

# The gestational age pattern of feto-infant mortality

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## 1. Introduction

The different segments of a birth cohort's mortality trajectory have been thoroughly mapped starting with the sudden decline in the risk of death after a peak at birth (e.g., Bourgeois-Pichat 1951; Galley and Woods 1999; Berrut et al. 2016), the arrival at minimum risk in late childhood (Ebeling 2018), the “hump-shaped” excess mortality in adolescence (e.g., Thiele 1871; Goldstein 2011; Remund et al. 2018) and the exponential increase in the mortality hazard over much of the adult life (e.g., Gompertz 1825) which eventually flattens (e.g., Perks 1932; Vaupel 1997; Horiuchi and Wilmoth 1998) and then plateaus among the oldest-old (e.g., Gampe 2010; Barbi et al. 2018). Similar investigations have been made concerning the changing mortality risk of the unborn child over the age of a pregnancy (e.g., Shapiro et al. 1962; Bakketeig et al. 1978; Goldhaber and Fireman 1991; Carlson et al. 1999; Woods 2009).

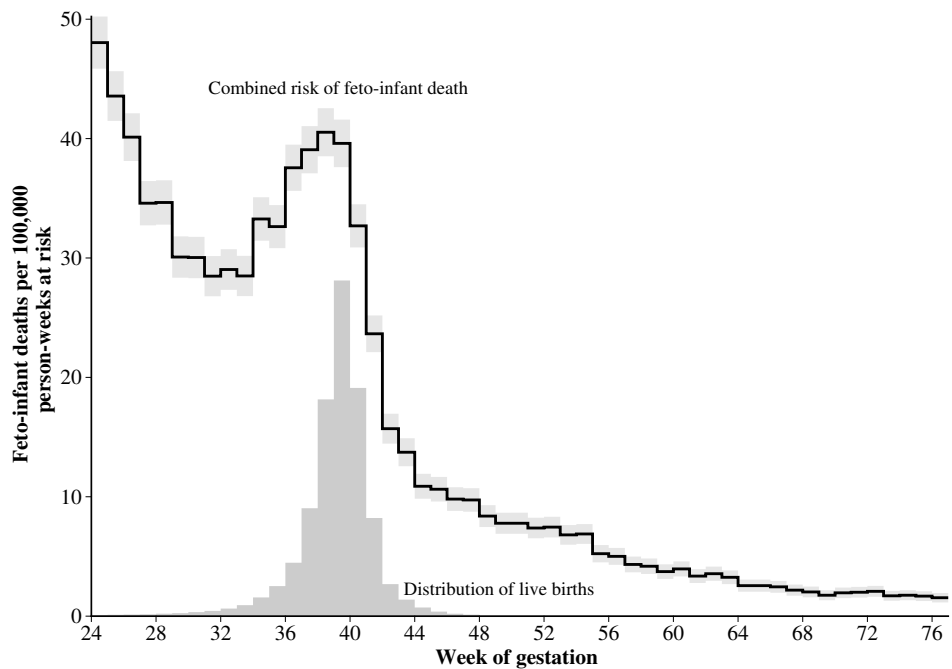
Both survival scenarios, fetal and infant, meet at the point of birth but are nonetheless fundamentally separated by the use of different timescales. While prenatal mortality is indexed by gestational age, commonly measured as the weeks since the last menstrual period of the pregnant woman, the survival of those born alive is followed over chronological age, i.e., time since birth. Such a strict separation of populations along the dividing line of birth makes this critical transition invisible in the study of mortality, delegating to it either the role of a right censoring or a point of entry into the risk set. An alternative perspective allows bridging the feto-infant gap by situating birth within the lifecycle of a cohort of unborn children whose survival is tracked over the age of gestation into infancy. By marking the vital events of fetal death, birth and infant death on a common age scale the risky transition of birth becomes an event *within* the temporal observation horizon and its effect on the survival of a cohort on the onset of life can be studied by defining a feto-infant mortality trajectory: the combined risk of fetal or infant death among all members of a conception cohort still alive at a given week of gestation.

Figure 1 shows gestation specific mortality rates for a cohort of children conceived in 2009 and either born or registered as an infant or fetal death in the U.S. The denominator of the rate is based upon all members of the conception cohort alive (either as fetus or infant) during a given week of gestation whereas the numerator includes all fetal- and infant deaths within the same week. The feto-infant mortality trajectory constructed from these rates thus measures the changing risk of *any* adverse pregnancy outcome for the 52 weeks

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**Figure 1:** The feto-infant mortality trajectory over gestational age for a U.S. cohort of fetuses conceived in 2009, surviving until fetal viability and followed over the next 52 weeks. The risk of feto-infant death among the survivors of the cohort declines exponentially over age interrupted only by a "birth hump."



following gestational age 24, commonly defined as the “limit of viability” (Seri and Evans 2008). The trajectory may be interpreted as the changing hazard of death for a cohort as its members pass the tumultuous transition period from fetus to infant. An exponential decline in mortality characterizes this period, interrupted only by hump-shaped excess mortality associated with the age-distribution of deliveries.

In this paper, descriptive findings on the feto-infant mortality trajectory and the associated phenomenon of a “birth hump” are presented.

Mortality trajectories that stretch across the feto-infant gap have been proposed several times but never entirely realized. In his seminal work on “ontogenescence,” – the declining risk of death over age before maturity – Levitis (2011) considers the hazard of death on an age continuum from conception until adolescence with the event of birth acting as a “transitional shock.” By using a negative time scale before birth and positive after, Levitis implicitly assumes all births to occur at the same time post-conception – a simplifying assumption which naturally leads to a spurious “birth spike” rather than a “birth hump.” Williamson and Woods (2003) and Woods (2009) model the cumulative risk of death of a cohort from conception until the first birthday and use weeks of gestation as time scale throughout the complete follow-up. As Williamson and Wood’s model is based on previously published disjoint fetal- and infant lifetables, they too assume all births to take place at full-term.

The joint consideration of fetal and infant death is perhaps most prominently expressed in the perinatal mortality rate, commonly defined as the sum of all fetal deaths and infant deaths during the first seven days of life over the number of births within a year (World Health Organization 2006). The necessity for such a measure arose from the uncertainty regarding the classification of a death as stillborn or infant due to both varying legal requirements and the subjective judgment of the pediatrician. Designed as a simple and therefore widely applicable indicator, the perinatal mortality rate does not consider the time dimension of pregnancy, nor does it permit a survival analytic interpretation as the denominator is not the population at risk.

Fetal lifetables add both a time dimension and a survival interpretation to the analysis of pregnancy outcomes via the introduction of an “ongoing pregnancies” denominator (e.g., Shapiro et al. 1962; French and Bierman 1962; Bakketeig et al. 1978; Goldhaber and Fireman 1991). These life-tables report the probability of fetal death among the intrauterine survivors to some week of gestation. In an influential article Yudkin et al. (1987) advertise the use of a “fetuses at risk” denominator for the analysis of perinatal mortality by gestational age leading to a range of age-specific mortality indices that include compound fetal- and infant death endpoints (Kristensen and Mac 1992; Smith 2001; Platt et al. 2004; Joseph 2004; Smith 2005; Joseph 2007). Kristensen and Mac (1992) follows a cohort of fetuses from week 31 of gestation into infancy until week 76 and calculates a corresponding survival curve. The ratio of fetal- and neonatal<sup>1</sup> deaths over fetuses at risk by week of gestation is advertised by Joseph (2007) as the proper “causal” framework to the study of perinatal mortality. Making a similar argument Platt et al. (2004) propose

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<sup>1</sup>In this paper I use the term “neonatal” as referring to the first week of life.

a Cox regression model over age of gestation featuring a combined feto-infant death endpoint.

