RESEARCH ARTICLE

Does recall time matter in verbal autopsies? Evidence from urban informal settlements in Nairobi, Kenya [version 1; peer review: 2 approved with reservations]

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Abstract

Background: To assign a cause of death to non-medically certified deaths, verbal autopsies (VAs) are widely used to determine the cause of death. The time difference between the death and the VA interview, also referred to as recall time, varies depending on social and operational factors surrounding the death. We investigated the effect of recall time on the assignment of causes of death by VA.

Methods: This is a secondary analysis of 2002-2015 survey data of the Nairobi Urban Health Demographic Surveillance System (NUHDSS). The independent variable recall time was derived from the date of death and the date when the VA was conducted. Univariate and multivariate logistic regression methods were used to calculate odds ratios of assigning a cause of death in defined categories of recall time.

Results: There were 6218 deaths followed up between 2002 and 2016, out of which 5495 (88.3%) had VAs done. Recall time varied from 1-3001 days (mean 115.5 days, sd216.8). Majority of the VAs (45.7%) were conducted between 1-3 months after death. The effect of recall time varied for different diseases. Compared to VAs conducted between 1-3 months, there was a 24% higher likelihood of identifying HIV/AIDS as the cause of death for VAs conducted 4-6 months after death (AOR 1.24; 95% CI 1.01-1.54; p-value = 0.043) and a 40% increased chance of identifying other infectious diseases as the cause of death for VAs conducted <1 month after death (AOR 1.4; 95% CI 1.02-1.92, p-value = 0.024).

Conclusions: Recall time affected the assignment of VA cause of death for HIV/AIDS, other infectious diseases and maternal/neonatal causes. Our analysis indicates that in the urban informal setting, VAs should be conducted from one month up to 6 months after the death to improve the probability of accurately assigning the cause of death.

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Any reports and responses or comments on the article can be found at the end of the article.

Keywords

Nairobi, Verbal autopsy, Cause of death, Recall, time

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Introduction

Mortality data collected as part of vital registries, disease surveillance systems and epidemiological studies is essential for decision making¹. The need for its accuracy and reliability cannot be overemphasized^{2,3}. Currently, the gold standard method for assigning the specific cause of death depends on the certification by a medical practitioner using guidelines stipulated in the International classification of diseases and related health problems (ICD), currently available in the eleventh version⁴.

In countries where the vital registration systems are not fully developed, or in situations where there are challenges in medical certification of deaths, such as a home death, verbal autopsies (VAs) are an important means of assigning the cause of death^{5,6}. The VA consists of interviews using standardized questionnaires with the deceased person's close relatives or caregivers who are aware of the circumstances leading to the death⁶. The most widely used VA questionnaires are by the World Health Organization⁶ and the Health Metrics Research Consortium⁷.

After completion of the VA interview, data from the VA questionnaire is then interpreted to assign the cause of death. Methods for assigning the cause of death from the VA interview data vary and include the use of either physician certified reviews or computer coded verbal autopsy systems such as inter-VA, random forest (RF), tariff, King-Lu (KL) or simplified symptom pattern (SSP) among others⁸. A systematic review comparing physician certified VAs with various computer coded VA systems found that although the methods differed in the cause of death output, none of the VA interpretation methods reviewed was superior to the others⁸.

A key element in the reliability of cause of death data as determined through VAs is the time between when the verbal autopsy was conducted and when the death occurred, also referred to as the recall time⁹. There are varying optimal recall time to achieve maximum validity of a VA available in literature ranging from as soon as the death occurs up to 12 months and beyond^{10–12}. Data from a study in South Asia indicates that the probability of assigning a correct cause of death by VA methods decreases by 0.55% per increasing month of the recall time⁹. In contrast, a study in South Africa found that apart from neonatal causes, there was no impact on the validity of VA for recall time as long as one year¹³. It is of importance that the timing of the VA not only takes account of the validity but also social and cultural factors¹⁴. As such, it may be imperative to set context-specific optimal timing for VAs. In this paper, we aim to determine the optimal timing of conducting a VA to achieve higher odds of assigning the accurate cause of death in a low-income setting. We use mortality data collected from the Nairobi Urban Health Demographic Surveillance System (NUHDSS). This NUHDSS is run by the African population and research Centre (APHRC) in the informal settlements of Korogocho and Viwandani in Nairobi, Kenya¹⁵.

Methods

Study setting

This paper utilizes mortality data collected from two informal settlements in Nairobi Kenya, that form the NUHDSS, a DSS system run by APHRC. Korogocho is located in Ruaraka Sub-County in Nairobi and covers an area of 0.9 square km with a total population of 36,900 and a density of 42,401 persons per km² as per the 2019 Kenya population and housing census¹⁶. Viwandani is located in Makadara Sub-County, it covers an area of 5 km² with a population of 43,070 and a density of 8554 persons per square km¹⁶. Viwandani forms part of Nairobi's industrial region and its inhabitants are mainly casual labourers within the industries in the area, while Korogocho is inhabited by residents mainly engaged within the informal job sector.

Key challenges in Korogocho and Viwandani, as in many other urban informal settlements, include minimal formal infrastructure such as roads and piped water networks as well as rising cases of communicable and non-communicable diseases, rampant insecurity, environmental pollution, alcohol and drug abuse¹⁵.

Design

This secondary analysis uses VA data collected through a series of surveys by the APHRC in the NUHDSS. The general methodology of the NUHDSS is summarized elsewhere¹⁷. In summary, these were a series of surveys conducted from 2002 to 2015 using methodologies described by The International Network for the Demographic Evaluation of Populations and their Health (INDEPTH) network¹⁸. Surveys were done every four months. Information on the events surrounding the death within the demographic health system was collected by field interviewers and later independently validated by three physicians to assign a probable cause of death. Data from the questionnaire was then used as inputs in the [InterVA-4 software](#), a tool that uses probabilistic models based on Bayes' theorem to interpret symptom data and determine possible causes of death¹⁹. The output from the InterVA-4 model consists of up to three likelihoods per case attributed to different causes. An indeterminate residual portion is assigned when the sum of the probability of the three likelihoods is less than 100%¹⁹. For this study, we utilized the output from the InterVA-4 model to carry out analysis of optimal recall time.

