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Clustering of infant deaths among Nigerian women: investigation of temporal patterns using dynamic random effects model

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Abstract

This study was conducted to estimate the magnitude of infant death clustering as well as the mortality risk associated with death of a preceding child and investigate how these have changed over three decades (1980–2013) in Nigeria. Birth history data from the Nigeria Demographic and Health Survey for 1990, 2003, 2008 and 2013 were analysed using dynamic random effects models. The effect of death of an immediate preceding child (sibling mortality correlation) was estimated by controlling for background characteristics and unobserved heterogeneity.

A total of 232,090 single births to 56,123 women were analysed. Results showed that 13.2% in the oldest maternal cohort (≤ 1969) experienced death of at least two infants and they accounted for 40.2% of all infant deaths. Among the 1970–1979 maternal cohort, it was 8.6% and 31.6% respectively. In the youngest maternal cohort (≥ 1980), 3.3% had recorded multiple infant deaths but accounted for 20.3%. Model results revealed that sibling mortality correlation increased the probability of infant death by 0.080 and 0.061 in the 1980–1989 and 2010 birth cohorts respectively.

There is a substantial level of infant death clustering in Nigeria, and this is closely driven by sibling mortality correlation both of which have declined very slowly over time. To achieve desired progress in child survival, death clustering should be addressed alongside other barriers to child survival.

Keywords: Infant mortality, Death clustering, Death concentration, Preceding child, Sibling mortality correlation, Dynamic models, Nigeria

Introduction

As a reflection of the global progress, the annual rate of under-five mortality reduction in sub-Saharan Africa (SSA) peaked at 4.1% during 2000–2015 compared to 1.6% in 1990–2000 (UNICEF, 2015). If current trends persist globally, 47 countries may not meet the Sustainable Development Goal (SDG) target of 25 deaths per 1000 live births by year 2030 (UNICEF, 2015). Thirty-four of these 47 countries are in SSA. Therefore, the rate of under-five mortality reduction must be accelerated to meet the SDG targets. This is very important especially in those countries with high fertility and population growth rates. One of the potential threats to the global efforts to accelerate childhood mortality reduction is a phenomenon called “death clustering”. In simple terms, childhood death clustering refers to the concentration of child deaths among certain women (Sastri, 1997b). Although it has been of research interest for some time, it was

first introduced into the demographic literature in the early 1990s (Das Gupta, 1990). Since then, it has been investigated using varieties of methodological approaches contingent on the type and source of data.

Childhood deaths could be clustered in certain women because children born to the same woman may share genetic, household and community characteristics. The factors found to influence childhood death clustering in mothers/families are also related to the established bio-demographic and socio-economic determinants of childhood mortality which include short birth spacing, poverty and maternal education among others (Omariba, Beaujot, & Rajulton, 2007; Sastry, 1997a). At the community, levels are socio-cultural norms that affect child healthcare and environmental characteristics such as the source of drinking water, sanitation and access to healthcare (Sastry, 1997a; Zenger, 1993). Several authors most of whom analysed data from health and demographic surveillance systems have also reported child death clustering from clusters of communities/villages (Adjuik, Kanyomse, Kodayire, Wak, & Hodgson, 2010; Alabi, Baloye, Doctor, & Oyedokun, 2016; Awini, Mattah, Sankoh, & Gyapong, 2010; Lutambi, Alexander, Charles, Mahutanga, & Nathan, 2010; Van Bodegom, Eriksson, Houwing-Duistermaat, & Westendorp, 2012). Few of these also identified poor infra-structural and environmental characteristics as the main predictors of spatial clustering of child mortality (Alabi *et al.*, 2016; Awini *et al.*, 2010).

With advancement in statistical methodologies which has continued to enhance empirical investigation and better understanding about childhood death clustering, explanations of the processes culminating in the phenomenon have evolved over the past two decades (Edvinsson & Janssens, 2012). One of the mechanisms that have been proposed to underlie death clustering is the higher risk of child mortality associated with death of a preceding child. Previously, this has been lumped together with the effect of unobserved variables (unobserved heterogeneity). It is now empirically feasible to investigate the mortality risk associated with death of the preceding child (or sibling mortality correlation) while controlling for unobserved heterogeneity. This requires the use of dynamic/transition models which take advantage of the sequence of births or deaths among children born to a woman. To date, the technique has been applied to single cross-sectional data from three developing countries—India (Arulampalam & Bhalotra, 2006, 2008), Kenya (Omariba, Rajulton, & Beaujot, 2008) and Bangladesh (Saha & van Soest, 2011). Infant death clustering was found to be high in 13 out of the 15 states in India (Arulampalam & Bhalotra, 2008). Sibling mortality correlation raised the risk of infant death by 29% and 40% in Bangladesh and Kenya respectively. Infant death clustering tends to be common in high fertility settings (Sastry, 1997a). Therefore, it is worthwhile to replicate studies on death clustering in different countries especially where fertility and childhood mortality remained high.