This paper contributes to the aforementioned literature on survival analysis *around* the onset of life by studying the age pattern of mortality in a cohort of fetuses as they transition into infancy. A distinct phenomenon of this perinatal mortality trajectory is the “birth hump.” Via simple decomposition analysis, I show how fetal-, neonatal, and post-neonatal mortality and the probability of live birth all act together to form the “hump.” In a second step, I quantify the magnitude of the “hump” by proposing that the distinctive shape of feto-infant mortality on a cohort level is the result of two competing hazards: An “ontogenescent” hazard, due to causes with a declining incidence, and a “transitional” component, due to birth-related causes. I propose the probability of a fetus at 24 weeks of age to survive the following 12 months as a summary of the feto-infant mortality trajectory and ask how differences in this indicator across cohorts and between population strata are driven by changes in the shape of the gestational age pattern of feto-infant mortality.

## 2. Data and Methods

In this paper, I analyze U.S. fetal deaths, births, and infant deaths over the age of gestation. The data basis for this analysis consists of the birth certificates for the U.S. birth cohorts 1989/1990, 1999/2000, 2009/2010, the linked infant death certificates where applicable, and the fetal death certificates for the years 1989/1990, 1999/2000, and 2009/2010. Digitized versions of the certificates are provided by the National Center for Health Statistics in the form of the “Birth Cohort Linked Birth – Infant Death Data Files” (National Center for Health Statistics 2016a) and the “Fetal Death Data Files” (National Center for Health Statistics 2016b, see also Martin and Hoyert (2002) for an introduction).

To analyze the gestational age mortality trajectory of a cohort of fetuses as they transition into life, I construct three conception cohorts of all fetuses conceived during the years 1989, 1999, and 2009 respectively. The 2009 cohort is further stratified by sex and maternal origin, two characteristics which are available on most birth-, fetal- and infant death certificates and serve to show how the phenomenon of the birth hump and the ontogenescent feto-infant mortality decline compares across key demographics.

Only fetuses who survived until week 24 of gestation, commonly defined as the “age of viability,” are considered in this study. Reporting guidelines and practice for fetal death vary across states. A left-truncation age of 24 serves to rectify these differences. It is chosen because based on evidence that under-reporting drastically increases already at week 23 (Greb et al. 1987)<sup>2</sup>. Furthermore, a relatively late left-truncation age serves to minimize the bias due to the unknown numbers of induced abortions.

The initial size of the fetal-cohort at the beginning of week 24 is calculated via the “extinct cohort” method (Bakketeig et al. 1978; Feldman 1992) by adding all life-births within a

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<sup>2</sup>While their study dates back to 1987 I found evidence for the continued under-registration of fetal deaths in the U.S. prior to week 24 in the form of declining fetal death rates going from week 23 to 20 which lacks a biological explanation.

conception cohort to all fetal deaths at weeks 24+. This is due to the simple observation that a life-birth at week of gestation  $t$  was a fetus prior to  $t$ .

Using a multi-state lifetable, I follow the initial fetal population at week 24 for 52 weeks counting for each week  $t$  fetal deaths, neonatal deaths, post-neonatal deaths, and the corresponding population of survivors and their distribution across these three states. Distinguishing fetuses, newborns and infants who survived the first week of life then allows to decompose week-to-week changes in the combined fetoinfant mortality rates  $m_t = \frac{\text{\# fetal or infant deaths at week } t}{\text{Total fetoinfant time at risk during week } t}$  into changes due to a shifting distribution of fetuses, vs. neonates vs. post-neonates, and changes due to declining or increasing mortality rates within each state. Such a decomposition explains the “birth hump” in terms of the perinatal population dynamics. The Kitagawa method (Kitagawa 1955) is used to perform the decomposition.

Assuming a competing risks model (Table 1) where death is either the result from causes which exhibit gradually declining incidence over gestation (e.g., extreme prematurity, in-utero fatalities due to congenital anomalies) or from causes increasing in incidence as full-term approaches (e.g., obstetric causes), I quantify the share of fetal- or infant deaths over the one-year follow-up from fetal viability which can be attributed to the “birth-hump.”

I define  $F(52)$ , the probability for a fetus alive at the 24th week of pregnancy to die in the following year, as a summary indicator of adverse pregnancy outcomes. In order to elucidate how the “shape” of the fetoinfant hazard trajectory determines population differences in overall fetoinfant death counts, I decompose differences in  $F(52)$  between two populations into differences due to the initial magnitude of mortality at the age of fetal viability, differences due to the rate of mortality decline over gestational age, and differences due to the location, shape, and magnitude of the “birth hump” component. This decomposition is performed via the Hourouchi decomposition of the differences in  $F(52)$  as predicted by the model outlined in Table 1.

**Table 1:** Parametric specification of the feto-infant mortality trajectory over age of gestation and derived quantities.

Ontogenescent component	Transitional component
<p><i>Ontogenescent hazard</i> The instantaneous risk of fetal or infant death at gestational age <math>t = x + 24</math> due to causes with a continuously declining incidence.</p> $h^O(x) = a_1 \exp(-bx)$ <p><i>Cumulative ontogenescent hazard</i></p> $H^O(x) = \int_0^x h^O(s) ds = \frac{a_1 - a_1 \exp(-bx)}{b}$ <p><i><math>a_1</math> Level of feto-infant mortality</i> The approximate hazard of feto-infant death at age of fetal viability.</p> <p><i><math>b</math> Rate of ontogenescence</i> The relative rate of feto-infant mortality decline over gestational age in absence of birth hump.</p>	<p><i>Transitional hazard</i> The instantaneous risk of fetal or infant death at gestational age <math>x</math> due to causes associated with the timing of onset of labor.</p> $h^T(x) = a_2 \exp\left(-\frac{(x-c)^2}{2\sigma^2}\right)$ <p><i>Cumulative transitional hazard</i></p> $H^T(x) = \int_0^x h^T(s) ds = a_2 \sigma \sqrt{\pi/2} [\operatorname{erf}(A) + \operatorname{erf}(B)],$ <p>where <math>A = \frac{c}{\sqrt{2}\sigma}</math>, <math>B = \frac{x-c}{\sqrt{2}\sigma}</math>, and <math>\operatorname{erf}(\cdot)</math> is the Gaussian error function.</p> <p><i><math>a_2</math> Magnitude of birth hump</i> The instantaneous risk of fetal or infant death contributed by the birth-hump component at its peak.</p> <p><i><math>c</math> Location of birth hump</i> The gestational age <math>t = c + 24</math> coinciding with the peak of the risk of fetal or infant death contributed by the birth-hump component.</p> <p><i><math>\sigma</math> Spread of transitional shock</i> The curvature of the risk of feto-infant death around its peak. Higher values flatten the birth hump.</p>
Combined hazard	
<p><i>Hazard of feto-infant death</i> The instantaneous risk of fetal or infant death <math>x</math> weeks past fetal viability.</p> $h(x) = h^O(x) + h^T(x)$ <p><i>Feto-infant survival curve</i> The probability of surviving <math>x</math> weeks past fetal-viability.</p> $S(x) = \exp\left(-H^O(x) - H^T(x)\right)$ <p><i>Cumulative incidence of feto-infant death</i> Probability of fetal or infant death <math>x</math> weeks past fetal-viability.</p> $F(x) = 1 - S(x)$	
Competing risks inference	
<p>Cumulative incidence of feto-infant death due to causes associated with the timing of onset of labor.</p> $F^T(x) = \int_0^x S(s)h^T(s) ds$ <p>Share of feto-infant deaths over <math>x</math> weeks following fetal viability contributed by the "birth hump".</p> $\rho(x) = \frac{F^T(x)}{F(x)}$	

### 3. Results

#### 3.1 Feto-infant population dynamics over gestational age

The gestational age trajectory of feto-infant mortality as shown in Figure 2A may be segmented into a decline from week 24 to 33, a steep increase from week 33 to 39, a steep decrease from week 39 to 45, and a more gradual decrease over weeks 45 to 72. Throughout these four segments, the population composition shifts from a cohort of fetuses to a cohort with a substantial share of neonates to a cohort entirely composed of postneonates. Fetal mortality rates decline until week 32 and start to increase drastically into post-term, neonatal mortality declines until week 40, and then plateaus and post-neonatal mortality declines continuously over the entire observation period (Figure 2B). As shown by the Kitagawa decomposition in 2, the particular shape of the feto-infant mortality trajectory is the result of both the aforementioned changes in composition and rates.