Data analysis

We used STATA version 15.1 in the analysis and included all records where a successful VA was conducted. The main independent variable of interest in the analysis, recall time, is derived from the date of death and the date when the VA was conducted. Based on consideration of the average mourning time, and the acceptable time to conduct a VA as per the Kenya Verbal Autopsy Guidelines²⁰, we categorized recall time into less than 1 month, 1 to 3 months, 4 to 6 months, 7 to 12 months and greater than 12 months. In the descriptive analysis, we tabulate the recall time against other background

characteristics to calculate their frequencies in the categories of recall time.

For each death, there were three possible causes as an output from the InterVA-4 model. In all our analysis, we used the most probable cause of death output. The dependent variables were derived from each specific probable cause of death, transformed into a binary variable of yes/no. There were 22 most probable causes of death as an output in the interVA-4 model. To reduce data sparsity in our modelling, we collapsed these 22 probable causes of death into 12 categories guided by their counts and “reasonable relatedness”. Frequently occurring diseases such as tuberculosis and HIV/AIDS were retained in their original categories. Details of this re-categorization of the probable causes of death are found in [Table 1](#). This approach to recategorization has also been utilised elsewhere^{13,17}. We determined the cause-specific mortality fractions using the most probable cause of death from the collapsed categories.

Table 1. Cause of death recategorization criteria.

Cause of death as per interVA-4	Re-classified cause of death groups
AIDS/HIV	AIDS/HIV
Asthma	Other chronic/NCDs
Anaemia	Malaria/Anaemia
Cardiovascular	Cardiovascular
COPD	Other chronic/NCDs
Diabetes mellitus	Other chronic/NCDs
Diarrhoeal diseases	Other infectious diseases
Direct obstetric	Neonatal/Maternal
Infectious diseases	Other infectious diseases
Injuries	Injuries/Other external
Liver disease	Other chronic/ NCDs
Malaria	Malaria/Anemia
Malnutrition	Other chronic/NCDs
Meningitis	Meningitis
Neonatal causes	Neonatal/Maternal
Malignancies	Malignancies
Respiratory tract infections	Respiratory tract infections
Tuberculosis	Tuberculosis
Other communicable disease	Other infectious diseases
Other non-communicable disease	Other chronic/NCDs
Other external cause	Injuries/Other external
Indeterminate	Indeterminate

To investigate the effect of recall time on determining the cause of death outcome of the VA, logistic regression methods were used to calculate odds ratios of assigning a probable cause of death using the categories of recall time defined above. The 1 to 3 months recall time was used as the reference category for the logistic regression. In the univariate logistic regression analysis, association of recall time and probable cause of death outcome of the VA was tested using chi square tests while in the multivariate analysis, likelihood ratio tests were carried out. Wald tests were used to determine the effect of various categories of recall time. The backward modelling approach was used in the multivariate analysis to adjust for the background characteristics.

Ethical considerations

Permission to use the dataset was obtained from APHRC. The original NUHDSS study obtained ethical clearance from Kenya Medical Research Institute (KEMRI) Ethical Review Committee (Ref number KEMRI/RES/7/3/1)

Results

Distribution of deaths in the NUHDSS 2002–2016

There were a total of 6218 deaths followed up between 2002 and 2016, of which the highest number were in 2011(533; 8.6%), out of which 5495 (88.3%) had VAs conducted. Most of the deceased were between the ages of 15–49 (3147, 50.6%). Distribution of deaths by year and age is shown in [Figure 1](#). The most common cause of death in the period of 2002 to 2016 was tuberculosis (16.93% n=930) and the least commonly identified cause of death was malaria and or anaemia. The cause-specific percentages are shown in [Figure 2](#).

Description of VA recall time

The range of recall time for deaths with VAs done was from 1 day to 3001 days, with a mean recall time of 155.5 days (standard deviation 216.8). Majority of the verbal autopsies (45.7%) were conducted between one to 3 months of death. Distribution of socio-demographic characteristics varied based on the various verbal autopsy recall periods as shown in [Table 2](#). There were more VAs conducted within the first month of death between 2010 and 2013 (26%) as compared to the other years under study.

The effect of recall time on VA cause of death assignment

The effect of recall time varied for different diseases ([Table 3](#)). We found strong evidence (p-value = 0.0124) that identifying HIV/AIDS as the likely cause of death varied based on the timing of the VA. Conducting the VA 4 to 6 months after the death increased the likelihood of identifying HIV/AIDS as the cause of death by 24% compared to conducting it between 1 to 3 months (AOR 1.24; 95% CI 1.01-1.54; p-value = 0.043).

Both the crude and the adjusted analysis showed that an assignment of malaria/anaemia, meningitis, malignancies and other chronic diseases as the probable cause of death did not depend on the timelines within which the VA was conducted. We observed that assignment of tuberculosis as the probable

Figure 1. Number of deaths over years by age categories in the NUHDSS.

Figure 2. Cause-specific mortality fractions in the NUHDSS between 2002-2016 derived from the interVA-4 model output (collapsed categories).

Table 2. Distribution of background characteristics by recall time.