Furthermore, there is a suspicion that childhood mortality reduction could be stalled where death clustering is prevalent (Arulampalam & Bhalotra, 2008). Although it is expected that childhood death clustering should decline with mortality levels, however, there is no conclusive evidence on the temporal pattern of infant death clustering and sibling mortality correlation in sub-Saharan Africa where childhood mortality rates are among the highest globally. Empirical evidence on the temporal trends in infant death clustering is very important for policies and programmes on child survival. The present study seeks to address this gap and provide further insights into

the temporal structure of sibling mortality correlation as a potential explanation for infant death clustering in Nigeria.

Nigeria is one of the most populous countries in sub-Saharan Africa with very slow progress in child survival (Akinyemi, Adebawale, Bamgboye, & Ayeni, 2015; UNICEF, 2015). Fertility has remained stagnated at above 5 children per woman since 1980s. This is not surprising because contraceptive prevalence has remained very low (Alkema, Kantorova, Menozzi, & Biddlecom, 2013). There is a wide differential in under-five mortality rates across the six geo-political regions, though partly attributed to socio-economic factors (Adedini, Odimegwu, Imasiku, Ononokpono, & Ibisomi, 2014b; Akinyemi, Bamgboye, & Ayeni, 2013). Several intervention programmes have been implemented to address issues related to child survival. These programmes cover topical issues such as child nutrition, immunisation and management of childhood illnesses. Though some progress has been made in the last one decade, mortality and other child health indicators are still below desired levels (Winter, Akinlo, & Florey, 2016).

Previous studies have identified individual-, household- and community-level determinants of childhood mortality (Adedini, Odimegwu, Imasiku, & Ononokpono, 2014a; Akinyemi et al., 2015; Antai & Moradi, 2010), but death clustering has not been previously explored. Reports from an operational research study conducted in three Northern states of Nigeria (Jigawa, Yobe and Zamfara) revealed that more than 80% of child deaths were reported by only 20% of the women studied (Klouda & Adamu, 2013). Similarly, analysis of the 2011–2013 data from Nuhche Health and Demographic Surveillance System (HDSS) in Zamfara State, North West Nigeria, revealed that about one third of under-five deaths were contributed by only 5% of all households in the HDSS (Alabi, Oyedokun, Doctor, & Adedini, 2017). In addition, the authors also confirmed the effects of many determinants of under-five mortality. These two Nigeria studies on childhood death clustering did not estimate the contribution of death of a preceding sibling to infant mortality neither was the data robust enough to investigate temporal changes in infant death clustering and sibling mortality correlation.

Given the fact that infant death clustering is common in high fertility settings (Arulampalam & Bhalotra, 2008), Nigeria is a suitable setting to further explore the phenomenon. Therefore, this study is aimed at addressing the following objectives: (i) to estimate the magnitude of infant death clustering among women and effect of death of a preceding child (sibling mortality correlation) in Nigeria and (ii) to investigate how these phenomena have changed over three decades. The main hypothesis investigated was that sibling mortality correlation is a significant contributor to infant death in Nigeria.

Literature overview

A survey of the literature revealed that most of the explanations for death clustering are closely connected to the proximate and distal determinants of childhood mortality.

These factors which are associated with concentration or clustering of child deaths could be observed or unobserved (unmeasured during data collection) depending on the study design and data collection instrument. Besides, they could operate at individual (child or mother), family (household) and community levels (Adedini et al., 2014b; Antai & Antai, 2008). Based on these premises, previous studies have estimated child death clustering using different approaches depending on the available data. The most

common type of data employed has been based on birth histories of women aged 15–49 years which are often collected during Demographic and Health Surveys (DHS) in many developing countries.

One analytical approach is that in which childhood death clustering is estimated as the excess observed deaths over the expected (Das Gupta, 1990; Ronsmans, 1995). Expected deaths are modelled using binomial or negative binomial models based on an underlying assumption that the risk of child death is the same for all women (or families). Unfortunately, this assumption is sometimes not true. Another weakness of the approach is the inability to investigate how the mortality of children in the same family affects inter-family variations (Zaba & David, 1996).

The second analytical technique which has so far been the most popular was developed around the concept of “unobserved heterogeneity” and has been used to infer death clustering. The idea emanates from the notion that death clustering implied wide variations in the distribution of child deaths between families. Death clustering was therefore estimated as the “unobserved heterogeneity” after all measurable/observed predictors of child death have been controlled. For example, see Madise, Mathews, and Whitworth (2014); Omariba et al. (2007); and Sastry (1997b). One limitation of the approach is that it mixed up death clustering in families/women with “unobserved heterogeneity” which could be at the level of the mother, household or community. Besides, correlated mortality risks among infants born to the same mother are not specifically controlled. Death clustering and unobserved heterogeneity are separate empirical concepts as demonstrated in some later studies (Arulampalam & Bhalotra, 2008; Saha & van Soest, 2011).

Two other measures proposed for child death clustering were proportion of children dead out of all births to a woman and a binary measure for multiple deaths (Kuate-Defo & Diallo, 2002). These have been subjected to linear and logistic regression respectively in order to identify the determinants of death clustering (Kuate-Defo & Diallo, 2002). Though simple and logical, the measures did not permit the investigation of shared dependence in the risk of death among children born to the same woman.