*Weeks 25 to 33: pre-term decline.* Feto-infant mortality declines by 34.6 percent over the nine weeks following the age of fetal-viability. The overwhelming share of the decline can be attributed to the lessening burden of prematurity reflected in the 95.7% decline of neonatal mortality: Infants born at week 25 have a risk of death elevated by a factor of 654 compared to the fetal population at the same age whereas at week 33 this neonate penalty is reduced to a factor of 35.8. However, the increasing share of neonates from less than a percent to 1.9 percent counterbalances the effect of the reduction in neonatal mortality on the differential in feto-infant mortality.

*Weeks 33 to 39: increase towards full-term.* Approaching full-term, feto-infant mortality reverses its trend and increases by 38.9 percent. Part of this increase is due to the increase in fetal death rates by more than 144 percent, an effect that is, in turn, mediated by the declining share of the fetal population in the cohort by 51.7 percent. Neonatal mortality rates continue to be higher compared to fetal mortality prior to week 39 (Figure 2C), and thus the rapidly increasing share of newborns contributes to the increase in combined feto-infant mortality.

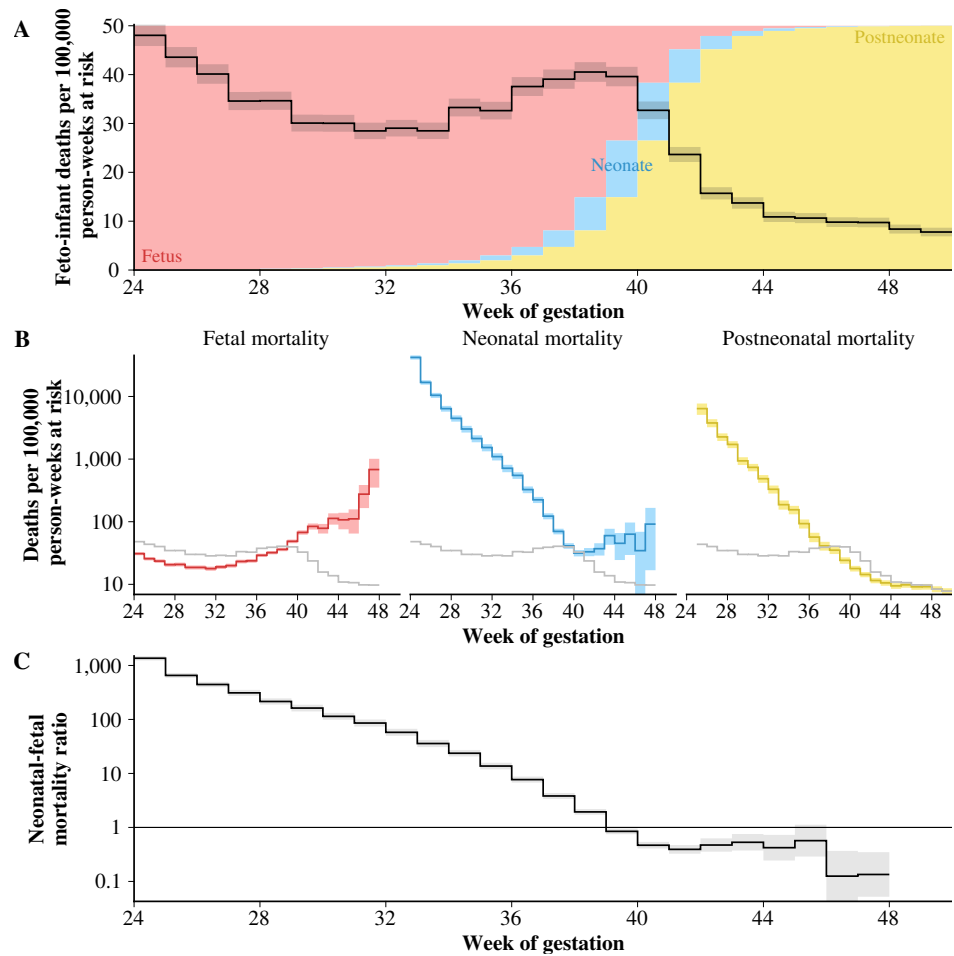
*Weeks 39 to 45: post-term decline.* While mortality increases post-term for both fetuses and neonates, it is the quickly vanishing share of both sub-populations that, along with declining rates of post-neonatal mortality, drives the steep decline in feto-infant mortality following full-term.

*Weeks 45 to 72: post-neonatal decline.* With no remaining fetuses or neonates in the population, the feto-infant mortality trajectory is completely determined by the declining mortality of the post-neonatal population.

#### 3.2 Stratum-specific feto-infant mortality trajectories

Males have a higher probability of death in the 52 weeks following fetal viability (Figure 3A). Out of 100,000 male fetuses of the 2009 U.S. conception cohort surviving until 24

**Figure 2:** Rates of fetal death, neonatal death, and post-neonatal death over weeks of gestational age as calculated for the cohort of U.S. fetuses conceived in 2009. The shaded background shows the distribution of survivors among the three states.





**Table 2:** Decomposition of the change in combined fetο-infant mortality over gestational age into contributions due to changing risk of death among fetuses, newborns, and post-neonates and the changing structure of the population along these dimensions. Percent relative change is given in parenthesis.

Week	25	→	33	→	39	→	45	→	72
<b>Mortality by stratum</b> in deaths per 100,000 person-weeks exposure									
Fetus	25.6	(-22.4)	19.9	(+144)	48.6	(+128)	111.1	.	.
Neonatal	16,774	(-95.7)	713	(-94.2)	41.2	(+53.5)	63.2	.	.
Postneonatal	6,392	(-97.1)	187	(-87.1)	24.1	(-59.3)	9.8	(-78.8)	2.1
<b>Relative exposure by stratum</b>									
Fetus	.998	(-2.6)	.973	(-51.7)	.470	(-98.9)	.005	(-100)	0
Neonatal	<.001	(+732)	.008	(+2901)	23.2	(-97.6)	.006	(-100)	0
Postneonatal	<.001	(+5,130)	.019	(+1443)	.298	(+232)	.989	(+1.1)	1
<b>Combined fetο-infant mortality</b>									
	43.6	(-34.6)	28.5	(+38.9)	39.6	(-73.2)	10.6	(-80.4)	2.1
<b>Absolute change in fetο-infant mortality</b> due to differences in									
Composition		+121.2		+96.7		-37.2		-0.39	
Rates		-136.3		-85.6		+8.2		-8.2	
Total Δ		-15.1		+11.1		-28.9		-8.6	

weeks of gestation 851–878<sup>3</sup> (one out of 114–116) will not survive the following year, compared to 739–763, (one out of 131–135) female deaths over the same period. Thus, the male probability of fetο-infant death is 12.6–17.8 percent higher than that of females. Most of this difference in survival is explained by a higher hazard level in males as measured by the  $a_1$  parameter: Ignoring the slightly earlier peak of the birth hump in males, the male hazard of fetο-infant death is consistently higher across the 52 weeks of post-viability gestational age. The hazard of fetο-infant death declines with a rate of 6.5–6.8 percent per additional week of gestation for males and 7.0–7.3 percent for females. The higher rate of ontogenescence among females substantially contributes to the sex-difference in one-year post-viability survival, while different magnitude and spread of the “birth hump” exhibit only a marginal and non-significant contribution. For additional parameter estimates see Tables 3 and 6.