Recall time	< 1 month	1–3 Months	4–6 Months	7–12 Months	>12 Months	Total
	N= 5,494(100%)					
	936 (17.04)	2,509 (45.67)	1,012 (18.42)	579 (10.54)	458 (8.34)	5,494 (100%)
Variable						
Gender						
Female	371 (15.85)	1,071 (45.77)	457 (19.53)	247 (10.56)	194 (8.29)	2340 (100)
Male	565 (17.91)	1438 (45.59)	555 (17.60)	332 (10.53)	264 (8.37)	3154 (100)
Age						
<1 year	167 (14.09)	551 (46.50)	223 (18.82)	144 (12.15)	100 (8.44)	1185 (100)
1–14years	108 (15.84)	294 (43.11)	126 (18.48)	89 (13.05)	65 (9.53)	682 (100)
15–49 years	509 (18.36)	1234 (44.52)	508 (18.33)	268 (9.67)	253 (9.13)	2772 (100)
50–64 years	95 (18.16)	264 (50.48)	93 (17.78)	47 (8.99)	24 (4.59)	523 (100)
65 plus	57 (17.17)	166 (50.00)	62 (18.67)	31 (9.34)	16 (4.82)	332 (100)
Residence						
Korogocho	523 (16.24)	1491 (46.29)	614 (19.06)	346 (10.74)	247 (7.67)	3221 (100)
Viwandani	413 (18.18)	1018 (44.81)	397 (17.47)	233 (10.26)	211 (9.29)	2272 (100)
Year of death						
2002–2005	148 (10.06)	636 (43.24)	286 (19.44)	161 (10.94)	240 (16.32)	1471 (100)
2006–2009	142 (9.91)	578 (40.33)	331 (23.10)	231 (16.12)	151 (10.54)	1433 (100)
2010–2013	462 (26.32)	917 (52.25)	245 (13.96)	80 (4.56)	51 (2.91)	1755 (100)
2104–2016	184 (17.04)	378 (45.27)	150 (17.96)	107 (12.81)	16 (8.34)	835 (100)
Place of death						
House	286 (15.04)	878 (46.19)	364 (19.15)	209 (10.99)	164 (8.63)	1901 (100)
Health facility	432 (16.88)	1162 (45.39)	479 (18.71)	271 (10.59)	216 (8.44)	2560 (100)
Enroute to health facility	65 (18.62)	150 (42.98)	63 (18.05)	44 (12.61)	27 (7.74)	349 (100)
Other	151 (22.47)	315 (46.88)	102 (15.18)	54 (8.04)	50 (7.44)	672 (100)
Missing	2 (16.67)	4 (33.33)	4 (33.33)	1 (8.33)	1 (8.33)	12 (100)
Respondent lives in the HH						
Yes	470 (19.00)	1118 (45.19)	441 (17.83)	272 (10.99)	173 (6.99)	2474 (100)
No	360 (18.58)	873 (45.05)	362 (18.68)	198 (10.22)	145 (7.48)	1938 (100)
Not in the universe	67 (10.29)	312 (47.93)	110 (16.90)	52 (7.99)	110 (16.90)	651 (100)
Missing	39 (9.05)	206 (47.80)	99 (22.97)	57 (13.23)	30 (6.96)	431 (100)
Sought health care						
Yes	662 (15.43)	1957 (45.62)	814 (18.97)	485 (11.31)	372 (8.67)	4290 (100)
No	263 (23.11)	524 (46.05)	186 (16.34)	87 (7.64)	78 (6.85)	1138 (100)
Don't know	4 (10.53)	16 (42.11)	8 (21.05)	3 (7.89)	7 (18.42)	38 (100)
Missing	7 (25.00)	12 (42.86)	4 (14.29)	4 (14.29)	1 (8.34)	28 (100)

Table 3. Crude and adjusted logistic regression analysis to examine the effect of recall time on the VA cause of death assignment groups.

Disease/Recall time	OR (95% CI)	P-Value ^a	P-value ^b	AOR (95%CI) ^y	P- Value ^a	P-Value ^b
HIV/AIDS						
<1 Month	0.7 (0.55-0.89)	0.004	<0.0001	0.85 (0.66-1.09)	0.198	0.0124
1–3 Months	Ref			Ref		
4–6 Months	1.31 (1.07-1.60)	0.008		1.24 (1.01-1.54)	0.043	
7–12months	1.26 (0.99-1.61)	0.060		1.16 (10.89-1.52)	0.256	
>12 months	0.98 (0.73-1.30)	0.876		0.77 (0.57-1.05)	0.106	
Malaria -Anaemia						
<1 Month	0.67 (0.33-1.34)	0.254	0.3285	0.72 (0.35-1.46)	0.359	0.3606
1–3 Months	Ref			Ref		
4–6 Months	0.93 (0.51-1.69)	0.808		0.91 (0.50-1.67)	0.767	
7–12months	0.86 (0.40-1.86)	0.710		0.72 (0.33-1.57)	0.408	
>12 months	1.66 (0.86-3.19)	0.128		1.65 (0.82-3.32)	0.158	
Tuberculosis						
<1 Month	0.85 (0.70-1.06)	0.148	0.0442	0.90 (0.71-1.13)	0.381	0.7997
1–3 Months	Ref			Ref		
4–6 Months	1.08 (0.89-1.31)	0.440		1.01 (0.81-1.24)	0.963	
7–12months	0.89 (0.70-1.15)	0.373		0.89 (0.67-1.18)	0.437	
>12 months	1.30 (1.02-1.67)	0.036		1.05 (0.79-1.41)	0.731	
Respiratory tract infections						
<1 Month	1.05 (0.81-1.37)	0.707	0.0714	1.17 (0.88-1.54)	0.280	0.0585
1–3 Months	Ref			Ref		
4–6 Months	1.11 (0.86-1.42)	0.438		1.12 (0.86-1.47)	0.380	
7–12months	1.28 (0.95-1.72)	0.105		1.12 (0.81-1.54)	0.493	
>12 months	1.54 (1.13-2.11)	0.006		1.70 (1.21-2.41)	0.002	
Meningitis						
<1 Month	0.82 (0.54-1.23)	0.333	0.4481	0.83 (0.55-1.26)	0.378	0.4409
1–3 Months	Ref			Ref		
4–6 Months	0.93 (0.64-1.36)	0.709		0.89 (0.60-1.31)	0.553	
7–12months	0.94 (0.59-1.51)	0.802		0.83 (0.51- 1.34)	0.447	
>12 months	1.38 (0.88-2.16)	0.164		1.35 (0.08-2.19)	0.217	
Other infectious diseases						
<1 Month	1.16 (0.86-1.55)	0.329	0.0953	1.40 (1.02-1.92)	0.038	0.0214
1–3 Months	Ref			Ref		
4–6 Months	0.80 (0.58-1.10)	0.176		0.75 (0.54-1.06)	0.102	
7–12months	1.34 (0.95-1.87)	0.091		1.11 (0.78-1.60)	0.560	
>12 months	0.85 (0.55-1.32)	0.474		0.75 (0.47-1.20)	0.227	