Other methodological studies argued that child death clustering is largely driven by the correlation in the risk of death among siblings. The basic argument was that the survival or otherwise of an infant depends on that of his/her immediate predecessor. Alternatively, the death of a previous child is believed to reduce the survival chances of an index child. Although a few previous studies have investigated the effect of the survival of the previous child on infant mortality (Ikamari, 2000; Zenger, 1993), they were however not within the context of clustering; neither did they quantify the extent of sibling mortality correlation net of unobserved heterogeneity. In this study, we attempt to bridge this gap by investigating the infant mortality risk associated with death of a preceding child (sibling mortality correlation) while controlling for unobserved heterogeneity using data from four rounds of demographic and health survey in Nigeria.

Methods

Data sources

Data for this study were extracted from the birth history records of the 1990, 2003, 2008 and 2013 Nigeria Demographic and Health Survey (NDHS). The NDHS is a

cross-sectional nationally representative household survey first conducted in 1990 while the most recent was in 2013. Selection of eligible women for interview involved a two-stage stratified cluster sampling technique. Detailed descriptions of the survey methodologies are available in the respective reports (National Population Commission (NPC) [Nigeria], 2014; National Population Commission [Nigeria], 2004).

Out of the five rounds conducted so far (1990, 1999, 2003, 2008 and 2013), the greatest decline in childhood mortality was observed between 2003 and 2013 (Akinyemi et al., 2015). Evidence actually shows that several other countries witnessed a faster decline in childhood mortality since 2000s (UNICEF, 2015). To undertake a robust exploration of infant death clustering among different birth cohorts of Nigerian children, birth history data from the four surveys (1990, 2003, 2008 and 2013) were merged together and analysed.

Study sample

Analysis of the birth history data was restricted to birth cohorts grouped as follows: 1980–1989, 1990–1999, 2000–2009 and 2010 onward. This classification was based on prior evidence on childhood mortality patterns in Nigeria which revealed that the levels were higher in the 1980s and 1990s but gradually declined since early 2000s (Akinyemi et al., 2015). Multiple births were excluded from the analysis because of their higher risk of death in early childhood (Uthman, Uthman, & Yahaya, 2008). The final weighted sample size was 232,090 births to 56,123 women.

Variables

The outcome variable analysed in this study was the risk of infant death which refers to death of a child within the first 11 months after birth. We focused on infancy because more than half of deaths among under-five children occur in the first few months after birth (Rajaratnam et al., 2010). In the DHS, birth history data were collected on every child a woman has given birth to. Variables collected include the date of birth, sex, survival status, preceding birth interval and age at death if the child is not alive at the time of the survey.

The primary explanatory variable which was also used to capture sibling mortality correlation was the survival status of the immediate preceding child. Other determinants of infant mortality which are either specific to each child or known to be constant once a woman starts childbearing were included as control variables. The following child characteristics were analysed: sex, birth interval and birth order. Birth interval is a major biological determinant of infant mortality (Rutstein, 2008b) and is important in this study because the death of a preceding child could result in a shorter birth interval for the next. In such circumstance, birth interval may partly explain sibling mortality correlation. Birth order is also important in a high-fertility setting like Nigeria in view of the established knowledge that the risk of infant mortality is higher among multiparous women (Akinyemi et al., 2013).

Maternal factors included in the analysis were the age of the mother at child's birth, maternal education, household wealth index, type of residence, religion and geo-political region. Maternal education is a critical factor in child survival because it is an indicator of the socio-economic status of the mother as well as her knowledge and

skill in child care (Smith-Greenaway, 2013). Also, previous studies have shown that child mortality rates in Nigeria are higher in rural areas and in the North West and North East regions (Adedini et al., 2014b; Antai, 2011). We also controlled for religious affiliation (Christianity, Islam or traditional) of the mother because it has been shown that certain childcare behaviour and related practices are influenced by religious beliefs (Antai & Antai, 2008). The household wealth index was derived from ownership of selected items using principal component analysis (Rutstein & Staveteig, 2014). It was categorised as poorest, poorer, middle, richer and richest. Wealth index has been used as an indicator of socio-economic status in previous studies on maternal and child health (Rutstein, 2008a). Other variables such as maternal and child healthcare utilisation were excluded because they were collected only for the index child in the surveys. The effects of these unobserved variables were appropriately captured in the random component of the fitted models.

Description of the statistical model

Following methodological developments for infant death clustering and unobserved heterogeneity (Arulampalam & Bhalotra, 2008; Saha & van Soest, 2011), we employed a dynamic random effects model to investigate the effect of death of the immediate preceding child (sibling mortality correlation) on infant mortality. Detailed explanations of the advanced statistical models can be found in the above-cited references. However, the basic theoretical underpinnings are briefly described. Issues of empirical and statistical interest are the need to account for sibling mortality correlation, observed maternal/children variables and unobserved heterogeneity due to characteristics shared by children from the same mother/family, household and community. The model equation was as follows:

$$y_{i,j}^* = x_{ij}'\beta + \gamma y_{ij-1} + \alpha_i + u_{ij} \quad (1)$$

where $y_{i,j}^*$ is the probability of death for infant j from mother i , x represents observed child-level and maternal covariates and β is the coefficient (effect) of the covariates. γ captured the sibling mortality correlation which is the effect of death of the previous child, y_{j-1} . α_i is a random parameter representing the unobserved heterogeneity (at family, household and community levels) while the last component, u_{ij} , is the error term which assumed a normal distribution with a mean of zero and variance, σ_u^2 . Hypothesis was tested for $\alpha = 0$ which implied that unobserved heterogeneity has no effect on the probability of infant death. Similarly, hypothesis $\gamma = 0$ means that there was no sibling mortality correlation or that death of the preceding child does not affect the risk of infant death in the index child.