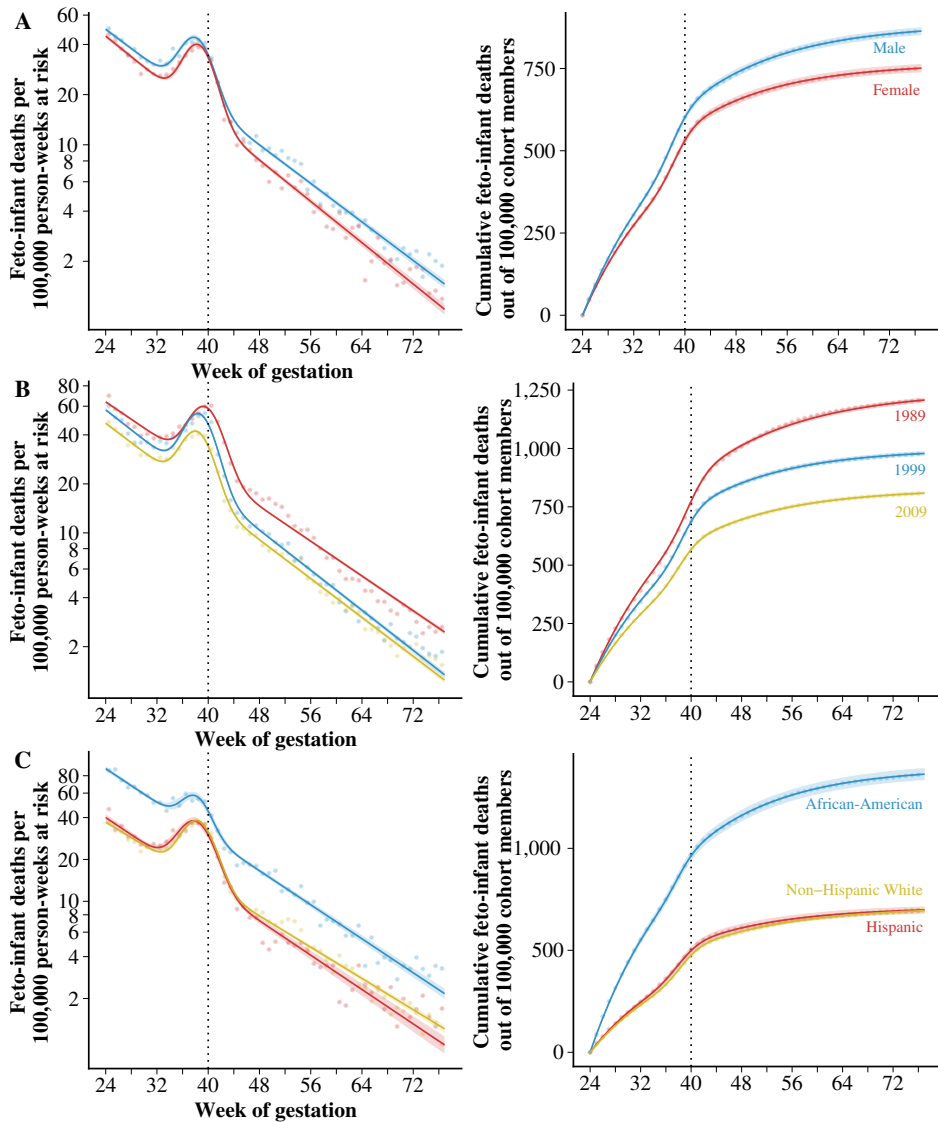
Fetο-infant survival improves considerably in the U.S. from 1989 to 2009. Out of 100,000 fetuses conceived in 1989 and reaching the age of viability 1,196–1,219 (one in 82–84), do not survive the following 52 weeks. This number drops to 969–990 (one in 101–103) deaths for the 1999 cohort and further down to 800–818 (one out of 122–125) deaths for conceptions in 2009. The 17.8–20.1 percent improvement in fetο-infant survival between 1989 and 1999 is mainly explained by an increase in the rate of ontogenescence from approximately 6.1–6.2 percent per additional week of gestation to 7.0–7.2 percent and by a drop in the level of fetο-infant mortality from 63–65 to 56–58 deaths per 100,000 person-weeks of exposure at fetal-viability. While the location of the peak “birth-hump” shifts into earlier gestation by about a week, neither magnitude nor spread of the hump change substantially between cohorts 1989 and 1999. Hence, the contribution of the transitional component to the overall improvement in one-year post-viability survival is small.

<sup>3</sup>I report the 95 percent credible interval around the estimates.

A different picture emerges for the 16.1–18.6 percent improvement in fetoinfant survival between conception cohorts 1999 and 2009, which is primarily driven by a decline in the level of fetoinfant mortality from 56–58 to 46–48 deaths per 100,000 person-weeks of exposure at fetal-viability. A substantial reduction in the magnitude of the birth hump from a peak value of 32–35 deaths per 100,000 person-weeks of exposure to 23–25 further contributes to the survival improvements while the rate of ontogenescence remains nearly constant.

There are considerable differences in fetoinfant survival by ethnicity of the mother with the hazard of fetoinfant death for the cohort of African-American origin consistently being greater than the hazards for the cohort of Hispanic or Non-Hispanic White origin (Figure 3C). In the U.S. conception cohort 2009, out of 100,000 fetuses of African-American origin at pregnancy week 24 an estimated 1,332–1,399 (one in 71–75) either die in-utero or as infants in the following year compared to 679–719 (one in 139–147), and 684–705 (one in 142–146) for fetuses of Hispanic and non-Hispanic white origin respectively. Virtually all of the differences in fetoinfant survival between the African-American stratum and the White/Hispanic origin strata are due to differences in the hazard level. Notably, the share of deaths attributable to the birth-hump is substantially lower in the African-American stratum (7.5–11.0 percent) compared to the Hispanic (18.7–24.4 percent) and non-Hispanic White (19.1–21.7 percent) strata. For further details see Tables 5 and 8.

**Figure 3:** Age trajectories of feto-infant survival A) by sex for the U.S. conception cohort 2009, B) by U.S. conception cohort, C) by maternal origin for the U.S. conception cohort 2009. Fitted (lines) versus lifetable estimates (points).



## 4. Discussion

With the advent of modern obstetric practice, the distinction between fetus and infant stage became malleable and a question of optimal choice: Should we deliver early? Should we perform surgery in-utero? Should we try to prolong the pregnancy? Better and more widely employed prenatal diagnostics made the uterus almost transparent<sup>4</sup> The fetoinfant distinction lost relevance as available knowledge about the degree of maturity, and the presence of congenital disorders informed the future survival prospects of the child, in-utero or not. Isaacson (1996), in a historical study of obstetric texts, identified the “creation of the fetus-infant” – a being separate from the pregnant woman and endowed with a history that begins before birth. The consideration of the fetoinfant mortality trajectory over age of gestation naturally arises from the continued blurring of lines that divide the stages of existence. But what does the fetoinfant as a concept contribute to the study of mortality? 1. a more realistic quantification of undesired pregnancy outcomes, 2. the phenomenon of a “birth hump,” and on a related note, 3. the possibility to model the risk associated with the transition of birth.