Disease/Recall time	OR (95% CI)	P-Value ^a	P-value ^b	AOR (95%CI) ^y	P- Value ^a	P-Value ^b
Maternal/neonatal						
<1 Month	0.61 (0.43-0.86)	0.005	0.0618	0.57 (0.40-0.82)	0.002	0.024
1–3 Months	Ref			Ref		
4–6 Months	0.97 (0.73-1.29)	0.820		1.01 (0.75-1.36)	0.962	
7–12months	0.97 (0.62-1.28)	0.527		0.98 (0.67-1.43)	0.91	
>12 months	0.85 (0.56-1.28)	0.431		1.02 (0.66-1.58)	0.938	
Malignancies						
<1 Month	1.14 (0.74-1.75)	0.562	0.7464	1.08 (0.69-1.71)	0.726	0.5190
1–3 Months	Ref			Ref		
4–6 Months	0.83 (0.52-1.33)	0.448		0.79 (0.48-1.29)	0.340	
7–12months	1.04 (0.61-1.78)	0.890		1.05 (0.59-1.86)	0.871	
>12 months	1.24 (0.72-2.16)	0.440		1.44 (0.79-2.65)	0.228	
Injuries/other external causes						
<1 Month	1.62 (1.34-1.94)	<0.001	>0.001	1.28 (0.96-1.70)	0.094	0.2626
1–3 Months	Ref			Ref		
4–6 Months	0.86 (0.70 -1.06)	0.152		1.06 (0.78-1.44)	0.701	
7–12months	0.84 (0.65- 1.09)	0.180		1.37 (0.93-2.011)	0.108	
>12 months	0.85 (0.64-1.13)	0.260		0.91 (0.58-1.42)	0.673	
Cardiovascular						
<1 Month	1.34 (0.99-1.82)	0.057	0.0202	1.36 (0.98-1.89)	0.064	0.0858
1–3 Months	Ref			Ref		
4–6 Months	1.19 (0.88-1.62)	0.252		1.15 (0.83-1.60)	0.394	
7–12months	0.77 (0.49-1.20)	0.240		0.78 (0.48-1.25)	0.299	
>12 months	0.64 (0.38-1.09)	0.100		0.70 (0.40-1.24)	0.226	
Other Chronic NCDs						
<1 Month	1.03 (0.72-1.49)	0.180	0.911	0.97 (0.67-1.41)	0.880	0.4877
1–3 Months	Ref			Ref		
4–6 Months	1.05 (0.74-1.49)	0.792		1.11 (0.78-1.60)	0.563	
7–12months	1.20 (0.79-1.82)	0.382		1.36 (0.88-2.09)	0.162	
>12 months	1.16 (0.73-1.85)	0.518		1.40 (0.86-2.28)	0.175	
Indeterminate						
<1 Month	0.88 (0.69-1.11)	0.268	0.0027	0.92 (0.71-1.17)	0.490	0.0339
1–3 Months	Ref			Ref		
4–6 Months	0.79 (0.62-0.99)	0.044		0.77 (0.60-0.98)	0.036	
7–12months	0.82 (0.62-1.10)	0.200		0.80 (0.59-1.09)	0.158	
>12 months	0.51 (0.35-0.74)	0.000		0.59 (0.39-0.88)	0.011	

^aWald p value.^bchi square p value^yVariables adjusted in the model: Gender, Age, Slum area, Year of death, Place of death, Respondent lives in household, Sought health care^dLikelihood ratio p value.

OR = odds ratio, CI = confidence interval, AOR = adjusted odds ratio, Ref = reference group, HIV = human immunodeficiency virus, AIDS = acquired immunodeficiency syndrome, NCDs = non-communicable diseases.

cause of death was dependent on the recall time in the unadjusted analysis (p value= 0.0442) but this effect was lost in the adjusted analysis.

In the crude analysis, recall time did not affect the identification of other infectious diseases as the probable cause of death but we found an effect of recall time in the adjusted analysis (p-value = 0.024). Compared to doing the VA one to three months after the death, conducting it in less than one month increased the chance of identifying other infectious diseases as the probable cause of death by 40 per cent (AOR 1.4; 95% CI 1.02-1.92). Similarly, identifying maternal/neonatal causes of death did not depend on the recall time in the crude analysis but in the adjusted analysis, there was an effect of recall time with a 43% lower chance of assigning probable cause of death to maternal/neonatal causes in VAs done less than one month compared to those done between 1 to 3 months (AOR 0.57; 95%CI 0.40-0.82; p-value=0.002).

Discussion

In this secondary analysis, we investigate the effect of recall time on the cause of death assigned through the InterVA-4 software. There was a notable variation in the timing of VAs with a higher percentage of VAs being conducted within three months of death in the latter years of follow up (2010 – 2016) as compared to the earlier years (2002–2009) suggesting an improved death notification and follow up system in the latter years.

The recall time in a VA is crucial as it determines if the respondent(s) can accurately recount the prevailing symptoms, signs and probable diagnosis before the deceased person's death²¹. The InterVA-4, as well as other VA systems, rely on the accuracy of this information to assign the possible cause of death. In our adjusted analysis, we found a significant difference in the odds of the assignment of a cause of death as HIV/AIDS, other infectious disease and maternal/neonatal causes by recall time. We found that there was a 40% higher chance of identifying other infectious diseases as the cause of death in our analysis for VAs conducted less than one month of death as compared to those conducted between one to three months.

The timing of VAs should consider prevailing circumstances to the death and ensure the appropriate mourning period be observed based on the accepted cultural norms and religion. In a study²² looking at effects of culture and ethics on VAs in Ghana, it was noted that the timing should take into consideration the respondent's emotional distress, for example from the death of an only child or a maternal death. The mourning period may also vary based on the relationship between the interviewee(s) and the deceased as well as the nature of the death and role of the deceased person in the household²². The Kenya verbal autopsy guidelines provide for a 30–40-day period of mourning before conducting a VA with a maximum allowable period of one year after the death²⁰. In our analysis, there was a 43% less probability of assigning maternal/neonatal cause

of death for VAs conducted less than one month after the death as compared to those conducted between one to three months.

Our results, in general, find no effect of recall time on identifying the cause of death by VA for most chronic diseases (except HIV/AIDS) as compared to acute conditions. We postulate that VA respondents in situations where the deceased had a chronic disease are more likely to remember the symptoms, signs and the probable cause of death overtime.