The expression in Eq. (1) is a form of a Markov chain model because of the assumption that the death of a child depends on that of its predecessor (Omariba et al., 2008). In this study, a first-order Markov chain model was implemented which means that the death of a child is only dependent on that of its immediate predecessor.

Inclusion of the term y_{ij-1} in the model imposed a constraint because the term (y_{ij-1}) was non-existent for firstborn children since they have no predecessors. In the econometrics literature where the model originated, this is known as the “initial condition problem”. Further explanations on this and different approaches to deal with it are provided in

Arulampalam et al. (2006). During modelling, this constraint was handled by specifying a separate equation for first births. The equation is as follows:

$$y_{i1}^* = z' \lambda + \theta \alpha_i + u_{i1} \quad i = 1, \dots, n \quad (2)$$

where z is a factor of covariates related to the survival of first births with λ representing their coefficient. The effect of unmeasured variables (unobserved heterogeneity) was represented by $\theta \alpha_i$ in Eq. (2) above. Equations (1) and (2) therefore represent a complete model for infant mortality with due control for sibling mortality correlation (death of preceding child), observed maternal and child variables and unobserved heterogeneity.

The final joint dynamic random effects model was fitted using a Stata programme called *redprob* (Stewart, 2006). The programme produced estimates for sibling mortality correlation, effect of covariates (x), variance due to unobserved heterogeneity and their standard errors. It was implemented in Stata MP version 14.

Analysis

The first stage of data analysis involved the use of descriptive statistical tools to summarise the explanatory variables according to child birth cohorts. This afforded the opportunity to assess variations in the maternal and other background characteristics of children in the four birth cohorts. Furthermore, to provide a clear indication of child death clustering among women, we described the distribution of infant deaths among the 56,123 mothers according to maternal cohort.

The probability of infant death was estimated using life table techniques as applied in previous studies (Akinyemi, Adedini, Wandera, & Odimegwu, 2016). Similar to previous studies (Arulampalam & Bhalotra, 2008; Saha & van Soest, 2011), the probability of infant death was estimated conditional on death of the immediate preceding child (p_1) and conditional on the survival of the immediate preceding child (p_0). The two probabilities were subsequently used to derive the average partial effect (APE) of sibling mortality concentration. APE was estimated as $p_1 - p_0$, and this gives an indication of the excess probability of infant death attributable to death of the preceding child. This measure was first estimated based on raw data and therefore not adjusted for background variables, correlated mortality risk and unobserved heterogeneity. Later in the analysis, after the dynamic random effects models were fitted, the same measures were re-estimated and adjusted by using parameters obtained from the models. Separate models were fitted for the four child birth cohorts to describe changes in sibling mortality correlation over time. We estimated the percentage of infant death explained by sibling mortality correlation. This was calculated as the ratio of the adjusted APE to the unadjusted APE.

To further explore how sibling mortality correlation affects infant mortality, four multivariable models were fitted to the pooled birth history data in a sequential manner. Model I included only the survival status of the preceding child. For model II, we included child birth cohort as dummy variables along with an interaction term for the survival status of the preceding child and child birth cohort. Model I assessed the overall influence of sibling mortality correlation on infant death, while the interaction term in model II was used to investigate the temporal trends in the effect of death of the preceding child (sibling mortality correlation). Coefficients for dummy-coded child

birth cohort are indicative of trend in infant mortality risk (which is not the focus in this paper), while the interaction term was used to assess changes in sibling mortality correlation over time. A statistically significant interaction term implies that sibling mortality correlation has changed.

In model III, all other background characteristics except birth interval were added to model II to investigate the extent to which these factors explain the effect of sibling mortality correlation on infant mortality. Lastly, birth interval was added in model IV to explore its role as potential pathway through which sibling mortality correlation affects infant death.

Ethical approval

Data analysed in this study were retrieved from the DHS online archive after due approval. Besides, NDHS 2008 and 2013 were approved by the Nigeria National Health Research Ethics Committee. DHS questionnaires and protocols were reviewed and approved by the ICF Institutional Review Board (The DHS Program, 2017) to ensure compliance with the U.S. Department of Health and Human Services regulations for the protection of human subjects (45 CFR 46).

Results

Background characteristics

Background characteristics of births to women are presented in Table 1 according to birth cohorts. The percentage of male births ranged from 52.3% in the 1980–1989 birth cohorts to 50.4% among the cohorts born in 2010 onward. The proportion of first births decreased gradually from 32.8% in the 1980–1989 birth cohort to 19.9% in the most recent cohort, but a reverse pattern was observed for births of order 6 and above. With respect to birth interval, the percentage of children born within 24 months after a preceding birth declined from 27.1% (1980–1989 birth cohort) to 17.1% (≥ 2010 birth cohort) while the percentage born after at least 36 months had increased from 11.7 to 29.9%.

