


Premature adult mortality in India: what is the size of the matter?

Chalapathi Rao ¹, Aashish Gupta,² Mamta Gupta,³ Ajit Kumar Yadav⁴

To cite: Rao C, Gupta A, Gupta M, *et al.* Premature adult mortality in India: what is the size of the matter? *BMJ Global Health* 2021;**6**:e004451. doi:10.1136/bmjgh-2020-004451

Handling editor Soumitra S Bhuyan

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/bmjgh-2020-004451>).

Received 14 November 2020
Accepted 3 May 2021



© Author(s) (or their employer(s)) 2021. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

¹Research School of Population Health, Australian National University College of Medicine Biology and Environment, Canberra, Australian Capital Territory, Australia

²Demography and Sociology, University of Pennsylvania, Philadelphia, Pennsylvania, USA

³Alchemist Research and Data Analysis, Chandigarh, India

⁴Gender research project, International Institute for Population Sciences, Mumbai, Maharashtra, India

Correspondence to

Dr Chalapathi Rao;
chalapati.rao@anu.edu.au

ABSTRACT

Background Reducing adult mortality by 2030 is a key component of the United Nations Sustainable Development Goals (UNSDGs). Monitoring progress towards these goals requires timely and reliable information on deaths by age, sex and cause. To estimate baseline measures for UNSDGs, this study aimed to use several different data sources to estimate subnational measures of premature adult mortality (between 30 and 70 years) for India in 2017.

Methods Age-specific population and mortality data were accessed for India and its 21 larger states from the Civil Registration System and Sample Registration System for 2017, and the most recent National Family and Health Survey. Similar data on population and deaths were also procured from the Global Burden of Disease Study 2016 and the National Burden of Disease Estimates Study for 2017. Life table methods were used to estimate life expectancy and age-specific mortality at national and state level from each source. An additional set of life tables were estimated using an international two-parameter model life table system. Three indicators of premature adult mortality were derived by sex for each location and from each data source, for comparative analysis

Results Marked variations in mortality estimates from different sources were noted for each state. Assuming the highest mortality level from all sources as the potentially true value, premature adult mortality was estimated to cause a national total of 2.6 million male and 1.8 million female deaths in 2017, with Bihar, Maharashtra, Tamil Nadu, Uttar Pradesh and West Bengal accounting for half of these deaths. There was marked heterogeneity in risk of premature adult mortality, ranging from 351 per 1000 in Kerala to 558 per 1000 in Chhattisgarh among men, and from 198 per 1000 in Himachal Pradesh to 409 per 1000 in Assam among women.

Conclusions Available data and estimates for mortality measurement in India are riddled with uncertainty. While the findings from this analysis may be useful for initial subnational health policy to address UNSDGs, more reliable empirical data is required for monitoring and evaluation. For this, strengthening death registration, improving methods for cause of death ascertainment and establishment of robust mortality statistics programs are a priority.

INTRODUCTION

Over the past three decades, there have been declines in child mortality across the world, as a result of global and local actions under the United Nations Millennium

Key questions

What is already known?

- Reliable measures of mortality at adult ages are required for evidence-based health policy, monitoring and evaluation of progress towards health-related United Nations Sustainable Development Goals (UNSDGs).
- In the absence of reliable data from Civil Registration and Vital Statistics systems in many countries including India, these measures are largely derived from alternate data sources, data synthesis or modelling methods.

What are the new findings?

- This article presents a comparative analysis of measures of premature adult mortality from several data sources for India and its 21 larger states, examining their reliability and correspondence.
- Following a conservative approach, the article proposes the maximum estimate of mortality between the ages of 30 and 70 years by sex for each location from any source as the potential baseline level of premature adult mortality around 2016–2017.

What do the new findings imply?

- Although all the six data sources or estimation methods demonstrated some weaknesses, the adequate quality of data from the Civil Registration System (CRS) in several states suggests that through the implementation of strategic interventions, the CRS could be developed into a reliable data source for tracking progress towards the UNSDGs.
- The methods and findings from our analyses would be relevant for other countries facing the need to reconcile local mortality data with modelled mortality estimates from international sources.

Development Goals campaign.^{1 2} Concomitantly, mortality at adult ages has been recognised as a growing component of population level disease burden.^{3 4} For this reason, the health-related United Nations Sustainable Development Goals (UNSDGs) for 2030 include specific targets to reduce adult mortality from major non-communicable diseases (NCDs), tuberculosis, HIV/AIDS, road traffic accidents, suicides, and occupational and environmental exposures, among

other health conditions and risk factors.⁵ Monitoring progress towards these UNSDG targets requires routine and reliable measures of population level cause-specific mortality at adult ages, for which Civil Registration and Vital Statistics (CRVS) systems with medical certification of cause of death are the optimal data source.⁶ Unfortunately, reliable CRVS systems are not yet in place in many parts of the world, which limits the monitoring of adult mortality trends^{7 8}

Good-quality CRVS data have been directly used to analyse national adult mortality trends in high-income countries.⁹ For several Latin American countries and South Africa, CRVS data were first adjusted for incomplete death registration, prior to similar trend analyses.^{10 11} The findings from these studies enabled an improved understanding of underlying diseases and risk factors for adult mortality in these countries. In India, although death registration systems have existed for over 150 years, not all deaths are currently registered in the national Civil Registration System (CRS).¹² Levels of death registration completeness vary by gender, age and location across the country, and these data gaps have limited the reliability of direct adult mortality measures.¹³

The Sample Registration System (SRS) and National Family Health Survey (NFHS) programmes were established in 1970 and 1992, respectively, as alternate sources of mortality data, and had been specifically designed to enable reliable measurement of indicators of infant and under-five mortality.¹⁴ Although the SRS and NFHS programmes also compile information on adult deaths, the samples are not adequately powered for precise adult mortality measurement, especially at subnational levels.^{13 14} Despite these limitations, there have been several attempts in the past decade to estimate national and subnational adult mortality rates using SRS, NFHS and CRS data, but with varying calculation methods, indicators and adult age categories, which limits the comparability of results across methods or over time.^{13 15 16} As a result, patterns and determinants of adult mortality are less well understood.

More recently, there have been two initiatives that employed data synthesis methods using various data sources to derive national and sub national estimates of population and deaths by age and sex. The state-level results from the 'National Burden Estimates' (NBE) Study for 2017 were based on national level estimates of population and deaths derived for the United Nations (UN) World Population Project (WPP).^{17 18} Similar estimates of populations and deaths for each state are also available from the Indian State-level Burden of Disease initiative, which is a component of the Global Burden of Disease (GBD) Study for 2016, carried out by a collaboration between the Indian Council of Medical Research, the Public Health Foundation of India and the Institute of Health Metrics and Evaluation (IHME), University of Washington, USA.^{19 20} The NBE and GBD synthetic estimates add to the available empirical data sources, to help understand levels of mortality at adult ages.

To understand and address biological, environmental, behavioural, socioeconomic or health system factors that potentially influence patterns of disease burden and adult mortality across the country, there is an urgent need for reliable and timely gender specific subnational measures of adult mortality. The aim of this study was to conduct a comparative analysis of premature adult mortality from different data sources at national and state level around the period 2013–2017, to improve overall understanding of subnational levels and differentials in adult mortality patterns. We defined the interval between ages 30 and 70 years as the life span of premature adult mortality, and used three indicators to compare the magnitude of mortality at these ages across the different data sources and analytical approaches. This age interval corresponds to the indicator definition for NCD mortality under the UNSDGs.²¹ We propose that our findings on the maximum plausible levels of total mortality from all causes in this age interval across all data sources could be used as a suitable proxy baseline measure for monitoring progress in India and its states towards the UNSDG NCD targets, till such time that detailed information on causes of death is available. The findings are also used to make recommendations for improving adult mortality monitoring levels in India, during and beyond the UNSDG era.