Delay of VA for periods beyond 12 months has been hypothesised to lead to a less accurate recall of symptoms¹². However, our analysis did not show any significant differences for any of the identified causes of deaths for recall periods above 12 months. In a study in South Africa, the only significant difference for recall periods over 12 months were for neonatal deaths¹³.

The ideal method of identifying the effect of recall time would have been a comparison of the cause of death generated from gold standard death certification to the VA assigned cause of death for the different recall times. However, in the absence of the gold standard comparator, as in our case, the alternative approach used in this paper was to compare probabilities or odds of making a specific VA cause of death assignment for different recall periods.

Conclusion

Recall time affected the assignment of VA cause of death for HIV/AIDS, other infectious diseases and maternal/neonatal causes. Our analysis indicates that in the urban informal setting, VAs should be conducted from one month up to 6 months after the death to improve the probability of accurately assigning the cause of death.

Data availability

The [KENYA - NUHDSS - Verbal Autopsy, Causes of deaths 2002–2015](#) dataset was used in this secondary analysis. The dataset is available upon request from the APHRC Microdata Portal <http://microdataportal.aphrc.org/index.php/catalog>. Access to the dataset can be obtained through submission of a written request in the APHRC portal following creation of an account. Further information on how to apply for access to the data can be found [here](#).

Author Contributions

Conceptualization: DM, JG; Data curation: DM, JG Formal Analysis: DM, JG; Writing-original draft preparation: DM, JG; Writing-Review & Editing: DM, JG

Acknowledgements

The authors wish to acknowledge APHRC as the source of the data utilized for the secondary analysis

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Aaron S. Karat

TB Centre, London School of Hygiene & Tropical Medicine, London, UK

Thank you for the opportunity to review this paper, which is generally well written and deals with an important aspect of verbal autopsy implementation. This is a secondary analysis of VA data from more than 5,000 deaths and, though the findings are interesting, I have some concerns around some of the methods and around the interpretation of results.

Major comments:

1. Calculation of CSMFs - The method used to calculate CSMFs is potentially problematic. InterVA output (up to three causes with likelihoods +/- residual likelihood) is intended to be converted into CSMFs in a way that takes account of all causes of death assigned (see Appendix 4 of the InterVA-4 user guide), not only the most likely cause, as described by the authors. The 'collapsing' process described seems perfectly justifiable but may have been better applied after the calculation of CSMFs to reflect better the true output of InterVA. Adopting this approach would likely have produced different CSMFs for each recall period, and may therefore have affected the results of the regression.
2. Logistic regression - Although most factors likely to affect the relationship between recall time and cause of death were adjusted for, time of year/seasonality was not. This is potentially very important, as it may have affected mortality patterns but also affected the ability of VA teams to access respondents (e.g. this may have been harder in the rainy season) and therefore affected time between death and VA. This may be less relevant in the urban setting where this study was conducted (compared with the rural setting in the study by Hussain-Alkhateeb *et al.*) but this should at least have been discussed and the omission justified by the authors.
3. Interpretation of results - My apologies if I have misunderstood, but I do not entirely follow some of the statements made in the discussion and I am not sure the conclusions flow from the results presented. The authors suggest that respondents may be more likely to remember the symptoms of an acute illness if the VA is done in a shorter time after death, yet in this analysis the odds of being assigned a respiratory tract infection CoD were higher

for VAs done after >12 months compared with 1–3 months.

4. I also do not entirely follow the possible explanation for why maternal deaths were less likely to be assigned <1 months after death than 1–3 months. Are the authors suggesting that fewer maternal deaths were assigned because VAs may have been delayed when a particular cause of death was suspected? It would seem important to know if this was done when any other cause of death was suspected, or if it was only for maternal deaths. The authors also state that the Kenyan guidelines recommend 30–40 days mourning prior to conducting a VA, but there is no description of how often these guidelines were followed (and/or if they were followed more closely in one district than another) – this is a potentially a major source of bias and should ideally have been explored in more detail in the analysis.

Minor comments:

Introduction:

1. **Paragraph 1:** The ‘gold standard’ for cause of death is one assigned by complete diagnostic autopsy, not by a medical practitioner, though of course for the majority of deaths a practitioner-assigned CoD is often the best that can be hoped for. It may be worth making this point a little more clearly in the first paragraph.
2. **Paragraph 2:** This is a very minor point, but the multiple plurals used here (‘interviews’, ‘questionnaires’, ‘relatives’, ‘caregivers’) suggests that a standard VA involves interviewing more than one person, when this is rarely the case.
3. **Paragraph 2:** I believe the full name of the organisation is the Population Health Metrics Research Consortium. A better reference when discussing the VA instrument (instead of the paper currently cited in Ref 7) may be the ‘VA tools’ page on the [PHMRC website](#).
4. **Paragraph 3:** the list of CCVA methods is slightly dated (the ref provided is a 2014 review). [This pre-print](#) may provide a more up-to-date list (InterVA is also misspelled in this paragraph, as “inter-VA”).

Methods:

1. If possible, please could the authors provide the exact cut-offs (in days) used for the recall time categories. At present it is not entirely clear if a VA conducted after ~3.5 months would be categorised as “1–3 months” or “4–6 months” (same for recall times between 6 and 7 months).
2. There may be a good reason for using 1–3 months as a reference instead of <1 month but the reason for this approach is not described in the methods. In my opinion this makes it more difficult to interpret the results and to look at trends across recall periods (regardless of statistical ‘significance’).

Results

1. **Page 4:** it may be more interesting to show the median + IQR for recall time, given that it is unlikely to be normally distributed?
2. **Table 2:** It is not clear what ‘Not in the universe’ means (under ‘Respondent lives in the HH’).
3. **Table 2:** Typo under ‘year of death (“2104”).

4. **Table 3:** It would make the results more transparent if the CSMF proportion for each cause was shown in this table, very ideally for each recall period for each cause, so readers are clear on the 'number' of outcomes associated with each OR/aOR.

5. **Table 3:** Typo in the 95% CI for the aOR for HIV/AIDS deaths with a recall time of 7–12 months ("10.89").

Is the work clearly and accurately presented and does it cite the current literature?

Partly

Is the study design appropriate and is the work technically sound?