Furthermore, there has been a gradual change in the age distribution of childbearing. Among the 1980–1989 birth cohorts, 37.9% of their mothers were aged below 20 years compared to 12.6% in the 2010+ cohort. Notable shift was also observed in maternal education such that the percentage of births to women with secondary/higher education rose from 15.3 to 32.5%. Rural-urban distribution remained almost the same with about two thirds of children in every birth cohort being born to women in rural areas.

Infant death clustering

Descriptive findings on infant death clustering among the 56,123 mothers are presented in Table 2 according to maternal birth cohorts. The percentage of women with no experience of infant death rose from 67.1% among the oldest cohort (born ≤ 1969) to 83.4% in the youngest cohort of mothers (born ≥ 1980). As an indication of infant death concentration among women, 13.2% of women in the oldest cohort (≤ 1969) have experienced death of at least two infants and these accounted for 40.2% of all infant deaths. Among the 1970–1979 maternal cohort, 8.6% had experienced multiple infant deaths and accounted for 31.6% of women whose infants had died. In the youngest maternal

Table 1 Background characteristics of births, Nigeria, 1980–2013

Variables	Birth cohorts			
	1980–1989 (n = 30,330)	1990–1999 (n = 74,211)	2000–2009 (n = 106,478)	≥ 2010 (n = 21,071)
Child-related variables				
Male sex	52.3	51.3	51.0	50.4
Birth order				
1	32.8	23.7	20.3	19.9
2–3	41.7	36.0	33.6	32.4
4–5	18.2	22.2	23.2	23.1
6+	7.4	18.1	22.9	24.7
Birth interval				
First born	32.8	23.7	20.3	19.9
< 24	27.1	26.1	23.8	17.1
24–36	28.5	30.5	31.9	33.2
> 36	11.7	19.6	24.0	29.9
Maternal variables				
Age at child's birth				
< 20	37.9	21.4	16.0	12.6
20–34	61.9	73.5	69.5	70.3
≥ 35	0.2	5.1	14.5	17.1
Education				
None	59.9	54.5	51.4	48.9
Primary	24.8	24.7	22.5	18.6
Secondary/higher	15.3	20.8	26.0	32.5
Household wealth quintile				
Poorest	24.4	25.0	24.9	23.4
Poorer	22.4	22.8	23.2	23.3
Middle	21.0	20.3	19.1	18.9
Richer	18.4	17.6	17.3	17.7
Richest	13.8	14.4	15.5	16.7
Religion				
Christianity	39.3	38.2	37.3	36.4
Islam	46.3	51.0	58.1	62.1
Others	14.4	10.8	4.6	1.5
Type of place of residence				
Rural	31.1	30.8	31.4	34.6
Urban	68.8	69.2	68.6	65.4
Region				
North Central	16.6	17.5	17.5	17.3
North East	34.5	35.1	35.9	37.5
North West	12.0	13.2	13.6	13.7
South East	10.4	9.4	8.6	8.9
South South	14.4	11.7	10.7	9.4
South West	12.0	13.1	13.8	13.2

Table 2 Infant death clustering among three maternal cohorts in Nigeria, 1980–2013

No. of infant deaths	Maternal cohorts					
	≤ 1969		1970–1979		≥ 1980	
	All women (%)	Women with infant deaths (%)	All women (%)	Women with infant deaths (%)	All women (%)	Women with infant deaths (%)
0	67.1		72.9		83.4	
1	19.7	59.8	18.5	68.4	13.3	79.7
2	8.1	24.5	5.7	21.1	2.7	16.2
3	3.1	9.5	1.8	6.7	0.5	3.0
4	1.4	4.3	0.7	2.4	0.1	0.8
5+	0.6	1.9	0.4	1.4	0.0	0.3
Number	12,529	4124	18,495	5008	25,099	4172

cohort (≥ 1980), 3.3% had recorded multiple infant deaths but accounted for 20.3% of all women who had lost infants to death. Overall, there was a notable decline in infant death concentration from the oldest to youngest maternal cohorts.

Figure 1 shows the distribution of infant death clustering across the six geo-political regions in Nigeria. Though death clustering was most prevalent in North East, North West and North Central, it was common in all the other three regions (South East, South South and South West). In addition, the graph also showed that the level of infant death clustering declined over time in all regions.

Table 3 provides further measures on the contribution of death clustering to infant mortality. To profile the temporal pattern from 1980s till 2010s, results were presented according to child birth cohort. The average partial effect (APE) which represents the excess probability of infant death given that the immediate preceding child also died in infancy declined from 0.17 in 1980–1989 to 0.08 in 2010 onward.