METHODS

Data sources

This analysis is based on life table derived summary mortality measures by sex for India and 21 large states, during the period 2013–2017. **Table 1** presents the general characteristics of data on age-specific population and deaths from the CRS, SRS, NFHS, GBD and NBE data sources^{13 17 19 22 23} that were used for life table analysis. Relevant additional details are also available from the Guidelines for Accurate and Transparent Health Estimates Reporting (GATHER) statement for this study (see online supplemental appendix 1). The SRS life tables were based on observed population and deaths in sample sites during 2017. For the NFHS life tables, the household survey data were used to create lifelines to estimate deaths and person-years lived in the exposure period. For the CRS life tables, recorded deaths in 2017 and population projections for 2017 developed by the National Commission on Population (NCP) were used.²⁴ The NBE and GBD studies reported estimates of deaths by age and sex for each state derived from the GBD and NBE internal modelling processes.^{17 19 25} These studies also reported state specific populations by age and sex, derived from the United Nations World Population Prospects and the IHME international population models, respectively.^{18 20} The NBE data did not include separate estimates for states of Andhra Pradesh, Himachal Pradesh, Telangana and Uttarakhand. We used these available GBD and NBE population and death estimates in our life table calculations to derive the GBD and NBE based life tables for each state. The SRS proportionate

Table 1 Characteristics of inputs used from different data sources to estimate life tables for India and states, 2013–2017

Data source	Reference Year	Deaths	Population	Remarks
CRS	2017	Death registration	Projections made by National Commission on Population	Death age groups standardised to SRS
SRS	2017	Dual record systems	Observed sample population	Multistage random sample of census blocks
NFHS	2013–2016	Household recall	Person years lived in each age group within households	Multistage random cluster sampling of households
NBE	2017	State specific SRS proportions and national level UN WPP mortality/population estimates		Pop and death age groups standardised to SRS
GBD	2016	IHME mortality models	IHME population projections	State-specific mortality model inputs not available
MODMATCH	2017	WHO Model Life Table system	Projections made by National Commission on Population	Empirical state-specific inputs of ${}_5q_0$ and ${}_{45}q_{15}$ (see table 2)

CRS, Civil Registration System; GBD, Global Burden of Disease; IHME, Institute of Health Metrics and Evaluation; NBE, National Burden Estimates; NFHS, National Family Health Survey; SRS, Sample Registration System; UN, United Nations; WPP, World Population Project.

age distributions of deaths were used to interpolate the coarser age categories of deaths from CRS and NBE to the standard age categories for our life tables, which were as follows: 0–1 year, 1–4 years, 5–9 years80–84 years and 85+years.

Since life table quantities are not affected by the age structure of a population, period life tables are standard demographic tools which enable mortality comparisons across space and time. Period life expectancy (LE) at birth, for instance, is the average lifespan of a hypothetical cohort if they experienced the current set of age-specific mortality rates throughout their lifetime. Population and death data from each of the five sources for each location were used as inputs to construct life tables, using a standard spreadsheet programme.²⁶ From each data source-location-sex specific life table, the following summary mortality risks were extracted for comparison:

1. LE at birth.
2. Risk of dying between birth and age 5 years (${}_5q_0$).
3. Risk of dying between 15 and 60 years (${}_{45}q_{15}$), conditional on survival up to age 15.
4. Risk of dying between 30 and 70 years (${}_{40}q_{30}$), (risk of premature adult mortality) conditional on survival up to age 30.

Uncertainty intervals were estimated for each of the above parameters applying bootstrap methods using a publicly available programmed spreadsheet specifically designed for such analysis²⁷ (see online supplemental appendix 2).

Model life table estimation

In addition, we estimated a sixth set of life tables by sex for each location, using the WHO Modified Logit Life Table System, implemented through its customised software tool named ‘MODMATCH’.^{28 29} This system uses input values of ${}_5q_0$ and ${}_{45}q_{15}$ in a statistical model, to predict a complete schedule of age-specific mortality risks for the location of interest. Mortality patterns in the MODMATCH model life tables approach are predicted

from relationships between observed levels of child and adult mortality and their associated age-specific mortality patterns across all ages. For MODMATCH, the regression equations for these relationships were derived using a historical series of reliable life tables.²⁸ For the current analysis, selected input values of ${}_5q_0$ and ${}_{45}q_{15}$ for each state were chosen from the three empirical data sources (CRS, SRS and NFHS). We generally selected the higher value for each variable as available from the CRS and SRS, with a few exceptions. From each MODMATCH derived life table, the LE at birth and risk of premature adult mortality were derived for comparative analysis. The same input parameters were also used in the two-dimensional Logarithmic Quadratic (Log-Quad) Model Life Table System developed by Wilmoth *et al* for a sensitivity analysis of MODMATCH outputs.³⁰ Given the little difference between these estimates, we used the MODMATCH results for further analysis. The estimated MODMATCH age-specific mortality risks were applied to respective age-specific populations from the NCP projections, to derive MODMATCH estimated numbers of deaths at each age.

Data quality evaluation

The quality of mortality data from each source was evaluated by comparing the trajectory and slope of graphs of age-specific risks of dying between 30 and 70 years by sex, across all sources. The natural log of mortality rates from ages 30 to 70 were plotted to evaluate if mortality rates from each source followed the Gompertz law of linear increase in log-mortality.^{31 32} Model plausibility was also assessed by confirming that the age-specific trajectories of MODMATCH derived mortality rates for each state reflected their input values of ${}_{45}q_{15}$ from the CRS, SRS or NFHS, respectively. The MODMATCH graphs were also compared with similar graphs by sex for each state that were derived from the Log-Quad Model, to evaluate model consistency.

Premature adult mortality burden

The absolute numbers of observed or estimated deaths between 30 and 70 years in each state by sex from each source in 2017 were compared, as cross-sectional measures of premature adult mortality burden. In addition, the ${}_{40}q_{30}$ risks from each source were applied to a cohort of individuals aged 30 years in 2017, to estimate the potential numbers of deaths that would occur in this cohort over a 40-year period till 2057. The cohort comprised one-fifth of the state population aged 30–34 years in 2017 from the NCP projections, and the estimated deaths are based on the assumption that mortality levels will remain constant at the 2017 level over the ensuing four decades.

