_0, t_0 + 1)$ does not depend on date of birth within the age interval $[x_0, x_0 + 1)$.

Then the average number of years lived within the age interval $[x_0, x_0 + 1)$ among people dying at that age $a(x_0)$ is equal to the share of number of deaths in the upper Lexis triangle in the total number of deaths at age x_0 .

Proof. Let b^Y denote the initial size of the birth cohort, and let l_x^Y stand for the cohort

survival function. It is obvious that $\int_{t_0}^{t_0+1} \beta(t) dt = b^Y$ and $l(x, t) = l^Y(x)$ for $t \in [t_0, t_0 + 1)$. It

follows that the number of cohort members attaining age x_0 by the end of calendar year t_0 is

equal to $b^Y \cdot {}_1L_{x_0}^Y$, where ${}_1L_{x_0}^Y = \int_0^1 l^Y(x_0 + \theta) d\theta$ represents the cumulated survivorship function

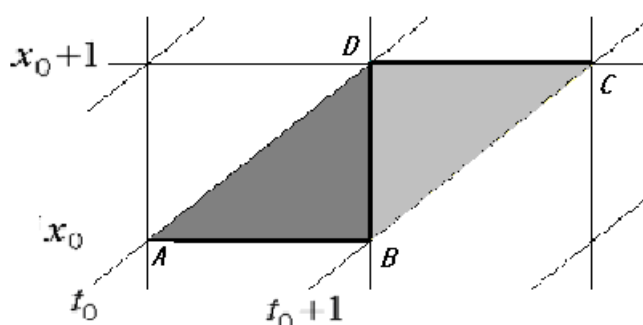
corresponding to $l^Y(x)$ in the age interval x_0 to $x_0 + 1$. In general, this number of people

is $\int_0^1 \beta(t_0 + \theta) \cdot l(x_0 - \theta + 1, t_0 + \theta) d\theta$. Taking into account the properties of the cohort assumed

above, this number of people is equal to $b^Y \int_0^1 l^Y(x_0 - \theta + 1) d\theta = b^Y \cdot \int_0^1 l^Y(x_0 + \xi) d\xi = b^Y \cdot {}_1L_{x_0}^Y$.

Let the parallelogram $ABCD$ (Figure 1-1) correspond to the cohort and age interval under consideration. We should prove that the number of deaths in the triangle BCD divided by the total number of deaths in the parallelogram $ABCD$ is ${}_1a_{x_0}^Y$.

Figure 1-1. Lexis parallelogram of deaths in a birth cohort.



The number of survivors at age x_0 is equal to $b^Y \cdot l_{x_0}^Y$ (the segment AB) and at age $x_0 + 1$ is equal to $b^Y \cdot l_{x_0+1}^Y$ (the segment DC). The segment BD corresponds to the number of survivors to the end of calendar years $Y + x_0$ and is equal to $b^Y \cdot {}_1L_{x_0}^Y$. Thus the number of deaths in the upper triangle, BCD , is $b^Y \cdot ({}_1L_{x_0}^Y - l_{x_0+1}^Y)$. If we replace ${}_1L_{x_0}^Y$ with the formula for its calculation ${}_1L_{x_0}^Y = l_{x_0+1}^Y + {}_1a_{x_0}^Y \cdot (l_{x_0}^Y - l_{x_0+1}^Y)$ we find that the number of deaths in the upper triangle is ${}_1a_{x_0}^Y \cdot b^Y \cdot (l_{x_0}^Y - l_{x_0+1}^Y)$. The product $b^Y \cdot (l_{x_0}^Y - l_{x_0+1}^Y)$ is exactly the number of deaths in the parallelogram $ABCD$. So, the ratio of BCD to $ABCD$ is $\frac{{}_1a_{x_0}^Y \cdot b^Y \cdot (l_{x_0}^Y - l_{x_0+1}^Y)}{b^Y \cdot (l_{x_0}^Y - l_{x_0+1}^Y)} = {}_1a_{x_0}^Y$

QED

Appendix 2. Possible influence of the transition to ICD 10 on indicators of infant mortality by cause of death.

Prior to the implementation of ICD 10, the US NCHS conducted a comparison study in which a sample of death records filed in 1996 was coded according to both the ICD 9 and ICD 10 classifications. From the crosstabulation of deaths by cause according to the two respective classifications, “comparability ratios”, each indicating the ratio of deaths in an aggregated cause category coded under the ICD 10 rules to the number of deaths in the same category coded under the ICD 9 rules, were calculated. (Anderson et al. 2001). These cross-classifications can also be used to develop “transition coefficients ” for converting underlying causes of death from ICD 9 categories into ICD 10 equivalents. We have opted not to employ these in our analysis, and have instead aggregated the deaths by detailed cause under the classification in effect in the respective years into a small number of broad groups of causes of death.