Partly

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Partly

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Tuberculosis, epidemiology, infectious diseases, verbal autopsy.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 03 Feb 2021

Donnie Mategula, Malawi-Liverpool-Wellcome Trust Clinical Research Programme , P.O Box 30096, Blantyre, Malawi

Dr Aaron S.Karat Many thanks for your helpful and thoughtful comments. We have re-submitted an updated manuscript which addresses your suggestions and concerns. Please find our detailed responses below.

Major comments:

1) Calculation of CSMFs - The method used to calculate CSMFs is potentially problematic. Inter-VA output (up to three causes with likelihoods +/- residual likelihood) is intended to be converted into CSMFs in a way that takes account of all causes of death assigned (see

Appendix 4 of the InterVA-4 user guide), not only the most likely cause, as described by the authors. The 'collapsing' process described seems perfectly justifiable but may have been better applied after the calculation of CSMFs to reflect better the true output of InterVA. Adopting this approach would likely have produced different CSMFs for each recall period, and may therefore have affected the results of the regression.

Response: Thank you for the comment. The cause specific mortality fractions have been re-calculated as suggested and are presented in table 2 of the revised paper. However, we have not used the CSMFs for the regression analysis as adding the cause of death specific weights/likelihoods to a model where the cause of death have been collapsed makes assumptions on how the cause of death weights/likelihoods combine before and after the re-categorization. To avoid making these assumptions, we prefer to use the most likely cause of death in the logistic regression analysis. However, we have included this limitation in the discussion.

2) Logistic regression - Although most factors likely to affect the relationship between recall time and cause of death were adjusted for, time of year/seasonality was not. This is potentially very important, as it may have affected mortality patterns but also affected the ability of VA teams to access respondents (e.g. this may have been harder in the rainy season) and therefore affected time between death and VA. This may be less relevant in the urban setting where this study was conducted (compared with the rural setting in the study by Hussain-Alkhateeb *et al.*) but this should at least have been discussed and the omission justified by the authors.

Response: Seasons and weather patterns in Nairobi have varied considerably over the period of analysis in the paper, it would have been less prudent to include seasonality in the model. Additionally, we agree with the reviewer that this variable is less relevant in this context unlike in the Hussain Alkhateeb *et al* paper. We have included text in the discussion section to reflect our rationale for the lack of inclusion of this variable in the model.

3) Interpretation of results - My apologies if I have misunderstood, but I do not entirely follow some of the statements made in the discussion and I am not sure the conclusions flow from the results presented. The authors suggest that respondents may be more likely to remember the symptoms of an acute illness if the VA is done in a shorter time after death, yet in this analysis the odds of being assigned a respiratory tract infection CoD were higher for VAs done after >12 months compared with 1–3 months.

Response: In our analysis, there is no overall association between recall time and RTI based on the likelihood ratio test (P value 0.0585) and that is why we did not focus on RTIs. Our general observation that respondents are more likely to remember acute illness if the VA is done in a shorter time still holds.

4) I also do not entirely follow the possible explanation for why maternal deaths were less likely to be assigned <1 months after death than 1–3 months. Are the authors suggesting that fewer maternal deaths were assigned because VAs may have been delayed when a

particular cause of death was suspected? It would seem important to know if this was done when any other cause of death was suspected, or if it was only for maternal deaths. The authors also state that the Kenyan guidelines recommend 30–40 days mourning prior to conducting a VA, but there is no description of how often these guidelines were followed (and/or if they were followed more closely in one district than another) – this is a potentially a major source of bias and should ideally have been explored in more detail in the analysis.

Response: Thank you for the comment. We have updated the discussion section to further explain the potential sources of delay to conducting a VA in the NUHDSS. This was not specific to maternal deaths but to the prevailing circumstances. We have edited the discussion to focus less on the maternal/neonatal finding. While the numbers for combined direct obstetric and neonatal causes were low, the majority of these were neonatal deaths. The effect observed could therefore have been a chance finding due to the few numbers or as a result of the higher proportion of neonatal deaths to direct obstetric causes (6 to 1) especially in cases where the neonatal deaths were not reviewed promptly.

Minor comments:

Introduction:

5) Paragraph 1: The 'gold standard' for cause of death is one assigned by complete diagnostic autopsy, not by a medical practitioner, though of course for the majority of deaths a practitioner-assigned CoD is often the best that can be hoped for. It may be worth making this point a little more clearly in the first paragraph.

Response: This has been updated

6) Paragraph 2: This is a very minor point, but the multiple plurals used here ('interviews', 'questionnaires', 'relatives', 'caregivers') suggests that a standard VA involves interviewing more than one person, when this is rarely the case.

Response: Multiple plurals have been reviewed and revised

7) Paragraph 2: I believe the full name of the organisation is the Population Health Metrics Research Consortium. A better reference when discussing the VA instrument (instead of the paper currently cited in Ref 7) may be the 'VA tools' page on the [PHMRC website](#).

Response: The changes have been made and the reference updated

8) Paragraph 3: the list of CCVA methods is slightly dated (the ref provided is a 2014 review). [This pre-print](#) may provide a more up-to-date list (InterVA is also misspelled in this paragraph, as "inter-VA").

Response: We have updated the list of CCVA methods based on the updated reference, however we have retained our original citation as well since Mapundo et al paper is in a preprint.

Methods:

9) If possible, please could the authors provide the exact cut-offs (in days) used for the recall time categories. At present it is not entirely clear if a VA conducted after ~3.5 months would be categorised as "1–3 months" or "4–6 months" (same for recall times between 6 and 7 months).

Response: Thank you for the comment, the highest cutoff integer month for our categories included all the possible decimals before the next category, for example 1-3 months includes all possible days up to 3.99. We have updated this in the methods section.

10) There may be a good reason for using 1–3 months as a reference instead of <1 month but the reason for this approach is not described in the methods. In my opinion this makes it more difficult to interpret the results and to look at trends across recall periods (regardless of statistical 'significance').

Response: The 1 to 3 months recall time was used as the reference category for the logistic regression for two reasons; the highest number of VAs were conducted within this period and secondly this falls within the Kenya VA guidelines on appropriate time to conduct a VA thus we set to compare all the results against this recall period. We have updated this in the methods section.

Results

11) Page 4: it may be more interesting to show the median + IQR for recall time, given that it is unlikely to be normally distributed?