Patients and public involvement

Patients or the public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

RESULTS

Evaluation of model inputs

The plausibility of risks of ${}_5q_0$ (child mortality) and ${}_{45}q_{15}$ (adult mortality) from different sources were evaluated for each location, to select the most viable inputs for our model life table analysis. The general approach was to select the highest plausible value of each input parameter from the three empirical sources, to reflect the potential maximal levels of mortality for the study reference time period. Table 2 shows that the CRS risks of ${}_5q_0$ are implausibly low, due to known under-registration of infant deaths in all states.^{12 13} The NFHS records higher levels of ${}_5q_0$ than the SRS in most states, except for males in Kerala, and for females in Andhra Pradesh, Gujarat, Haryana, Kerala, and Rajasthan. However, the NFHS rates relate to a period of 3 years prior to the survey (2013–2016), in addition to being subject to sampling error and recall bias. In comparison, the SRS has a much larger sample size, and is a continuous recording system with half yearly check surveys, therefore less subject to recall bias.^{33 34} For these reasons, the SRS ${}_5q_0$ values were chosen as inputs for MODMATCH in all states except Bihar, Jharkhand and Uttarakhand, for which the NFHS values were chosen, since the SRS values are implausibly low, as observed from other studies.³⁵ The modelled GBD and NBE ${}_5q_0$ risks demonstrate a relative difference $>\pm 5\%$ when compared with the SRS values in most states. Since the NBE and GBD mortality risks are essentially outputs derived from other modelling processes, they were not considered as inputs for our model life table analysis.

The CRS ${}_{45}q_{15}$ values are the highest of the three empirical sources for six states in males, and for two states in females, and were chosen as the most plausible inputs for MODMATCH for these states. For most of the other states, the SRS ${}_{45}q_{15}$ estimates were considered more plausible than those from the NFHS, due to its larger sample size, and for the compatibility of SRS measures with the

time period of other data sources, for comparison. For Bihar and Jharkhand, the NFHS ${}_{45}q_{15}$ measures were deemed more plausible, given the recognised under-reporting of SRS deaths in these states.³⁵ Similar to the comparisons of under-5 mortality risks, the GBD and NBE modelled values of ${}_{45}q_{15}$ differed from SRS values by $>\pm 5\%$ in three-fourths of all states,

Model evaluation

Figure 1 shows that MODMATCH derived mortality risks show a smooth exponential rise in mortality risks after age 40 years for Tamil Nadu, Punjab and Bihar, and correlate with respective ${}_{45}q_{15}$ inputs from the CRS, SRS and NFHS respectively. This is less obvious for Bihar, potentially due to the relative instability of NFHS mortality risk trends by age, as a result of low sample size. These graphs generally support the plausibility of MODMATCH derived age-specific trends as compared with the SRS, GBD and NBE trend lines, which do not appear to conform to the Gompertz law of smooth exponential increase in age-specific mortality (linear increase on the log-scale). Similar aberrations were observed in the graphs for all states (see online supplemental appendix 3). Using the same input parameters, the Log-Quad model outputs of LE at birth and of age-specific mortality trends by sex for each state were very closely correlated to those from MODMATCH (see online supplemental appendix 4).

Comparison of population exposures

Table 3 shows the estimated total populations and deaths as well as life expectancies at birth by sex for each location from the different sources. National-level and state-level differences in population exposures between the NBE derivations from WPP estimates, the GBD population estimates, and the NCP projections (used for the MODMATCH estimates) were evaluated to assess the likely impact of these denominators on the magnitude of deaths from each source. At the national level, the GBD population estimates are relatively higher than those from the NCP projections by 5.2%. Across states, these relative differences for males range from 1% in Kerala to 13% in Odisha, and in females from 2% in Kerala to 8% in Telangana. Similarly, the NBE populations also exceed the NCP projections by 2% at national level, with concomitantly lower orders of state level differentials, except for Jammu and Kashmir (12% for males, and 8% for females). These population differentials, although mostly in single digit percentages, actually translate into large population counts in most states, and strongly influence the comparisons of estimated deaths across data sources for each state.

Overall mortality

CRS mortality reports were excluded from the comparisons of overall mortality shown in Table 3, in view of the known under-registration of deaths in several states. Using the highest plausible empirically observed levels of ${}_5q_0$ and ${}_{45}q_{15}$ as model inputs, the MODMATCH LEs at

Table 2 Estimated risks of dying before age 5 years and between ages 15 and 60 years from various sources for India and states during 2014–2017

Location	Risk of dying between 15 and 60 years																			
	Risk of dying between birth and 5 years							Risk of dying between 15 and 60 years												
	Male			Female				Male			Female									
	CRS	SRS	NFHS	NBE	GBD	CRS	SRS	NFHS	NBE	GBD	CRS	SRS	NFHS	NBE	GBD	CRS	SRS	NFHS	NBE	GBD
India	11	39*	51	43	39	9	42	47	45	43	182	203	211	213	210	110	144	129	137	155
Andhra Pradesh	16	34	45	NA	29	13	29	27	NA	28	257	229	249	NA	205	153	153	134	NA	144
Assam	8	38	61	45	54	7	49	53	52	55	178	235	221	213	237	154	166	142	132	171
Bihar	5	34	61	42	47	5	40	53	43	55	107	160	225	151	151	69	137	174	132	152
Chhattisgarh	21	55	69	47	42	16	46	62	50	43	242	249	236	255	288	171	194	186	149	212
Gujarat	17	45	48	45	32	15	44	38	43	33	213	213	240	216	215	114	127	112	120	150
Haryana	4	37	35	42	32	4	45	41	44	38	196	237	216	242	227	132	130	119	132	143
Himachal Pradesh	11	31	43	NA	31	10	24	31	NA	28	165	167	220	NA	205	73	85	90	NA	96
Jammu and Kashmir	14	32	42	34	33	18	29	35	34	36	68	158	147	158	166	65	126	104	101	119
Jharkhand	9	32	51	38	35	8	38	56	42	37	138	170	190	178	168	83	154	128	146	182
Karnataka	16	31	36	33	29	13	30	29	37	27	249	212	193	241	246	132	142	96	150	181
Kerala	8	13	10	11	11	7	13	6	12	11	162	137	143	172	160	73	77	57	81	90
Madhya Pradesh	11	58	66	66	43	10	58	62	65	45	190	221	209	248	237	124	144	137	136	151
Maharashtra	11	19	32	21	24	10	21	21	22	24	164	170	183	193	190	106	121	91	127	140
Odisha	23	44	49	56	51	18	42	46	54	45	212	202	228	208	197	137	159	146	143	153
Punjab	6	24	29	26	30	6	27	37	31	34	198	241	193	208	203	134	142	118	125	143
Rajasthan	9	48	55	51	44	7	57	50	56	51	257	229	197	248	246	114	126	105	121	151
Tamil Nadu	12	23	34	21	15	11	22	24	21	15	239	201	209	206	218	127	111	117	119	143
Telangana	11	33	29	NA	24	12	34	40	NA	23	166	204	306	NA	184	124	135	148	NA	128
Uttar Pradesh	6	59	77	59	54	6	70	82	64	64	121	255	218	262	251	78	206	149	187	201
Uttarakhand	5	31	54	NA	28	5	26	49	NA	29	109	252	250	NA	237	52	155	115	NA	131
West Bengal	14	26	39	27	31	11	29	24	28	30	143	154	175	167	176	106	143	127	116	139

* Numbers in bold font indicate input values selected for MODMATCH life tables.

CRS, Civil Registration System; GBD, Global Burden of Disease; NA, not available; NBE, National Burden Estimates; NFHS, National Family Health Survey; SRS, Sample Registration System.