In our analysis we employed 4 broad categories of cause of death, plus a residual category of “all other causes”. These broad categories were not developed initially from either the ICD9 and ICD10 classifications, although both classifications were mapped at the 4-digit level into the 5 broad categories. Table 2-1 presents a crosstabulation of infant deaths coded on both the ICD9 and ICD10 classification from the NCHS comparison study conducted in 1996. The detailed causes of death for each of the two ICD versions have been grouped into the 5 broad categories used in the present analysis. Only infant deaths which were assigned valid codes on both ICD9 and ICD10 have been tabulated. The bivariate data are presented as deaths per thousand.

Table 2-1 reveals a high degree of compatibility between the ICD9 and ICD10 classifications at the very high level of aggregation represented by the 5 categories. The overwhelming majority of deaths are in cells on the main diagonal, amounting to 92 percent of all classified

infant deaths. According to the data, discrepancies due to the transition from ICD9 to ICD10 can distort the results of an intertemporal comparison of data classified under the two respective ICD versions at the level of 8 percent of all infant deaths.

Table 2-1. Correspondence between deaths classified according to IC9 and ICD10 classifications of causes of death, USA (per 1000).

	ICD10	ICD9				
		B45	B44	B31, B32	B466	All other
Certain conditions originating in the perinatal period	P00 - P96	449	21	5	0	20
Congenital malformations, deformations and chromosomal abnormalities	Q00 - Q99	4	199	1	0	7
Diseases of the respiratory system	J00 - J99	1	1	21	0	1
Sudden infant death syndrome	R95	1	1	1	102	4
All other and unknown		6	6	2	0	149

Note: Only record axis codes are considered. Causes of death for which there was no occurrence in 1996 are not represented for obvious reasons. This tabulation refers to all infant deaths which could be coded on both ICD9 and ICD10, which are less than the number of infant deaths registered in 1996.

On the basis of table 2-1 it would be possible to calculate hypothetical transition coefficients to redistribute the deaths coded under the ICD9 version and grouped into our 5 categories into the categories they would fall under if coded under ICD10. If these transition coefficients based on Table 2-1 are applied to the cohorts 1983-1987 then we can estimate possible errors

connected with changes in classification at the level of our broad groups of categories. We took the results of such a recalculation of the data for the birth cohorts 1983-1987, originally coded under ICD9, and compared them to the data for birth cohorts 2000-2004 coded under ICD10. The comparison indicates that for all races, mortality decline from certain conditions originating in the perinatal period and sudden infant death syndrome would be less than what was observed by 0.3 and 0.1 per 1000, respectively, and infant deaths in the other broad groups would be greater by 0.1-0.2 per thousand. The degree of discrepancy occasioned by the shift in ICD versions would be about the same among Whites and among the race category "Other". For the Black subpopulation the degree of discrepancy would be twice as much. For instance, mortality decline from certain conditions originating in the perinatal period would be 0.7 per thousand greater than what is indicated in Table 5. However all cause specific death probabilities for the Black subpopulation are about two times greater than the average for all races combined. This is evidence that the change in ICD versions would not alter our conclusions concerning the roles of the broad cause of death categories in the dynamics of average age of infant death. In other words, by grouping deaths into our broad categories, we subsume most of the difference in classification of causes of death within these categories.

Appendix 3. Practical recommendations for calculation of the average age at death in infancy ${}_1a_0$ based on the central death rate ${}_1M_0$ at age 0.

The equations for ${}_1a_0$ are 3-segment piecewise linear splines on ${}_1q_0$. At the each ${}_1q_0$ -segment ${}_1a_0$ is linear function of ${}_1q_0$. In terms of ${}_1M_0$ the average age is a relatively exotic function that can be derived from the two simultaneous equations

$$\begin{aligned} {}_1a_0 &= A + B_1q_0 \\ {}_1q_0 &= \frac{{}_1M_0}{1 + (1 - {}_1a_0) \cdot {}_1M_0} \end{aligned}$$

This function is the solution of the quadratic equations in ${}_1a_0$ which coefficients are linear functions of ${}_1M_0$. However the function that describes dependency ${}_1a_0$ from ${}_1q_0$ is an approximate empirical function. Therefore, excessive precision is unnecessary and it is sensible to use an approximate function for describing dependency ${}_1a_0$ from ${}_1M_0$. The same method was used by Arriaga et al. (1976) to obtain the counterpart of the Coale-Demeny formulas for ${}_1M_0$. We can presume that segments and knots of the spline function from ${}_1M_0$ correspond to segments and knots of the spline function from ${}_1q_0$. The formulas for calculation ${}_1a_0$ from ${}_1M_0$ are presented in the Table 3-1.

Table 3-1. Parameters of 3-segment piecewise linear splines recommended for estimation of average age of death in infancy based on the central death rate ${}_1M_0$.

Lower limits ${}_1M_0$	Upper limits ${}_1M_0$	Equation
Male		
0	0.02300	$0.14929 - 1.99545 \cdot {}_1M_0$
0.02300	0.08307	$0.02832 + 3.26021 \cdot {}_1M_0$
0.08307	+	0.29915
Female		
0	0.01724	$0.14903 - 2.05527 \cdot {}_1M_0$
0.01724	0.06891	$0.04667 + 3.88089 \cdot {}_1M_0$
0.06891	+	0.31411

We offer these formulas for use in circumstances where only ${}_1M_0$ is available.