*Quantifying undesired pregnancy outcomes:* A sole focus on rates of infant mortality, fetal mortality or perinatal mortality hides the true incidence of adverse pregnancy outcomes, and the incompatible definition of these measures prohibits to form a simple sum which more truthfully reflects on the loss of life within a pregnancy cohort. By employing the methods of survival analysis, I have calculated that one out of 73 African-American women who conceived in 2009 and continued their pregnancy to the 24th week of gestation are expected to lose their child in the following year. Not only shows this number the continued racial disparity in the prospects of survival at the onset of life when compared to the population average of one in 124; would pregnancy come with the same warnings as prescription drugs, fetoinfant death past the age of fetal viability would have to be labeled as a “common” side effect according to the standards put forth in CIOMS Working Groups III and V (1999).

*The phenomenon of the “birth hump”:* Former bio-demographic analyses and descriptions of the age trajectories of combined fetoinfant mortality suffered either from a lack of data, the authors being forced to rely instead on rough assumptions and combinations of observations from various populations (Williamson and Woods 2003; Woods 2009; Levitis 2011; Berrut et al. 2016). The detailed micro-data on fetal deaths, births, and infant deaths in the U.S. allowed me to calculate the mortality trajectory of various fetal cohorts as they transition into infancy. Interestingly, the asymmetric sigmoid-shape for the cumulative distribution of fetoinfant deaths around term, proposed by Williamson and Woods (2003) on the grounds of theoretical considerations, is supported by the data presented here and it can be derived from the assumption of an exponentially declining hazard component added to a hazard component with a Gaussian shape. Under a simple competing risks model, less than 20 percent of fetoinfant deaths during the one-year follow-up from the age of fetal viability are contributed by the “birth hump.” Notably, among the conception cohort of African-American maternal origin, this contribution is only 9 percent, while the general level of fetoinfant mortality is much higher among this

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<sup>4</sup>In quite a real sense if one considers the images generated from a 3D ultrasound.

stratum compared to the population average. This suggests that the “hump” is an additive phenomenon, which in turn implies an increasing share of deaths during or shortly after labor on all fetal and infant death as fetoinfant mortality continues to decline.

*Risks associated with the transition of birth:* A formidable challenge for future research is to connect the aggregate phenomenon of the “birth-hump” to the transitional shock experienced by an individual as it moves from the intrauterine environment into infancy. In perinatal epidemiology, the same estimation problem is motivated by the desire to maximize the survival chances of an unborn child. Should labor be induced at a given age of gestation, or is it better to wait until labor sets in naturally? At heart lies the question of the risk of death prior to and after delivery, but such a change in risk can only ever be estimated indirectly as no one born alive ever died in utero. Current approaches to calculating the “risk of birth” thus are based on the strong assumption that children born in a given week of gestation  $t$  experienced the same risk of fetal death as the complete cohort of unborn children until  $t$ . But given that some fetal conditions are associated with a higher risk of stillbirth, pre-term birth, and infant death, this assumption must be deemed very crude, and instead, one would expect prematurely born infants to have had a higher risk of fetal death compared to infants born on full-term. Including exactly those information on fetal condition into the model, which are strongly associated with the timing of birth as well as fetal and infant death, would alleviate this issue and allow for a more precise estimate of the change in mortality across the fetoinfant transition conditional on the timing of birth. The methodological framework for such an analysis could be given by the multi-timescale and multi-state approach to survival analysis, as proposed by Iacobelli and Carstensen (2013).

## A. Description of methods

### Determining gestational age

For the purposes of this paper, a key field on the birth and death certificates is the estimated age of gestation upon delivery as a proxy for the length of a pregnancy. This measure is subject to certain biases, which are crucial to keeping in mind when interpreting the results. Gestational age is defined as the weeks since the first day of the last menstrual period of the mother (LMP) and commonly measured by subtracting the date at LMP as reported by the women from the date of (still-)birth. This method suffers from recall bias, which may lead to digit preference in the reported dates. An alternative is to derive the age of a pregnancy from ultrasound measurements of the unborn child. While the ultrasound method generally allows for a more precise estimate of the date of life-birth, it can be systematically biased in cases where the fetus is growth restricted. As abnormal fetal growth is a risk factor for fetal death, the gestational age at stillbirth may be severely biased under the ultrasound method. Additionally, the age of gestation at fetal death is positively biased by the time-lag between intrauterine death and the delivery of the dead fetus, which in-part explains the observation of fetal deaths at the implausible gestational ages of 46 and 47 weeks. In analyzing the age pattern of feto-infant mortality, I will utilize the gestational ages at delivery as they are reported on the birth and fetal death certificates and point out the role of the aforementioned biases whenever they are relevant for the interpretation of the results.

### Delineating conception cohorts

When considering the survival of fetuses into infancy, a straightforward definition of a cohort is all subjects who have been conceived during the same time period, i.e., a “conception cohort.” To determine the year of conception  $y$  for every single subject in the data I subtract the estimated weeks of gestation at delivery from the date at delivery and add two weeks to account for the average delay between the date of the last menstrual period of the mother (the time origin of the gestational age) and the date of fertilization. In this paper, I compare conception cohorts 1989, 1999, 2009.

### Assembling the multi-state feto-infant life table

After delineating the conception cohorts  $y$  and determining  $N_{24}^F$ , the number of fetuses at risk at the start of observation, I calculate a multi-state feto-infant life table across the five states of fetus  $F$ , neonate  $N$ , post-neonate  $P$ , dead  $D$  and censored  $C$ . This requires the aggregation of transition counts  $T$  among states over age. For each single week of gestation let  $T_t^{F \rightarrow D}$  denote the number of fetal deaths,  $T_t^{F \rightarrow N}$  the number of births,  $T_t^{N \rightarrow D}$  the number of neonatal deaths,  $T_t^{N \rightarrow P}$  the number of recent survivors of the first week of life,  $T_t^{P \rightarrow D}$  the number of post-neonatal deaths, and  $T_t^{P \rightarrow C}$  the number of censorings

at week 77. The fetal, neonatal and post-neonatal population at risk at the beginning of gestational age  $t$  are then given by the recurrence equations

$$\begin{aligned} N_t^F &= N_{t-1}^F - T_{t-1}^{F \rightarrow D} - T_{t-1}^{F \rightarrow N}, \\ N_t^N &= N_{t-1}^N + T_{t-1}^{F \rightarrow N} - T_{t-1}^{N \rightarrow D} - T_{t-1}^{N \rightarrow P}, \\ N_t^P &= N_{t-1}^P + T_{t-1}^{N \rightarrow P} - T_{t-1}^{P \rightarrow D} - T_{t-1}^{P \rightarrow C}. \end{aligned}$$

To calculate the population exposures, I assume a uniform distribution of births and fetal deaths within each week of gestation where  $E_t^F$  is the total time spent in the fetal state over week of gestation  $t$  by the conception cohort under observation. The exposure times for the neonate and post-neonate state,  $E_t^N$  and  $E_t^P$ , are additionally informed by the chronological age of the infant at the time of transition measured in days.

Writing  $S = \{F, N, P\}$  for the set of fetal, neonatal and post-neonatal states with  $s \in S$  I calculate, for every week  $t$ , state-specific mortality rates  $m_t^{s \rightarrow D} = \frac{T_t^{s \rightarrow D}}{E_t^s}$ , total exposure times  $E_t = \sum_S E_t^s$ , state-specific relative exposures  $p_t^s = \frac{E_t^s}{E_t}$ , the combined fetoinfant mortality rate  $\bar{m}_t = \sum_S m_t^{s \rightarrow D}$ . The empirical distribution of life-births given by  $\pi_t^{F \rightarrow N} = \frac{T_t^{F \rightarrow N}}{\sum_t T_t^{F \rightarrow N}}$ .