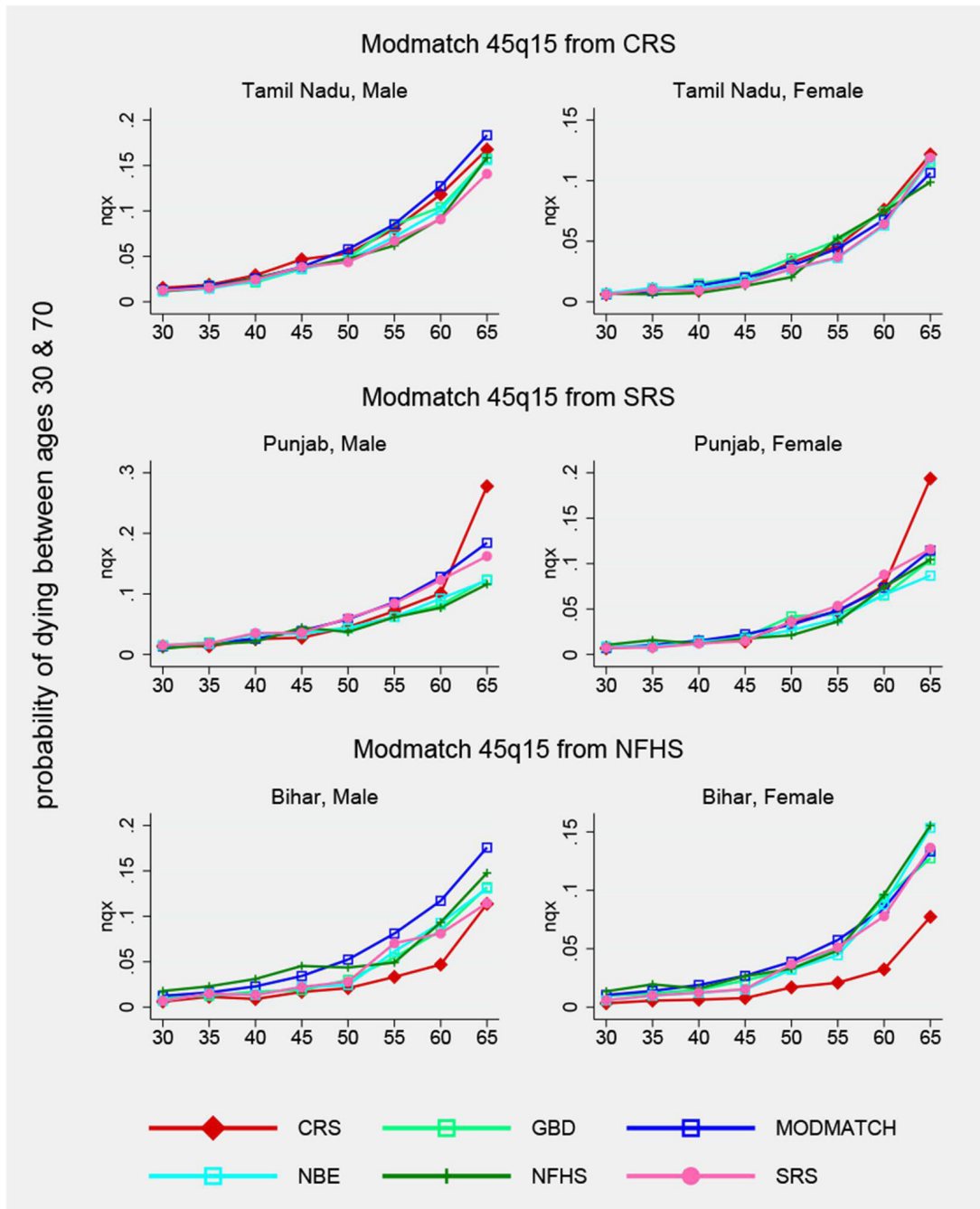


Figure 1 Mortality risk (30–70 years) from different sources for selected states, 2017. CRS, Civil Registration System; GBD, Global Burden of Disease; NBE, National Burden Estimates; NFHS, National Family Health Survey; SRS, Sample Registration System.

birth were found to be lowest out of all the sources at national level. They were also the lowest in 17 states for males, and in 13 states for females. The MODMATCH age-specific mortality predictions are based on historical empirical mortality schedules from countries with complete mortality data, derived from modelled relationships between ${}_5q_0$ – ${}_{45}q_{15}$ and overall mortality. Their plausibility is also supported by the findings from figure 1 and online supplemental appendix 3, and suggest that LEs at birth could be potentially lower than what is known from the SRS, which is the national standard source for LE estimates for India.³⁶ Even if the MODMATCH LE estimates

are not considered, there is a range of 1 year or more in LEs from the NBE, GBD and SRS for males in 13 states, and for females in 11 out of 21 states. These considerable differences in LE at birth across the various sources of mortality estimates create uncertainty as to the true mortality levels for each state in 2017.

At the national level, MODMATCH estimated the highest number of total male deaths, while the GBD estimated the highest numbers of total female deaths. At state level, there were substantial differences in estimated deaths from each source, arising from variations in age-specific death rates as well as in estimated population

Table 3 Estimated population, total deaths and life expectancy at birth by sex from various sources for India and states during 2014–2017

Location	Female																			
	Male						Female													
	NBE		GBD		MODMATCH		SRS		NBE		GBD		MODMATCH		SRS					
Pop (m)	Deaths ('000s)	LE (years)	Pop (m)	Deaths ('000s)	LE (years)	Pop (m)	Deaths ('000s)	LE (years)	Pop (m)	Deaths ('000s)	LE (years)	Pop (m)	Deaths ('000s)	LE (years)	Pop (m)	Deaths ('000s)	LE (years)			
India	694	5298	67.4	708	5230	68.0	675	5369	67.2	68.3	645	4354	70.6	672	4680	70.3	638	4275	70.5	71.0
Andhra Pradesh	NA	NA	NA	27	219	69.1	26	268	65.4	68.2	NA	NA	NA	27	213	71.8	26	195	71.3	71.4
Assam	18	114	68.4	18	134	66.0	17	130	65.9	67.2	17	97	70.5	18	122	67.7	17	101	68.7	68.9
Bihar	62	374	69.6	64	374	69.8	61	471	64.6	69.5	56	360	69.1	59	364	68.6	56	333	68.0	69.4
Chhattisgarh	15	133	63.8	15	149	62.1	14	120	64.0	63.3	14	105	68.1	15	118	68.4	14	104	67.7	67.2
Gujarat	35	250	67.8	36	248	69.1	35	278	66.3	67.4	32	182	73.6	33	225	72.7	32	197	71.2	72.2
Haryana	15	117	67.2	15	109	68.3	15	130	65.9	67.7	13	83	72.4	14	90	71.5	13	90	70.7	72.7
Himachal Pradesh	NA	NA	NA	4	34	68.4	4	31	69.5	70.9	NA	NA	NA	4	21	77.1	4	22	75.7	76.7
Jammu and Kashmir	8	41	73.5	7	40	72.0	7	43	69.8	74.2	7	26	81.7	7	33	74.6	6	33	72.7	78.2
Jharkhand	19	112	69.3	20	111	70.0	19	127	66.9	70.3	18	107	69.8	19	119	68.8	18	105	68.7	68.8
Karnataka	34	316	66.3	34	302	66.5	33	319	65.9	67.6	33	278	69.6	34	274	69.7	32	226	71.7	71.1
Kerala	17	143	72.1	17	163	71.6	17	150	71.3	72.5	18	118	78.1	19	141	77.3	18	115	77.6	77.8
Madhya Pradesh	42	382	63.4	44	324	66.4	42	339	65.0	65.5	39	299	67.9	41	307	68.7	39	264	69.0	69.2
Maharashtra	65	445	71.1	64	449	71.0	63	473	70.4	71.8	59	391	73.6	60	426	73.0	58	377	73.8	73.9
Odisha	23	197	66.6	24	202	67.2	21	199	66.4	68.4	22	154	70.1	24	170	70.1	22	163	69.7	70.6
Punjab	16	111	71.5	16	115	71.3	16	163	66.8	69.4	14	83	75.4	15	94	73.9	14	108	72.0	73.8
Rajasthan	40	323	64.8	42	301	66.1	39	323	64.2	66.5	37	221	71.3	39	247	70.8	37	237	70.0	71.6
Tamil Nadu	40	333	69.4	40	347	69.3	38	395	67.0	70.2	40	256	74.0	40	292	72.8	38	260	73.3	74.1
Telangana	NA	NA	NA	20	127	71.2	19	153	67.7	68.1	NA	NA	NA	20	118	74.3	18	122	71.9	70.5
Uttar Pradesh	116	989	64.2	122	937	65.2	115	1023	63.5	65.2	105	802	66.6	112	860	66.1	106	857	65.1	66.4
Uttarakhand	NA	NA	NA	6	5	67.0	6	55	64.0	66.5	NA	NA	NA	6	35	72.8	5	41	69.3	72.8
West Bengal	52	349	70.5	52	370	69.8	49	343	70.6	71.1	48	304	72.4	50	312	72.2	47	303	71.8	72.4