Parameter estimates from the full dynamic random effects model were also used to generate adjusted APE (Table 4). The results show that after adjusting for selected observed variables and unobserved heterogeneity, sibling mortality correlation increased the probability of infant death by 0.080 and 0.061 in the 1980–1989 and 2010 birth

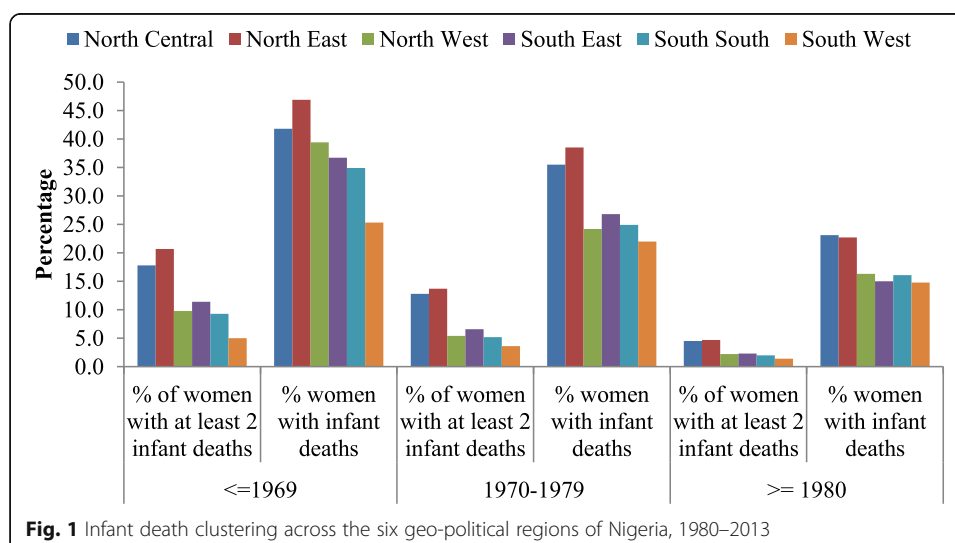


Table 3 Unadjusted average partial effect of sibling mortality correlation on probability of infant death in Nigeria, 1980–2013

Row no.	Probabilities	Computation	1980–1989 (<i>n</i> = 30,330)	1990–1999 (<i>n</i> = 74,211)	2000–2009 (<i>n</i> = 106,478)	≥ 2010 (<i>n</i> = 21,071)
1	Probability of infant death conditional on death of the preceding sibling (p_1)	Obtained from the life table technique using the Kaplan-Meier method	0.2435	0.247	0.1982	0.1262
2	Probability of infant death conditional on survival of the preceding sibling (p_0)	Obtained from the life table technique using the Kaplan-Meier method	0.0763	0.0752	0.0643	0.0499
3	Average partial effect ($p_1 - p_0$)	Row 1 – row 2	0.1672	0.1718	0.1339	0.0763

cohorts respectively. In other words, with death of the preceding child, the infant mortality level was higher by 61 deaths per 1000 live births among children in the 2010 birth cohorts. The percentage of infant death explained by sibling mortality correlation ranged from 48.1% in 1980–1989 birth cohort to 79.9% in the 2010–2013 birth cohort.

Detailed results from models based on full birth history for 1980–2013 are presented in Table 5. The coefficient for death of the preceding child was positive and statistically significant in all four models which showed that the risk of infant death was significantly higher when the immediate preceding child also died during infancy (effect of sibling mortality correlation). In model II, the coefficient of interaction term for death of the preceding child and child birth cohort was negative and statistically significant. This is an indication that the effect of death of the preceding child (sibling mortality correlation) had declined over time.

Changes in the coefficient for death of the preceding child between model II and model III showed that the adjusted variables partly explained the effect of sibling mortality correlation which remained statistically significant. The percentage change in the coefficient of death of the preceding child between model II and model IV was 10%. Both birth interval and death of the preceding child remained statistically significant as determinants of infant death. The effects of other variables such as male sex, birth order, maternal education and age at child's birth maintained their expected direction

Table 4 Adjusted contribution of death clustering to infant mortality in Nigeria, 1980–2013

Row no.	Probabilities	Computation	1980–1989 (<i>n</i> = 30,330)	1990–1999 (<i>n</i> = 74,211)	2000–2009 (<i>n</i> = 106,478)	≥ 2010 (<i>n</i> = 21,071)
1	Predicted probability of infant death conditional on death of the preceding sibling excluding the first births (p_1)	Conditional on the death of the preceding sibling; this was obtained from the probabilities predicted using model parameters	0.1658	0.166	0.1406	0.0929
2	Predicted probability of infant death conditional on survival of the preceding sibling excluding the first births (p_0)	Conditional on the survival of the preceding sibling; this was obtained from the probabilities predicted using model parameters	0.0854	0.0849	0.0664	0.0319
3	Adjusted APE [excluding first births] ($p_1 - p_0$)	Row 2 – row 1	0.0804	0.0811	0.0742	0.061
4	% infant death explained by clustering	(Row 3/row 3 Table 3) × 100	48.1	47.2	55.4	79.9

Table 5 Results from dynamic random effects models for infant mortality, Nigeria, 1980–2013