Response: Thanks, we concur. The Median and IQR for recall time have been included.

12) Table 2: It is not clear what 'Not in the universe' means (under 'Respondent lives in the HH').

Response: Implies that the response given did not apply to the context of the deceased under question. We have added a footnote in the table to explain

13) Table 2: Typo under 'year of death ("2104").

Response: This has been amended, thank you

14) Table 3: It would make the results more transparent if the CSMF proportion for each cause was shown in this table, very ideally for each recall period for each cause, so readers are clear on the 'number' of outcomes associated with each OR/aOR.

Response: We have added the number of outcomes associated with each OR/aOR. Please also refer to our response to comment number 1 to justify our choice.

14 Table 3: Typo in the 95% CI for the aOR for HIV/AIDS deaths with a recall time of 7–12 months ("10.89").

Response: This has been amended, thank you

Competing Interests: No competing interests were disclosed.

Reviewer Report 02 December 2020

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Lucia D'Ambruoso

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² University of the Witwatersrand, Johannesburg, South Africa

Overall comments:

This is an interesting paper examining recall times on the outcomes of VA surveys and contributes to the evidence base on this methodology. The paper is generally well written and clearly presented. Overall comments are as follows: first, the paper would be improved with fuller justification of key assumptions and aspects of the analytical approach around ideal recall times and CoD categories (see specific comments). Second, while I am not well-qualified to comment on statistical aspects, it would appear that not all the statistically significant findings are considered in the narrative. Third, the discussion could be further developed by considering the plausibility of the main findings, offering reasons as to why the recall time effects on some cause of death categories were found. Fourth, further attention to the social and cultural aspects of recall time, and in the context of Covid-19/other infectious outbreaks would add to the current relevance of the paper. Finally, a statistician should review the paper. As a qualitative researcher, I have not commented on the statistical approach in anything more than general terms. The paper is likely to be worthy of indexing, with attention to these issues.

Specific comments:

Introduction:

1. Paragraph 3: the authors may wish to acknowledge that computer-coded VA addresses observed inconsistencies within and between physician coders, as well as obviating the time implications of physician coding: in the settings where VA is needed, using physician time for VA interpretation is ethically problematic.
2. Paragraph 4: the need to account for social and cultural factors as a key factor determining recall period is convincingly introduced – how the authors deal with this issue is not clear however and could usefully be expanded on in the paper.

Methods:

1. Paragraph 1, Study setting: A map would be good to include, if possible.
2. Paragraph 3, Design: The term demographic health system may be misleading, suggest replace and use HDSS consistently.
3. Paragraph 3, Design: InterVA5 model is currently available, it may be worth considering applying InterVA5 for these data.
4. Page 4, Table 1: the title of this table might be rephrased - strictly speaking, criteria are not presented. Recategorization of InterVA CoDs to constructed CoD categories might be more accurate.
5. Page 4, Table 1: The collapsed CoD categories are not quite clear. The authors state that category classifications were guided by counts and reasonable relatedness - this requires some clarification, especially around amnesia and malaria. The authors also state that the collapsed categories are based on classifications used in other work, however there are 8 categories in the Hussain *et al.* study and 15 in the Oti and Kyobutungi paper. The relationship of these to the 12 categories used in this analysis similarly would benefit from some clarification.
6. Page 4, Table 1: It is not customary to cite AIDS/HIV as a category, rather HIV/AIDS, as is done later in the paper – suggest consistency with commonly used term.
7. Page 4, paragraph 1: CSMFs typically account for all probable CoD assigned per case. Suggest recalculating the CSMF using all probable CoDs from InterVA.
8. It is not entirely clear why 1-3 months recall time is used as the reference period. The paper indeed presents research from the same region where 12 months recall is found to be suitable. The authors may wish to clarify this element of the analysis.

Results:

- While I am not well-qualified to comment on the statistical analyses, it would appear that not all significant findings from the adjusted regression analysis are highlighted in the narrative, please see (d) and (e) below. The main findings would appear to be as follows: Compared to a recall period of 1-3 months.
 - (a) Cases with recall periods of 4-6 months are 24% more likely to be classified as due to HIV;
 - (b) Cases with recall periods of <1 month 40% more likely to be classified as due to other infectious causes.
 - (c) Cases with recall periods of <1 month 43% less likely to be classified as due to maternal causes.
 - (d) Cases with recall periods of >12 months 70% more likely to be classified as due to RTI.
 - (e) Cases with recall periods of >12 months 41% less likely to be classified as indeterminate.

Discussion:

1. The discussion could be further developed by considering the plausibility of the main findings offering reasons as to why the recall time effects on some cause of death categories were found. For example, that cases with recall periods of <1 month are 43% less likely to be classified as due to maternal causes is counter-intuitive – maternal deaths are relatively rare, generally avoidable and therefore likely to be highly memorable.
2. Page 9, paragraph 5: the statement 'our analysis did not show any significant differences for any of the identified causes of deaths for recall periods above 12 months' needs to be checked. Again, from a general perspective, Table 3 appears to suggest that deaths investigated >12 months are (a) 70% more likely to be classified as due to RTI (AOR 1.70 (CI 1.21-2.41) P=0.002) and (b) 41% less likely to be diagnosed by InterVA as indeterminate (AOR 0.59 (CI 0.39-0.88) P=0.011).
3. Further attention to social and cultural influences on recall times, and how these vary by country/region would be useful in this section. Thorough assessment of social and cultural norms prior to VA surveys is of critical importance, whether these factors have an impact on the findings presented in this paper would be useful to consider.
4. Some attention to how the analysis and recommendations on recall periods related to the Covid-19 pandemic and other infectious outbreaks, e.g., Ebola, is likely to be of relevance to readers.
5. The authors may wish to include strengths and limitations section, where a sensitivity analysis with e.g., up to 12 months as an ideal recall period (as is indicated in other research) is applied.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Partly

Are sufficient details of methods and analysis provided to allow replication by others?

Partly

If applicable, is the statistical analysis and its interpretation appropriate?

I cannot comment. A qualified statistician is required.

Are all the source data underlying the results available to ensure full reproducibility?

Partly

Are the conclusions drawn adequately supported by the results?

Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Global public health; qualitative methods; health policy and systems research; South Africa

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 03 Feb 2021

Donnie Mategula, Malawi-Liverpool-Wellcome Trust Clinical Research Programme , P.O Box 30096, Blantyre, Malawi

Dr Lucia D'Ambruoso, Many thanks for your helpful and thoughtful comments. We have re-submitted an updated manuscript which addresses your suggestions and concerns. Please find our detailed responses below:

Introduction:

Paragraph 3: the authors may wish to acknowledge that computer-coded VA addresses observed inconsistencies within and between physician coders, as well as obviating the time implications of physician coding: in the settings where VA is needed, using physician time for VA interpretation is ethically problematic.

Response: Thank you for the suggestion. This has been included.

Paragraph 4: the need to account for social and cultural factors as a key factor determining recall period is convincingly introduced – how the authors deal with this issue is not clear however and could usefully be expanded on in the paper.

Response: We have considered the social factors that were available in the dataset in the regression analysis. In the discussion, we have expounded on some of the potential social-cultural factors around recall period not available in the dataset.

Methods:

Paragraph 1, Study setting: A map would be good to include, if possible. A reference to where the NUHDSS map can be accessed has been provided in the revised version.

Paragraph 3, Design: The term demographic health system may be misleading, suggest replace and use HDSS consistently.

Response: This has been updated.

Paragraph 3, Design: InterVA5 model is currently available, it may be worth considering applying InterVA5 for these data.

Thank you for the comment. The data set was provided following the Inter-VA4 analysis therefore it would not be possible to analyze it using Inter-VA5. InterVA-5 was developed to accommodate the WHO 2016 standard VA tool that was not utilized for this data set.

Page 4, Table 1: the title of this table might be rephrased - strictly speaking, criteria are not presented. Recategorization of InterVA CoDs to constructed CoD categories might be more accurate.

Response: Thank you. This has been updated

Page 4, Table 1: The collapsed CoD categories are not quite clear. The authors state that category classifications were guided by counts and reasonable relatedness - this requires some clarification, especially around anaemia and malaria. The authors also state that the collapsed categories are based on classifications used in other work, however there are 8 categories in the Hussain *et al.* study and 15 in the Oti and Kyobutungi paper. The relationship of these to the 12 categories used in this analysis similarly would benefit from some clarification.

Response: We have updated the text to further detail how we did the recategorization. The approach to this classification has been used by other studies as cited, however the final number of categories varies depending on data and specific circumstances in the study areas. The collapsing of malaria and anaemia is based on the premise that malaria is a major cause of anaemia in tropical areas.

Page 4, Table 1: It is not customary to cite AIDS/HIV as a category, rather HIV/AIDS, as is done later in the paper – suggest consistency with commonly used term. This has been updated. Thank you.

Page 4, paragraph 1: CSMFs typically account for all probable CoD assigned per case. Suggest recalculating the CSMF using all probable CoDs from InterVA. CSMFs have been recalculated as suggested. It is not entirely clear why 1-3 months recall time is used as the reference period. The paper indeed presents research from the same region where 12 months recall is found to be suitable. The authors may wish to clarify this element of the analysis.

Response: The 1 to 3 months recall time was used as the reference category for the logistic regression for two reasons; the highest number of VAs were conducted within this period and secondly this falls within the Kenya VA guidelines on appropriate time to conduct a VA thus we set to compare all the results against this recall period. We have updated this in the methods section.

Results:

While I am not well-qualified to comment on the statistical analyses, it would appear that not all significant findings from the adjusted regression analysis are highlighted in the narrative, please see (d) and (e) below. The main findings would appear to be as follows: Compared to a recall period of 1-3 months.

- (a) Cases with recall periods of 4-6 months are 24% more likely to be classified as due to HIV;
- (b) Cases with recall periods of <1 month 40% more likely to be classified as due to other infectious causes.

- (c) Cases with recall periods of <1 month 43% less likely to be classified as due to maternal causes.
- (d) Cases with recall periods of >12 months 70% more likely to be classified as due to RTI.
- (e) Cases with recall periods of >12 months 41% less likely to be classified as indeterminate.

Response: Thank you. We have updated the missing significant results for the indeterminate, however we have left out the RTIs as the association with recall time is not significant on the overall likelihood ratio test of association.

Discussion:

The discussion could be further developed by considering the plausibility of the main findings offering reasons as to why the recall time effects on some cause of death categories were found. For example, that cases with recall periods of <1 month are 43% less likely to be classified as due to maternal causes is counter-intuitive – maternal deaths are relatively rare, generally avoidable and therefore likely to be highly memorable.

Response: While the numbers for combined direct obstetric and neonatal causes were low, the majority of these were neonatal deaths. The effect observed could therefore have been a chance finding due to the few numbers or as a result of the higher proportion of neonatal deaths to direct obstetric causes (6 to 1) especially in cases where the neonatal deaths were not reviewed promptly.

Page 9, paragraph 5: the statement 'our analysis did not show any significant differences for any of the identified causes of deaths for recall periods above 12 months' needs to be checked. Again, from a general perspective, Table 3 appears to suggest that deaths investigated >12 months are (a) 70% more likely to be classified as due to RTI (AOR 1.70 (CI 1.21-2.41) P=0.002) and (b) 41% less likely to be diagnosed by InterVA as indeterminate (AOR 0.59 (CI 0.39-0.88) P=0.011).

Response: Thank you for the observation. We have revised the statement and included the findings on the indeterminate causes of death. In our analysis, there is no evidence of an overall association between recall time and RTI based on the likelihood ratio test (P value 0.0585) and that is why we did not focus on RTIs in the results section

Further attention to social and cultural influences on recall times, and how these vary by country/region would be useful in this section. Thorough assessment of social and cultural norms prior to VA surveys is of critical importance, whether these factors have an impact on the findings presented in this paper would be useful to consider.

Response: Thank you for the suggestion. We have included an assessment of the potential social and cultural circumstances prior to a VA.

Some attention to how the analysis and recommendations on recall periods related to the Covid-19 pandemic and other infectious outbreaks, e.g., Ebola, is likely to be of relevance to readers.

Response: Thank you for the suggestion, however, given the period within which the VA were conducted (2002-2016) it might be difficult to extrapolate the findings to the current situation of Covid-19.

Competing Interests: There no competing interests