Highlighted value is the lowest LE at birth from all sources for each location.

GBD, Global Burden of Disease; LE, life expectancy; NA, not available; NBE, National Burden Estimates; SRS, Sample Registration System.

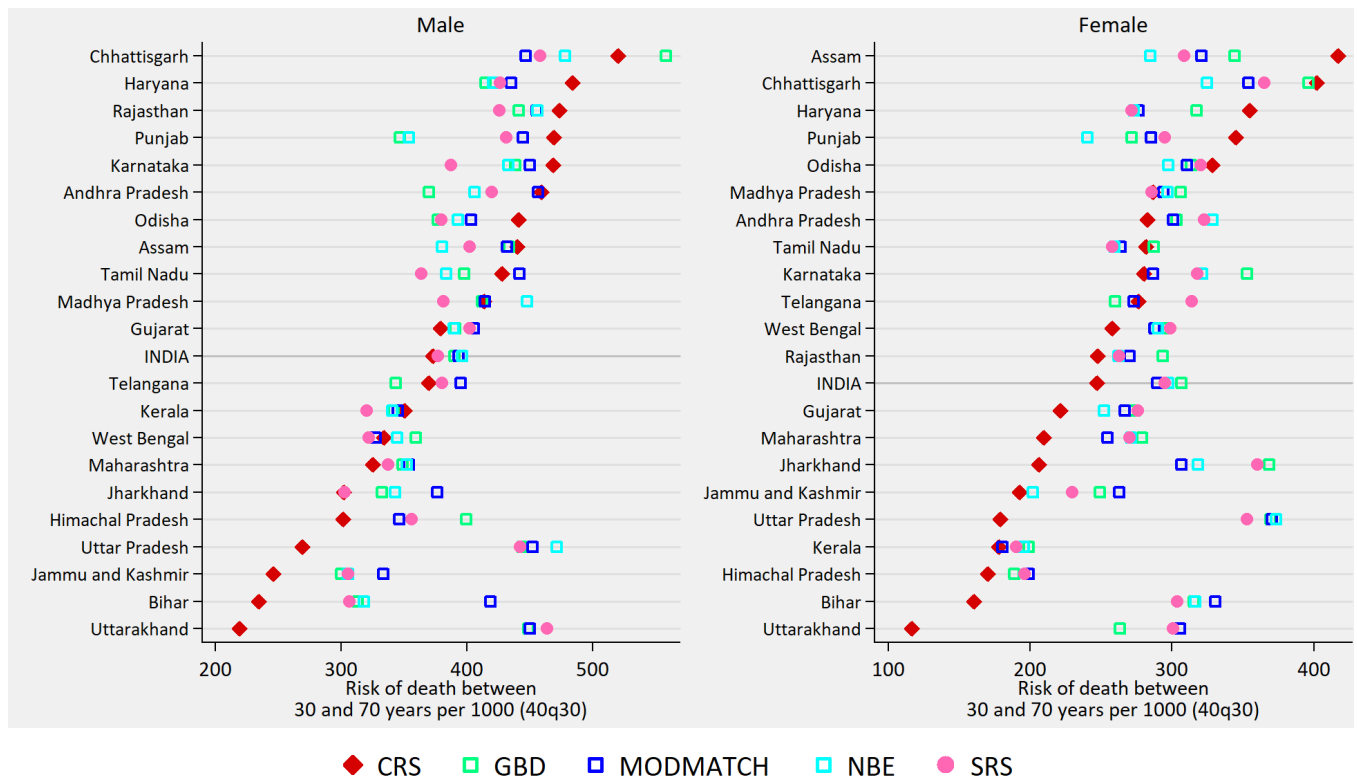


Figure 2 Estimated risk of dying between 30 and 70 years from different sources for India and states, 2017. CRS, Civil Registration System; GBD, Global Burden of Disease; NBE, National Burden Estimates; SRS, Sample Registration System.

exposures. The net effect of these variations was assessed in terms of the dispersion of estimated total deaths across sources. The dispersion was >10000 deaths in 18 out of 21 states for males, and 15 out of 21 states in females. Very high dispersions (>40000 deaths) were observed for males in six states, and for females in four states. Such variations in total estimated deaths from different sources for each state also create uncertainty about the likely true levels of overall mortality at sub national level.

Premature adult mortality risk

For interpreting the findings on premature adult mortality across different sources, a conservative approach was used in nominating the highest estimate for each state as the most likely value. **figure 2** shows considerable heterogeneity in the maximum values of $_{40}q_{30}$ mortality risks across the states. For males, these values range from 351 per 1000 in Kerala to 558 per 1000 in Chhattisgarh, and for females from 198 per 1000 in Himachal Pradesh to 409 per 1000 in Assam.

On comparing estimated risks across data sources, **figure 2** shows that despite a general perception of incomplete death registration, CRS values of the $_{40}q_{30}$ mortality risks were the highest out of all sources for males in eight states (representing 28% of the national male population), and for females in five states (13% of national female population). The MODMATCH values were the highest values for males in eight states (42%), and for females in four states (11%). The GBD modelled

$_{40}q_{30}$ estimates were the highest values for males in three states (10%), and females in seven states. (37%)

Another important comparison is between the CRS and SRS values. The observed CRS values of $_{40}q_{30}$ for males were higher than the SRS values in 13 states, and for females in 7 states, indicating the likelihood of bias in SRS data, from sampling error and/or from problems with data quality. The graphs in **figure 2** show limited convergence in risks of premature adult mortality across data sources in several states for both males and females, even if the low CRS values for some states are not considered, due to incomplete death registration.

Premature adult mortality burden

A cross-sectional perspective of the magnitude of premature adult mortality is provided in the left panel of **table 4**, in terms of the maximum estimate of deaths between 30 and 70 years in each state during 2017, colour coded to its data source. At the national level, a total of 2.6 million male and 1.8 million female deaths were expected to have occurred in this age group during 2017. Bihar, Maharashtra, Tamil Nadu, Uttar Pradesh and West Bengal account for approximately half of all these deaths (see online supplemental appendix 5A for detailed estimates).