	Coefficients (standard error)			
	Model I	Model II	Model III	Model IV
Child-related variables				
Death of preceding child, γ (SE)	0.4859 (0.0159)*	0.7069 (0.0423)*	0.6773 (0.0423)*	0.6318 (0.0426)*
Child birth cohort				
1980–1989 (ref)				
1990–1999		0.0214 (0.0166)	0.00913 (0.0183)	0.0513 (0.0185)*
2000–2009		– 0.0664 (0.0170)*	– 0.0903 (0.0233)*	– 0.135 (0.0236)
≥ 2010		– 0.2342 (0.0239)*	– 0.2538 (0.0319)*	– 0.1235 (0.0324)*
Interaction term				
Death of preceding child $\times$ child birth cohort		– 0.0932 (0.0163)*	– 0.0886 (0.0163)*	– 0.0999 (0.0164)*
Male sex				
			0.0832 (0.0094)*	0.0832 (0.0095)*
Birth order				
1 (ref)				
2–3			– 0.1846 (0.0146)*	– 0.1329 (0.0148)*
4–5			– 0.0979 (0.0134)*	– 0.0729 (0.0136)*
6+				
Birth interval				
< 24				0.4681 (0.0137)*
24–36				0.2434 (0.0133)*
> 36 (ref)				
Maternal variables				
Age at child's birth				
< 20 (ref)				
20–34			– 0.1836 (0.0169)*	– 0.1267 (0.0170)*
≥ 35			– 0.1620 (0.0258)*	– 0.0582 (0.0261)*
Education				
None (ref)				
Primary			– 0.0323 (0.0149)*	– 0.0299 (0.0149)*
Secondary/higher			– 0.0516 (0.0189)*	– 0.0526 (0.0191)*
Household wealth quintile				
Poorest (ref)				
Poorer			– 0.0021 (0.0138)*	– 0.0043 (0.0139)*
Middle			– 0.1226 (0.0162)*	– 0.1201 (0.0164)*
Richer			– 0.1459	– 0.1420

Table 5 Results from dynamic random effects models for infant mortality, Nigeria, 1980–2013
(Continued)

	Coefficients (standard error)			
	Model I	Model II	Model III	Model IV
			(0.0194)*	(0.0195)*
Richest			– 0.2122 (0.0253)*	– 0.2129 (0.0255)*
Religion				
Christianity			– 0.0589 (0.0219)*	– 0.0657 (0.0221)*
Islam			– 0.0728 (0.0192)*	– 0.0866 (0.0194)*
Others (ref)				
Type of place of residence				
Rural			0.0339 (0.0141)*	0.0329 (0.0142)*
Urban (ref)				
Region				
North Central			– 0.0358 (0.0226)	– 0.0529 (0.0229)*
North East			0.0644 (0.0266)*	0.0269 (0.0229)
North West			0.0792 (0.0225)*	0.0456 (0.0228)*
South East			0.1067 (0.0247)*	0.0573 (0.0249)*
South South			0.0297 (0.0240)*	0.0052 (0.0242)
South West (ref)				
Other model parameters				
θ (SE)	0.8939 (0.0708)*	0.8691 (0.0698)*	0.9169 (0.0807)*	0.9133 (0.0807)*
Family-level heterogeneity (SE)	0.1246 (0.0065)*	0.1257 (0.0067)*	0.1116 (0.0068)*	0.1126 (0.0069)*
Log-likelihood	– 64,816.322	– 64,666.778	– 64,145.23	– 63,508.255

* $p < 0.05$

which is well known in the literature. The results for unobserved heterogeneity also retained their statistical significance in all the models.

Discussion

Infant death clustering or concentration of infant death in certain women has the potential to slow down the desired progress in child survival. Although not a new phenomenon, methodological advancement has made it possible to clearly demonstrate that infant death clustering is different from unobserved heterogeneity. Apart from monitoring the reduction of childhood mortality levels, it is also necessary to pay attention to death clustering. Using birth history data from four DHSs in Nigeria, this study was conducted to contribute evidence in this regard.

Infant death clustering was found to be prevalent in Nigeria. The degree of child death clustering found in this study is higher than observed in other countries such as Kenya (Omariba et al., 2008), Bangladesh (Saha & van Soest, 2011) and India

(Arulampalam & Bhalotra, 2008). This is not surprising given the fact that both fertility and child mortality levels in Nigeria are higher than these countries. Since there is evidence that childhood mortality levels are higher in countries with higher fertility (Silva, 2012), the same pattern may have been mirrored in the level of infant death clustering. Furthermore, results for different maternal cohorts suggest that the general level of infant death clustering in Nigeria has declined over time. This pattern is consistent with prior evidence across several countries in SSA which showed that childhood death clustering reduced with mortality levels (Kuate-Defo & Diallo, 2002). Further exploration of infant death clustering patterns revealed that though the magnitudes differ, the phenomenon is present across the six geo-political regions. This agrees with a previous study which showed that determinants of childhood mortality are virtually the same across the different regions in Nigeria (Akinyemi et al., 2015).

The results confirmed that death clustering (as indicated by sibling mortality correlation) is a major contributor to overall levels of infant mortality. In fact, the percentage of infant death explained by clustering in Nigeria was found greater than that obtained in Kenya (40%) and India (15%). Besides, in conformity with the existing literature (Arulampalam & Bhalotra, 2008; Saha & van Soest, 2011), the effect of sibling mortality correlation remained statistically significant even after observed and unobserved variables (heterogeneity) have been controlled. Although birth interval partly explained the effect of sibling mortality correlation, however, both factors remained statistically significant in the full model. Death of an immediate preceding child could shorten the length of postpartum amenorrhea due to cessation of breastfeeding, thereby making a woman become pregnant within a shorter interval. This explanation was termed “fecundity hypothesis” by Arulampalam et al. (Arulampalam & Bhalotra, 2006). Death of the preceding child could also trigger what is known as the replacement effect (Rutstein & Winter, 2014) which implied that a woman intentionally makes effort to replace a dead child notwithstanding the prevailing circumstances that led to his/her death. Exploration of the replacement effect will require detailed analysis of contraceptive use history in the inter-birth periods. Unfortunately, such data are not available in the Nigeria DHS.