Expressing the combined fetoinfant mortality rate at week of gestation  $t$  as a weighted average of fetal-, neonatal-, and post-neonatal mortality rates,  $m_t = \sum_S p_t^s m_t^{s \rightarrow D}$  allows to explain the birth hump in terms of the shifting population proportions and mortality rates among the three groups over time. An application of the Kitagawa decomposition [Kitagawa1955] to the difference in fetoinfant mortality between weeks  $t_1$  and  $t_2$  yields

$$\Delta \bar{m}_{t_1, t_2} = \underbrace{\sum_S \frac{p_{t_1}^s + p_{t_2}^s}{2} \Delta m_{t_1, t_2}^s}_{\Delta r = \Delta r^F + \Delta r^N + \Delta r^P} + \underbrace{\sum_S \frac{m_{t_1}^s + m_{t_2}^s}{2} \Delta p_{t_1, t_2}^s}_{\Delta c = \Delta c^F + \Delta c^N + \Delta c^P}$$

The approach of further decomposing the rate component  $\Delta r$  and the compositional component  $\Delta c$  in their sub-population contributions  $\Delta r^F, \Delta r^N, \Delta r^P$ , and  $\Delta c^F, \Delta c^N, \Delta c^P$  has been proposed by (Chevan and Sutherland 2009).

## A latent competing risks model of fetoinfant mortality

In order to describe the shape and magnitude of the apparent “birth hump,” it is useful to separate the fetoinfant hazard trajectory into two components: a monotonically declining “ontogenescent hazard”  $h^O$  and a hump-shaped “transitional hazard”  $h^T$ . Both sources of risk add up to

$$h(x) = h^O(x) + h^T(x), \quad (1)$$

the total hazard of fetoinfant death at gestational age  $t = x + 24$ , where  $x$  measures the weeks since fetal viability. The primary purpose of model (1) is to facilitate comparisons

between populations by providing a parsimonious and informative parametrization of the fetoinfant mortality trajectory, and to that end, I choose simple parametric expressions for both components. A negative-Gompertz hazard,

$$h^O(x) = a_1 \exp(-bx),$$

captures the overall trend of log-linearly declining fetoinfant mortality over weeks of gestation with  $a_1$  being the level of the ontogenescent hazard at fetal viability, and  $b$  the rate of ontogenescence, that is the relative rate of fetoinfant mortality decline over the 52 weeks post-viability when any excess mortality contributed by the birth hump has been separated out.

The transitional hazard component reflecting the “birth hump” is specified to follow the kernel of a normal distribution resulting in the “Gaussian” hazard

$$h^T(t) = a_2 \exp\left(-\frac{(t - c)^2}{2\sigma^2}\right),$$

scaled by  $a_2$ , which measures the mortality level at the mode of the hump at age  $x = c$ . The width of the hump is controlled via parameter  $\sigma$  with larger values resulting in flatter peaks.

The above decomposition of the overall hazard of fetoinfant death into two components follows a long tradition of “competing risks” modeling of population mortality (Makeham1867, Siler1979, Heligman1980, Remund2018) where the overall risk of death is the sum of cause-specific hazards, each following a different age-trajectory. This begs the question about the nature of the risks competing with each other. The defining feature of the ontogenescent component is the continuous decline in hazard. Potential prenatal drivers of this decline are conditions that tend to lead to early fetal death such as severe congenital anomalies, e.g., anencephaly, or complications of fetal health that are all the more lethal, the earlier in pregnancy they occur, such as in-utero infection, placental dysfunction or abruption, abnormalities of the umbilical cord or rupture of the uterus. Correlated with these conditions is the risk of pre-term delivery, either induced in an attempt to save the fetus, or spontaneous. In either case, the survival chances of the pre-term child improve dramatically with the age of gestation. The post-term decline in the hazard of death may result from the continuing maturity of the cohort of infants and their increasing ability to resist the challenges posed by infection and accidents as well as the successful management of congenital malformations and chronic health conditions. Hazards due to complications of labor in late pre-term or term infants are captured by the transitional component and may be strictly birth-related such as intrapartum asphyxia or birth trauma or the consequence of severe fetal malformations that do not allow for survival outside of the uterus.

The fetoinfant mortality trajectory informs about the development of a cohort’s mortality risk, as its member’s transition into life. Complementary to this risk perspective is the incidence of fetoinfant death as measured by the probability of fetal- or infant death within  $x$  weeks following fetal-viability. The cumulative incidence of fetoinfant death can be derived from the two hazard components via the well-known relationship



$$F(x) = 1 - \exp\left(-\int_0^x h(s) ds\right) = 1 - \exp\left(-\int_0^x h^O(s) + h^T(s) ds\right).$$

### Level, ontogenescent and transition components of mortality differentials

Evaluating  $F(x)$  at  $x = 52$  gives the probability of fetal or infant death within one year of reaching the age of fetal viability, and thus  $F(52)$  is a summary measure of adverse pregnancy outcomes combining fetal and infant deaths. The difference in  $F(52)$  between two populations may be decomposed into three effects: 1) differences due to different levels of feto-infant mortality as measured by parameter  $a_1$ , 2) ontogenescent differences due to different rates of mortality decline over age of gestation as measured by parameter  $b$ , and 3) transitional differences due to the different magnitude, location and shape of the birth hump as measured by parameters  $a_2$ ,  $c$  and  $\sigma$ . Given parameter vector  $\theta = (a_1, b, a_2, c, \sigma)$ , for populations  $A$  and  $B$ , I perform a Horiuchi decomposition (Horiuchi et al. 2008) to explain how the between-population difference in each parameter contributes to the overall difference  $F(52, \theta_A) - F(52, \theta_B)$ . The level and ontogenescent contributions to the difference in one-year survival are given by the  $a_1$  and  $b$  parameter contributions respectively whereas the  $a_2$ ,  $c$ , and  $\sigma$  contributions sum up to the transitional contribution, e.g., the difference in one-year survival due to difference in the magnitude, location, and shape of the birth hump.

### Competing risks inference

How many members of a cohort fail to overcome the “birth hump” on their way to infancy? Following the calculus of competing-risks, one can derive the share of infant deaths contributed by the transitional hazard. The cumulative probability of feto-infant death due to causes associated with the transitional component is  $F^T(x) = \int_0^x S(s)h^T(s) ds$  which can be evaluated using numerical integration techniques. The share of deaths attributable to the transitional hazard up until post-viability age  $x$  then is  $\rho(x) = F^T(x)/F(x)$ .