There is no single data source that consistently estimates the maximum number of deaths in all states. Although the GBD estimates the maximum deaths for females in two-thirds of all states, and for males in seven states, these

Table 4 Estimated deaths for India and states at ages 30–70 years in 2017, and expected cohort deaths during 2017 to 2057

Location	Maximum estimates of premature adult mortality risks and deaths in 2017				Expected no of deaths in a cohort of 30-year-old men and women over the period 2017–2057			
	Men		Women		Men		Women	
	40q30 per 1000	Deaths ('000s)	40q30 per 1000	Deaths ('000s)	Cohort* ('000s)	Deaths ('000s)	Cohort* ('000s)	Deaths ('000s)
India	396	2609	306	1802	10340	3859–4098	10045	2911–3078†
Andhra	459	135	323	85	434	160–199	448	126–135
Assam	440	66	417	50	269	102–118	277	79–116
Bihar	419	192	330	138	763	239–320†	762	241–252†
Chhattisgarh	558	79	402	53	212	95–119	209	68–84
Gujarat	406	130	276	83	572	217–232	515	130–141†
Haryana	484	63	355	39	243	101–117	217	59–77
Himachal Pradesh	400	16	199	7	59	18–24	60	10 -- 12
Jammu and Kashmir	334	17	263	12	110	27–37	98	19–26
Jharkhand	376	55	369	56	268	81–101	263	81–97†
Karnataka	469	166	353	116	567	246–266	556	156–196
Kerala	351	74	199	43	242	82–85	276	49–55
Madhya Pradesh	448	164	306	99	614	253–275	575	165–176
Maharashtra	354	227	279	161	1071	348–379	979	249–273†
Odisha	441	96	329	70	327	123–144	355	106–117
Punjab	469	71	345	46	257	91–121	240	58–83
Rajasthan	474	152	293	85	563	248–267	538	133–158
Tamil Nadu	442	197	287	126	624	248–276	652	169–187
Telangana	395	68	314	47	318	109–125	327	85–90
Uttar Pradesh	471	468	373	322	1540	684–726†	1465	541–547†
Uttarakhand	464	24	306	14	83	37–38†	85	22–26†
West Bengal	359	182	299	131	798	262–287	785	226–231†

CRS; GBD; NBE; MODMATCH; SRS.

*Based on NCP population estimates.²⁴

†CRS values not included due to low completeness of death registration.

CRS, Civil Registration System; GBD, Global Burden of Disease; NBE, National Burden Estimates; NCP, National Commission on Population; SRS, Sample Registration System.

are largely a result of the higher GBD population exposures. Similarly, NBE derived deaths are the highest at national level and five other states for males, also driven by higher population bases. On the other hand, the directly observed CRS deaths are the maximum for males in six states and for females in three states, and MODMATCH death estimates are maximum for males and females in three states each. The CRS and MODMATCH deaths are derived using the NCP predicted populations, which are the lowest of all three sources of population data.

The potential future burden from premature adult mortality till 2057 was calculated by applying the ${}_{40}q_{30}$ risks to a cohort of individuals from each state who were aged 30 years in 2017. On eliminating the differences from varying baseline population exposures, it was found that the maximum cohort estimates were mostly based on the MODMATCH and CRS risks, particularly for males (see online supplemental appendix 5B for detailed estimates). While these estimates of expected deaths offer some value in understanding the broad targets for mortality reductions in each state, the broad ranges of these estimates (approximately 20 000 deaths or more) across different sources for some states are a matter of concern, and will hamper the exact quantification of mortality reductions achieved, in the future, over the 40-year period till 2057.

DISCUSSION

The principal findings from this analysis are the measures of premature adult mortality between ages 30 and 70 years for India and states from various data sources and methods around the commencement of the UNSDG era in 2017, as shown in figure 2 and table 4. Based on the assumption that the highest values of estimated mortality by sex for each state from any source represents this baseline measure for UNSDGs, a total of 2.6 million male and 1.8 million female deaths were expected to have occurred in India from this age group, during 2017. Bihar, Maharashtra, Tamil Nadu, Uttar Pradesh and West Bengal account for approximately half of all these deaths. In terms of mortality risks, there was considerable heterogeneity across states, with estimated values for males ranging from 351 per 1000 in Kerala to 558 per 1000 in Chhattisgarh, and for females from 198 per 1000 in Himachal Pradesh to 409 per 1000 in Assam.

Our findings show that at the national level, premature adult mortality accounts for nearly half of all deaths in males, and about a third of all female deaths, and these proportions are consistent across all the sources and for all states (details in online supplemental appendix 5A). These large proportions actually translate into an annual burden of considerable numbers of deaths in each state each year, as shown in table 4. These deaths are likely to be mostly from NCDs, injuries, tuberculosis and infectious hepatitis, which can be addressed through strengthening of ongoing programmes for disease and injury prevention and control.^{37–40} Also, our proposed estimates of deaths at these ages could be used as an outer bound

of deaths to which cause-specific proportional mortality distributions from reliable epidemiological data sources could be fitted, to estimate the population level mortality burden from specific diseases and conditions.

Although the results have mostly focused on the comparisons and variations of estimates from different sources, our overall aim was to estimate the potentially true magnitude of premature adult mortality around 2016–2017. This is of critical importance, since these are required as baseline levels for monitoring the effectiveness of strategies to address health-related UNSDGs that target this specific age category. Since each data source has limitations which vary in scope and effect across locations, it is not possible to nominate any single source as the optimal source for all states in 2017. Hence, we have chosen to report the maximum mortality estimate at these ages for each state, under the principle that health policies should be based on ‘worst case’ scenarios, for pragmatic health sector priority setting and resource allocation.

In this regard, we believe that the incorporation of mortality data from the CRS in 2017 in our estimation process has had an important influence on deriving our maximal mortality estimates. This is evidenced from the direct use of CRS derived values of ${}_{45}q_{15}$ risks as inputs for our model life tables for some states, as well as in applying the observed CRS ${}_{40}q_{30}$ risks in certain states to estimate the magnitude of premature adult deaths. Data from the CRS were not used for mortality estimation in the GBD and NBE analyses.

Three indicators were used to evaluate premature adult mortality, namely the risk of dying between the 30 and 70 years, the cross-sectional annual number of deaths at a baseline point in time, and a prediction of potential deaths in a cohort over a period of 40 years. Evaluating the risks of dying helps understand the underlying factors driving mortality, while cross-sectional estimates of deaths guide resource allocation for health services and clinical management to prevent death in the diseased, and predicted deaths serve as targets for disease control and mortality reduction strategies. Planning for disease prevention and control to reduce premature adult mortality is critical, since in addition to enabling individuals to fulfil their personal life goals, mortality prevention at these ages has important positive ramifications for their families, as well as society in general.

It is anticipated that the heterogeneity in mortality patterns is likely to be associated with varying disease profiles and causes of death patterns across the states. Hence, there would be a need for customised health programmes for each state, and related indicators and targets to monitor and evaluate their impact. While the GBD and NBE studies have published state level epidemiological profiles, they differ in their respective cause-specific mortality patterns for individual states.^{17 19} Moreover, our analysis proves that even at a gross level, the GBD and NBE total estimated numbers of deaths for each state are not the same, and these are influenced

by differing patterns of modelled age-specific mortality risks, and differing population exposures, resulting in disparate premature adult mortality risks (see online supplemental appendix 6) These myriad explanations for differences in the GBD and NBE deaths for each state, and the absence of specific evidence to assess and justify the reliability of one source over the other, limits the utility of either of these sources, for establishing baseline levels of premature adult mortality to monitor progress towards the UNSDGs.