Although sibling mortality correlation was found to have decreased over time, its contribution to infant death rose from 55.4% in the 2000–2009 child cohort to 79.9% in the most recent cohort (2010). This shows a distinction between the magnitude and impact of death clustering on infant mortality levels. A substantial impact of sibling mortality correlation as found in this study is a re-awakening on the urgent need to prevent infant death to forestall the risk it constitutes for the survival of succeeding births. It also underscores how infant death clustering could be triggered by the death of a child. In addition, increased impact of sibling mortality correlation may also be related to the poor state of maternal and child healthcare utilisation in the country (Winter et al., 2016). Death clustering was almost non-existent in a rural area of Bangladesh where health service coverage was optimal (Saha & van Soest, 2011). Investigation of the maternal and child healthcare continuum in Nigeria revealed that the level of drop-out at different stages is very high (Akinyemi, Afolabi, & Awolude, 2016). This can no doubt contribute to the persistence of infant death clustering.

A significant net effect of sibling mortality correlation after adjustment for birth interval, background socio-demographic characteristics and unobserved heterogeneity

implied that apart from the fecundity hypothesis, other processes or factors may be involved. Child healthcare practices, environmental and heredity cannot be ruled out. Recent empirical evidence on the causes of childhood deaths in Nigeria showed that sepsis, birth injury and pneumonia are prevalent (Adewemimo et al., 2017). Incidentally, data from the NDHS also revealed that coverage of medical treatment for fever and symptoms of acute respiratory tract infections remained stagnated at low levels between 2003 and 2013 (Winter et al., 2016). Without sustained improvement in maternal and child healthcare utilisation, infant death clustering might remain high and eventually slow down the progress desired for child survival.

An obvious limitation in this study stems from the fact that there was no data to permit exploration of other possible explanations for sibling mortality correlation. Such unobserved factors include maternal knowledge and competence in child care, cultural and environmental factors and other biological/genetic factors. Although our analytical method aptly captured these as unobserved heterogeneity, proper identification of the effects of these factors in child health and survival would need further research. Also, some variables such as maternal and childcare utilisation were not controlled in the analysis. These variables were not captured in the cross-sectional DHS birth histories. Although analysis could have been restricted to 5-year birth history so that some of these variables can be included in the models, this would have made it difficult to properly explore the temporal pattern in death clustering because the data would have covered only 5 years.

Another source of limitation is the non-availability of time-varying covariates in the DHS data. For example, mothers might have migrated from rural to urban areas during the course of child bearing. Change in place of residence poses different risks to child survival. Also, children born to a woman might have different fathers as union instability and remarriage is common (Isiugo-Abanihe, 1998). Many of these time-varying characteristics could not be adjusted. Rather, they were captured as unobserved heterogeneity for the purpose of statistical efficiency.

Conclusion

There is a substantial level of infant death clustering in Nigeria. Analysis of birth history data within a space of about three decades (1980–2013) revealed that there has been a little decline over time. Mortality risks shared by infants born to the same mother are a major contributor to death clustering and overall levels of infant mortality. These results have important implications for research, policies and programmes. Beyond the known determinants of childhood mortality, further research needs to unravel the unknown factors driving death concentration in certain women. By implication, it has become imperative to understand the steps or actions taken by parents and especially women to prevent repeated child deaths. To unravel these, longitudinal studies using a mix of data collection methodologies would be necessary.

A slow decline of infant death clustering throws up a new challenge to child survival programmes. If death clustering should remain stagnated, it means that inequity in child health and survival will persist and a stage would be reached that further decline in childhood mortality becomes difficult. This definitely negates the current sustainable development goal whose overarching intention is to “live no

one behind” (WHO, 2015). It is therefore important to continue to prevent child deaths and if a woman ever experiences a child loss, repeat occurrence should be prevented. Advocacy programmes could sensitise the population on the need for extra efforts to ensure that women with experience of infant death be supported to avoid a recurrence. Adequate coverage and uptake of maternal and child healthcare services is a potential strategy in this regard.

Abbreviations

APE: Average Partial Effect; HDSS: Health and Demographic Surveillance System; NDHS: Nigeria Demographic and Health Survey; NPC: National Population Commission, Nigeria

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Availability of data and materials

The dataset(s) supporting the conclusions of this article is(are) available in the online DHS archive [<https://dhsprogram.com/data/available-datasets.cfm>]. Permission from the DHS program is required to access the data.

Authors' contributions

JOA contributed to the conceptualisation and study design, analysis and interpretation and drafting of the manuscript. COO, OOB and BMG contributed to the interpretation of results and revision of intellectual content. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

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