### Censored likelihood

I fit model (1) via maximum likelihood with the likelihood function constructed from the probability of observing  $D_j$  fetal or infant deaths in age group  $j$  given model parameters  $\theta = (a_1, b, a_2, c, \sigma)$ , written as  $D_j[S(x_j|\theta) - S(x_{j+1}|\theta)]$ , and the probability of observing  $C_j$  censored survivors at the end of age group  $j$ ,  $C_j S(x_j + 1|\theta)$ , hence reflecting the fact that observations are both interval-censored, as the timing of combined fetal and infant deaths is only known to lie within some week of gestation, and right-censored, because observation stops one year after fetal-viability when most members of a cohort are still alive. Taking the product over all age groups  $j = 1 : J$  yields the likelihood function

$$L(\boldsymbol{\theta}|D_j, C_j) = \prod_j \left[ S(x_j|\boldsymbol{\theta}) - S(x_{j+1}|\boldsymbol{\theta}) \right]^{D_j} S(x_{j+1}|\boldsymbol{\theta})^{C_j},$$

with corresponding log-likelihood

$$\log L(\boldsymbol{\theta}|D_j, C_j) = \sum_j D_j \log \left[ S(x_j|\boldsymbol{\theta}) - S(x_{j+1}|\boldsymbol{\theta}) \right] + C_j \log S(x_{j+1}|\boldsymbol{\theta}).$$

## B. The nosplit algorithm for memory-efficient aggregation of event history data

The state-of-the-art for aggregation of multi-state life history data into age-period and cohort intervals is the “split-aggregate” method, whereby first, the individual level data-set is expanded into a single row per individual per time interval visited. In a second step, transition counts and state occupancy times per time interval are calculated from this expanded data set. The split step expands an already large data set even further and thus can be extremely costly in memory usage and processor time.

I present an episode-split free method to aggregate multi-state life-history data into time intervals. The “nosplit-aggregate” method first produces three summary tables from the unaltered individual-level data set and then derives the interval and state-specific risk sets, exposure times, and transition counts via elementary calculations on the aggregated tables.

The code listing below gives an implementation of nosplit in the R language (R Core Team 2020).

```
# Aggregate Transitions Counts and Occupancy Times
#
# Episode-split-free Risk-set and Exposure Time Calculation
# from Event History Data
#
# @param df
#   A data frame.
# @param t_in
#   Entry time into state.
# @param d_in
#   State being entered.
# @param t_out
#   Exit time from state.
# @param d_out
#   State being exited into.
# @param breaks
#   A numeric vector of break points for time-scale.
# @param wide
#   Output table in wide format (default=TRUE)?
# @param closed_left
#   Time intervals closed to the left and open to the right (default=TRUE)?
# @param disable_input_checks
#   Should input checks be disabled (default=FALSE)?
#
# @return
#   A data frame with columns
#   orig: origin state
#   j:   age group index
#   x:   starting age of j
#   n:   width of j
#   Z:   number of entries into origin state during j
#   W:   number of exits from origin state during j
#   P:   population number in origin state at beginning of j
#   O:   total observation time of population visiting origin state in j
#   (if wide = FALSE)
```

```

#   dest: destination state
#   W_k: number of exits from origin state to destination state during j
#   (if wide = TRUE)
#   to_*: number of exits from origin state to state * during j
AggregateStateTransitions <- function (
  df,
  t_in, d_in, t_out, d_out,
  breaks,
  wide = TRUE, drop0exp = TRUE,
  closed_left = TRUE,
  disable_input_checks = FALSE
) {

  require(tidyverse)

  t_in = enquos(t_in); d_in = enquos(d_in);
  t_out= enquos(t_out); d_out = enquos(d_out)

  # input checks

  if (identical(disable_input_checks, FALSE)) {
    # check if all transition times are contained in
    # range of breaks
    t_range = c(min(pull(df, !!t_in)), max(pull(df, !!t_out)))
    breaks_range = range(breaks)
    if ( identical(closed_left, TRUE) ) {
      if (any(
        t_range[1] < breaks_range[1] |
        t_range[2] >= breaks_range[2]
      )) {
        stop(paste0(
          'Transition time outside range of breaks. Ensure that all t_ >='),
          breaks_range[1], ' and <', breaks_range[2]
        )
      )
    }
  }
  if ( identical(closed_left, FALSE) ) {
    if (any(
      t_range[1] <= breaks_range[1] |
      t_range[2] > breaks_range[2]
    )) {
      stop(paste0(
        'Transition time outside range of breaks. Ensure that all t_ >'),
        breaks_range[1], ' and <=', breaks_range[2]
      )
    )
  }
}

# total number of age intervals
J_ = length(breaks)-1
# index of age intervals
j_ = 1:J_
# width of age intervals
n_j_ = diff(breaks)
# unique origin states

```

```

k_in_ = unique(pull(df, !!d_in))
# unique destination states
k_out_ = unique(pull(df, !!d_out))

# find the index of an interval defined by
# <breaks> each element in <x> is contained in
# returns NA if x outside breaks
FindIntervalJ <-
  function(x, breaks, cl = closed_left) {
    if (identical(cl, TRUE)) {
      # [a, b)
      right = FALSE; lowest = FALSE
    } else {
      # (a, b] with [a0, b0]
      right = TRUE; lowest = TRUE
    }
    .bincode(
      x = x, breaks = breaks,
      right = right, include.lowest = lowest
    )
  }

# 1. Aggregation

# tabulate exits by age, origin and destination state
W_k_tab <-
  df %>%
  select(t_out = !!t_out, d_in = !!d_in, d_out = !!d_out) %>%
  mutate(
    # add age interval index to each exit
    j = FindIntervalJ(pull(., t_out), breaks, closed_left),
  ) %>%
  # for each observed combination of
  # age and
  # origin state and
  # destination state...
  group_by(d_in, d_out, j) %>%
  summarise(
    # ...total number of exits
    W_k = n(),
    # total time lost in age due to exit
    Lw_k = sum(breaks[j+1]-t_out)
  ) %>%
  ungroup()

# tabulate exits by age and origin state
# based on prior tabulation on destination specific exits
W_tab <-
  W_k_tab %>%
  # for each observed combination of
  # age and
  # origin state...
  group_by(j, d_in) %>%
  summarise(
    # ...total exits
    W = sum(W_k),

```

```

    # ...total time lost in interval due to exit
    Lw = sum(Lw_k)
  ) %>%
  ungroup() %>%
  # add rows for missing combinations
  # of age interval and origin state
  complete(
    d_in = k_in_, j = j_,
    fill = list(W = 0, Lw = 0)
  )

# tabulate entries by age and state entered into
Z_tab <-
df %>%
  select(d_in = !!d_in, t_in = !!t_in) %>%
  mutate(
    j = FindIntervalJ(pull(., t_in), breaks, closed_left),
  ) %>%
  group_by(j, d_in) %>%
  summarise(
    # ...total entries
    Z = n(),
    # ...total entries right at start of interval
    Z0 = sum(t_in==breaks[j]),
    # ...total time lost in interval due to late-entry
    Lz = sum(t_in-breaks[j])
  ) %>%
  ungroup() %>%
  complete(
    d_in = k_in_, j = j_,
    fill = list(Z = 0, Z0 = 0, Lz = 0)
  )

# tabulate concurrent entries and exits by interval
ZW_tab <-
df %>%
  select(t_in = !!t_in, t_out = !!t_out, d_in = !!d_in) %>%
  # aggregate individual level entry
  # and exit times into predefined age groups
  mutate(
    # add interval index to each entry
    j = FindIntervalJ(pull(., t_in), breaks, closed_left),
    # are entries and exits in same interval?
    zw = j == FindIntervalJ(pull(., t_out), breaks)
  ) %>%
  # for each combination of
  # state and
  # interval
  group_by(d_in, j) %>%
  summarise(
    # ...total concurrent entries and exits
    # there may be NAs in logic vector <zw> when
    # and entry or exit falls outside the range
    # of all intervals. as those cases don't have to
    # be counted na.rm=TRUE is applied
    ZW = sum(zw, na.rm = TRUE)
  )