We have highlighted the lowest LEs at birth and the highest estimates of premature adult mortality by sex from all sources, as the most appropriate values to describe mortality levels for each state. The mortality measures from the CRS are known to be affected by bias due to under-registration, but our analyses did not include any measurement of registration completeness, and related adjustments. Therefore we chose not to use life expectancies from the CRS for this comparison in Table 3. The MODMATCH LEs at birth for most states were observed to be lower than what was previously understood from the SRS and GBD reports.^{19 36} These findings have implications for the computation and comparative interpretation of the Human Development Index across the states of India, since LE at birth is a key component of this index.

Our evaluation of plausibility of patterns of age-specific mortality risks (figure 2) show that the GBD, NBE and SRS age-specific mortality curves do not adhere to the Gompertz law that describes an exponential rise in mortality beyond 40 years, and such violations are generally considered to indicate problems with underlying data.³² In contrast, the MODMATCH outputs of age-specific risks are compliant with the Gompertz law. Further, the results from MODMATCH were validated using the Log Quad Model Life Table System. We chose to use the MODMATCH results for our comparative analysis, since the details on state-specific inputs used in MODMATCH, and references to the software tool and relevant documentation are available in the public domain, to replicate and verify our results for each state. Nevertheless, although the MODMATCH ${}_{40}q_{30}$ risks are generally higher than the SRS values for most states (see figure 2), they are still below the same risks derived from observed deaths in the CRS, for both males and females in some states. These MODMATCH predicted mortality patterns are based on the expectation that current mortality in the study population reflects international historical trends, and this is not the case for these states with higher ${}_{40}q_{30}$ risks from empirical CRS data. Hence, the MODMATCH too potentially underestimates premature adult mortality, and the degree of such under-estimation cannot be ascertained for several states with incomplete CRS death registration, particularly for females.

Another limitation of this analysis was our inability to establish the potentially minimum value of ${}_{40}q_{30}$ risk in Indian populations, due to all the uncertainty underlying

our estimates. A reliable minimum risk could have been used as a counterfactual level, to estimate the potentially avoidable burden from premature adult mortality in each state.⁴¹ For this estimation, the counterfactual minimum risk would be applied to each cohort of 30 years in 2017, to yield the minimum expected deaths for each state over the 40-year period till 2057. Subtracting these minimum expected deaths from the cohort mortality estimates presented in table 4 would yield the potentially avoidable burden from premature adult mortality in each state, if the state were to experience the same population health status and health system attributes as the population with the counterfactual risk. While it is possible to use an estimate of ${}_{40}q_{30}$ from a reliable international source as the counterfactual risk, we did not choose to do so, since it may not be epidemiologically coherent with Indian health experience.

From an overall perspective therefore, none of the available data sources or analytical methods appears to serve as the single appropriate source for understanding the true levels of premature adult mortality in India. Our approach to compare and report the maximum estimate for each state as the likely value is only borne out of expediency, to focus attention of health policy analysts and agencies on the potential magnitude of this component of disease burden around 2017, as baseline measures for the UNSDGs. In highlighting the gaps in empirical data, along with the variations in estimates across sources, we hope to draw the attention of public health bureaucrats to the urgent action required to strengthen the CRS as the optimal source for subnational mortality statistics in India. As described in detail elsewhere, the CRS is based on a sound legal framework and administrative structure, with adequate infrastructure and operations for basic registration of deaths by age and sex across the country, which have led to gradual improvements in data quality till date.^{12 13 42} The CRS is also the optimal source since its statistical compilations report data for all the smaller states and territories of India. Further interventions are required to improve completeness of death registration in some locations, and attribution of causes of death more generally, based on an incremental sampling approach supported by adequate capacity building, over the next decade.^{13 43 44} Also, CRS statistical reports should provide district level data with more detailed age-groups, to directly estimate abridged life tables and other age-related mortality measures. There is also a need to coordinate population projection exercises and methods across different groups, so that there is a unified set of consensus-based population exposures for evaluating mortality risks. The methods and findings from these analyses should be taken into account when using data from the CRS to evaluate and monitor the impact of the COVID 19 pandemic in India during 2020-2021, and into the future.⁴⁵ This analytical approach would also be relevant for other countries facing the need to reconcile local mortality data with modelled mortality estimates from international sources.

Twitter Aashish Gupta @aashishg_

Contributors CR conceptualised the research, led the analysis and drafted the initial version of the manuscript. AG collaborated on the analysis and design of figures and tables and contributed to writing the manuscript. MG and AKY provided inputs for the review of data quality, interpretation of results and documentation of implications of the study findings in the discussion. All authors contributed to the preparation of the final version of the manuscript.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient consent for publication Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Estimates of population and deaths by age and sex in 2016 for India and states from the Indian State-level Disease Burden Initiative were obtained through direct request to the authors of reference 19 in the manuscript. All other data are available in public open access repositories as listed in relevant references.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iD