```

```

) %>%
ungroup() %>%
complete(
  d_in = k_in_, j = j_,
  fill = list(ZW = 0)
)

# 2. Determine risk-sets and exposure times

# exit counts for all possible combinations
# of origin state, destination state and
# age interval
# intrastate transitions are 0 now
# but are added later
W_k_tab_complete <-
  W_k_tab %>%
  select(-Lw_k) %>%
  complete(
    d_in = k_in_, d_out = k_out_, j = j_,
    fill = list(W_k = 0)
  )

# occurrence-exposure table
oe_tab <-
  bind_cols(W_tab, Z_tab[,-(1:2)], ZW_tab[,-(1:2)]) %>%
  mutate(
    x = breaks[j],
    n = n_j_[j]
  ) %>%
  # for each entry state...
  group_by(d_in) %>%
  mutate(
    # number of observations entering j via j-1
    #  $R_{(j+1)} = R_j + Z_j - W_j$ 
    R = c(0, head(cumsum(Z) - cumsum(W), -1)),
    # population at risk at x_j
    P = R + Z0,
    # number of observations in j that did neither start
    # nor end during j
    Q = R - W + ZW,
    # number of observations entering j
    # that do not end during j
    U = Z - ZW,
    # total observation time during j
    O = Q*n + (Z + W - ZW)*n - Lz - Lw,
    # number of intrastate transitions
    I = Q + U,
  ) %>%
  ungroup() %>%
  left_join(W_k_tab_complete, by = c('d_in', 'j')) %>%
  # intrastate transitions
  mutate(
    W_k = ifelse(d_in == d_out, I, W_k)
  ) %>%
  select(orig = d_in, dest = d_out, j, x, n, Z, W, P, O, W_k)

```

```

# drop intervals with 0 exposure
if (identical(drop0exp, TRUE)) {
  oe_tab <-
    oe_tab %>%
    filter(0 > 0)
}

# convert to wide format
if (identical(wide, TRUE)) {
  oe_tab <-
    oe_tab %>%
    mutate(dest = paste0('to_', dest)) %>%
    spread(key = dest, value = W_k)
}

return(oe_tab)
}

```



## C. Tables of parameter estimates

**Table 3:** Table of estimated parameters of feto-infant hazard trajectory over gestational age by sex.

	Male	Female
$a_1$	4.9e-4 (4.8e-4, 5.1e-4)	4.5e-4 (4.3e-4, 4.6e-4)
$b$	6.7e-2 (6.5e-2, 6.8e-2)	7.1e-2 (7.0e-2, 7.3e-2)
$a_2$	2.5e-4 (2.3e-4, 2.6e-4)	2.4e-4 (2.2e-4, 2.6e-4)
$c$	1.4e+1 (1.4e+1, 1.4e+1)	1.4e+1 (1.4e+1, 1.5e+1)
$\sigma$	2.4e+0 (2.2e+0, 2.6e+0)	2.3e+0 (2.2e+0, 2.5e+0)
$F(52) \times 10e5$	864 (851, 878)	751 (739, 763)
$F(52)^{-1}$	116 (114, 118)	133 (131, 135)
$\rho(52)\%$	17.0 (15.9, 18.2)	18.4 (16.9, 19.7)

**Table 4:** Table of estimated parameters of feto-infant hazard trajectory over gestational age by conception cohort.

	1989	1999	2009
$a_1$	6.4e-4 (6.3e-4, 6.5e-4)	5.7e-4 (5.6e-4, 5.8e-4)	4.7e-4 (4.6e-4, 4.8e-4)
$b$	6.2e-2 (6.1e-2, 6.2e-2)	7.1e-2 (7.0e-2, 7.2e-2)	6.9e-2 (6.8e-2, 7.0e-2)
$a_2$	3.5e-4 (3.4e-4, 3.6e-4)	3.4e-4 (3.2e-4, 3.5e-4)	2.4e-4 (2.3e-4, 2.5e-4)
$c$	1.5e+1 (1.5e+1, 1.6e+1)	1.5e+1 (1.4e+1, 1.5e+1)	1.4e+1 (1.4e+1, 1.4e+1)
$\sigma$	2.5e+0 (2.4e+0, 2.6e+0)	2.3e+0 (2.3e+0, 2.4e+0)	2.4e+0 (2.2e+0, 2.5e+0)
$F(52) \times 10e5$	1,207 (1,196, 1,219)	979 (969, 990)	809 (800, 818)
$F(52)^{-1}$	82.8 (82.1, 83.6)	102 (101, 103)	124 (122, 125)
$\rho(52)\%$	18.0 (17.3, 18.7)	20.1 (19.2, 21.0)	17.6 (16.7, 18.5)

**Table 5:** Table of estimated parameters of feto-infant hazard trajectory over gestational age by maternal origin.

	African-American	Hispanic	Non-Hispanic White
$a_1$	9.0e-4 (8.7e-4, 9.4e-4)	4.0e-4 (3.8e-4, 4.2e-4)	3.7e-4 (3.6e-4, 3.8e-4)
$b$	7.0e-2 (6.8e-2, 7.3e-2)	7.1e-2 (6.8e-2, 7.4e-2)	6.5e-2 (6.3e-2, 6.6e-2)
$a_2$	2.4e-4 (2.1e-4, 2.8e-4)	2.3e-4 (2.0e-4, 2.6e-4)	2.3e-4 (2.2e-4, 2.5e-4)
$c$	1.4e+1 (1.4e+1, 1.4e+1)	1.4e+1 (1.4e+1, 1.4e+1)	1.4e+1 (1.4e+1, 1.5e+1)
$\sigma$	2.1e+0 (1.7e+0, 2.5e+0)	2.6e+0 (2.4e+0, 2.9e+0)	2.4e+0 (2.3e+0, 2.6e+0)
$F(52) \times 10e5$	1,364 (1,332, 1,399)	698 (679, 719)	695 (684, 705)
$F(52)^{-1}$	73.3 (71.5, 75.1)	143 (139, 147)	144 (142, 146)
$\rho(52)\%$	9.2 (7.5, 11.0)	21.5 (18.7, 24.4)	20.4 (19.1, 21.7)

**Table 6:** Shape decomposition of differences in  $F(52)$  between the sexes.

	Male - Female
$\Delta F(52) \times 10e5$	113 [95.3, 131]
Level contribution	62.1 [33.9, 89.2]
Ontogenescent contribution	41.4 [20.9, 61.2]
Birth hump contribution	9.23 [-5.67, 23.7]
$\Delta F(52)\%$	15.0 [12.6, 17.8]

**Table 7:** Shape decomposition of differences in  $F(52)$  between the conception cohorts.

	1989 - 1999	1999 - 2009
$\Delta F(52) \times 10e5$	-228 [-244, -213]	-170 [-183, -156]
Level contribution	-97 [-121, -74.2]	-136 [-156, -116]
Ontogenescent contribution	-109 [-126, -92]	20.1 [6.0, 34.2]
Birth hump contribution	-21 [-33.8, -9.2]	-54.0 [-65.1, -42.8]
$\Delta F(52)\%$	-18.9 [-20.1, -17.8]	-17.4 [-18.6, -16.1]

**Table 8:** Shape decomposition of differences in  $F(52)$  between maternal ethnicities.

	African-American - White	Hispanic - White	Hispanic - African-American
$\Delta F(52) \times 10e5$	668 (637, 701)	3.3 (-18.6, 25.4)	-665 (-701, -630)
Level contribution	747 (704, 797)	41.7 (10.3, 75.8)	-680 (-738, -623)
Ontogenescent contribution	-63.5 (-92.9, -35.0)	-46.9 (-72.3, -20.5)	-8.8 (-50.6, 32.7)
Birth hump contribution	-15.2 (-40.5, 11.9)	8.5 (-12.7, 32.8)	23.7 (-8.2, 56.7)
$\Delta F(52)\%$	96.4 (91.2, 102)	0.04 (-2.7, 3.7)	-48.8 (-50.7, -47.0)

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