Chalapati Rao <http://orcid.org/0000-0002-9554-0581>

REFERENCES

- 1 Cha S. The impact of the worldwide Millennium Development Goals campaign on maternal and under-five child mortality reduction: 'Where did the worldwide campaign work most effectively?'. *Glob Health Action* 2017;10:1267961.
- 2 UN Inter-agency Group for Child Mortality Estimation (UN-IGME). *Levels and trends in child mortality: report 2020*. New York: UNICEF, 2020. <https://www.unicef.org/media/79371/file/UN-IGME-child-mortality-report-2020.pdf.pdf>
- 3 Global Health Observatory (GHO) Data. Adult mortality rate 2000 - 2016: Situation and trends: World Health Organization, 2020. Available: https://www.who.int/gho/mortality_burden_disease/mortality_adult/situation_trends_text/en/
- 4 Murray CJL, Yang GH, Qiao X. Patterns and Causes. In: Feachem RGA, Kjellstrom T, Murray CJL, et al, eds. *The health of adults in the developing world*. New York: Published for the World Bank : Oxford University Press, 1992.
- 5 United Nations Statistics Division. SDG indicators: United Nations global SDG database: UN Department of Economic and Social Affairs, 2020. Available: <https://unstats.un.org/sdgs/indicators/database/>
- 6 Mills SL, Abouzahr C, Kim JH. *Civil registration and vital statistics (CRVS) for monitoring the sustainable development goals*. Washington D.C.: World Bank Group, 2017. <http://documents.worldbank.org/curated/en/979321495190619598/Civil-registration-and-vital-statistics-CRVS-for-monitoring-the-Sustainable-development-goals-SDGS>
- 7 Rao C. Elements of a strategic approach for strengthening national mortality statistics programmes. *BMJ Glob Health* 2019;4:e001810.
- 8 Rao C, Kelly M. Empiricism in non-communicable disease mortality measurement for the Asia-Pacific: lost in translation. *BMJ Glob Health* 2020;5:e003626.
- 9 Verguet S, Jamison DT. Performance in rate of decline of adult mortality in the OECD, 1970-2010. *Health Policy* 2013;109:137-42.
- 10 Calazans JA, Queiroz BL. The adult mortality profile by cause of death in 10 Latin American countries (2000-2016). *Rev Panam Salud Publica* 2020;44:1.
- 11 Pillay-van Wyk V, Msemburi W, Laubscher R, et al. Mortality trends and differentials in South Africa from 1997 to 2012: second National burden of disease study. *Lancet Glob Health* 2016;4:e642-53.
- 12 Gupta M, Rao C, Lakshmi PVM, et al. Estimating mortality using data from civil registration: a cross-sectional study in India. *Bull World Health Organ* 2016;94:10-21.
- 13 Rao C, Gupta M. The civil registration system is a potentially viable data source for reliable subnational mortality measurement in India. *BMJ Glob Health* 2020;5.
- 14 Pandey A, Roy N, Bhawsar RD. Health information system in India: issues of data availability and quality. *Demogr India* 2010;39:239-45.
- 15 Dubey M, Mohanty SK. Age and sex patterns of premature mortality in India. *BMJ Open* 2014;4:e005386.
- 16 Ram U, Jha P, Gerland P, et al. Age-specific and sex-specific adult mortality risk in India in 2014: analysis of 0-27 million nationally surveyed deaths and demographic estimates from 597 districts. *Lancet Glob Health* 2015;3:e767-75.
- 17 Menon GR, Singh L, Sharma P, et al. National burden estimates of healthy life lost in India, 2017: an analysis using direct mortality data and indirect disability data. *Lancet Glob Health* 2019;7:e1675-84.
- 18 United Nations Population Division. *World population prospects: the 2017 revision*. New York: United Nations Department of Economic and Social Affairs, 2017. <https://www.un.org/development/desa/publications/world-population-prospects-the-2017-revision.html>
- 19 India State-Level Disease Burden Initiative Collaborators. Supplement to: nations within a nation: variations in epidemiological transition across the states of India, 1990-2016 in the global burden of disease study. *Lancet* 2017;390:1-141.
- 20 Murray CJL, Callender C, Kulikoff XR, et al. Population and fertility by age and sex for 195 countries and territories, 1950-2017: a systematic analysis for the global burden of disease study 2017. *Lancet* 2018;392:1995-2051.
- 21 World Health Organization. Noncommunicable diseases global monitoring framework: indicator definitions and specifications, 2014. Available: https://www.who.int/nmh/ncd-tools/indicators/GMF_Indicator_Definitions_FinalNOV2014.pdf
- 22 Office of the Registrar General and Census Commissioner of India. *Sample registration system: statistical report 2017*. New Delhi: Office of the Registrar General of India, Ministry of Home Affairs, 2019. http://censusindia.gov.in/vital_statistics/SRS_Reports_2017.html
- 23 International Institute for Population Sciences (IIPS) and ICF. *National family health survey (NFHS-4) 2015-2016*. Mumbai: International Institute for Population Sciences, 2017. <http://rchiips.org/nfhs/NFHS-4Reports/India.pdf>
- 24 National Commission on Population. *Population projections for India and States 2011 - 2036: Report of the technical group on population projections*. New Delhi: Ministry of Health and Family Welfare, Government of India, 2019. https://nhm.gov.in/New_Updates_2018/Report_Population_Projection_2019.pdf
- 25 Dandona L, Dandona R, Kumar GA, et al. Nations within a nation: variations in epidemiological transition across the states of India, 1990-2016 in the global burden of disease study. *Lancet* 2017;390:2437-60.
- 26 Taylor R, Morell S. *Life table and probability of dying calculation tool*. Sydney, Australia: School of Public Health and Community Medicine, University of New South Wales, 2015. <https://sphcm.med.unsw.edu.au/node/303112492>
- 27 Andreev EM, Shkolnikov VM. Spreadsheet for calculation of confidence limits for any life table or healthy-life table quantity. MPIDR Technical Reports 2010 - 005 [Internet], 2010. Available: <https://www.demogr.mpg.de/papers/technicalreports/tr-2010-005.pdf>
- 28 Murray CJL, Ferguson BD, Lopez AD, et al. Modified logit life table system: principles, empirical validation, and application. *Population Studies* 2003;57:165-82.
- 29 World Health Organization. MODMATCH (version 1.5): modified logit life table system, 2012. Available: https://www.who.int/healthinfo/global_burden_disease/tools_software/en/
- 30 Wilmoth J, Zureick S, Canudas-Romo V, et al. A flexible two-dimensional mortality model for use in indirect estimation. *Popul Stud* 2012;66:1-28.
- 31 Benjamin Gompertz. X. On the nature of the function expressive of the law of human mortality and on a new mode of determining the value of life contingencies 1825;115:513-83.
- 32 Heuveline P, Clark SJ. Model schedules of mortality. In: Rogers RG, Crimmins EM, eds. *International Handbook of adult mortality*. Dordrecht: Springer, 2011: 511-32.
- 33 Office of the Registrar General and Census Commissioner of India. *Sample registration system*. New Delhi: Vital Statistics Division,

- Office of the Registrar General of India, 2020. http://censusindia.gov.in/Vital_Statistics/SRS/Sample_Registration_System.html
- 34 Mahapatra P. The Sample Registration System (SRS) in India: An Overview, as of 2017. Workshop on National CRVS Verbal Autopsy Sampling Strategies [Internet]. 2017, 2017. Available: https://www.researchgate.net/publication/326098440_The_Sample_Registration_System_SRS_in_India_An_Overview_as_of_2017
- 35 Yadav AK, Ram F. There Is a Glaring Gender Bias in Death Registrations in India. *Economic and Political Weekly*. 2019;54(51):Available from. Available: <https://www.epw.in/engage/article/there-glaring-gender-bias-death-registrations>
- 36 Office of the Registrar General and Census Commissioner of India. *Sample registration system: abridged life tables 2013-2017*. New Delhi, 2019. https://censusindia.gov.in/Vital_Statistics/SRS_Life_Table/SRS%20based%20Abridged%20Life%20Tables%202013-17.pdf
- 37 National Health Mission. Communicable disease control programme. Ministry of health and family welfare, government of India, 2020. Available: <https://nhm.gov.in/index1.php?lang=1&level=1&sublinkid=1056&lid=616>
- 38 National Health Mission. Non communicable disease control programmes. Ministry of health and family welfare, government of India, 2020. Available: <https://nhm.gov.in/index1.php?lang=1&level=1&sublinkid=1041&lid=614>
- 39 Mohan P, Mohan SB, Dutta M. Communicable or noncommunicable diseases? Building strong primary health care systems to address double burden of disease in India. *J Family Med Prim Care*
- 40 Trauma and Burns Division Program Management Unit. National programme for Prevention & Management of Trauma & Burn Injuries (NPPMT&BI). Ministry of Health and Family Welfare, Government of India, 2020. Available: <https://main.mohfw.gov.in/basicpage-6>
- 41 Gavurova B, Vagasova T. Assessment of Avoidable Mortality Concepts in the European Union Countries, Their Benefits and Limitations. *Advances in Health Management* [Internet]., 2017. Available: <https://www.intechopen.com/books/advances-in-health-management/assessment-of-avoidable-mortality-concepts-in-the-european-union-countries-their-benefits-and-limita>
- 42 Mahapatra P, Chalapati Rao PV. Cause of death reporting systems in India: a performance analysis. *Natl Med J India* 2001;14:154–62.
- 43 Begg S, Rao C, Lopez AD. Design options for sample-based mortality surveillance. *Int J Epidemiol* 2005;34:1080–7.
- 44 Rao C, Usman Y, Kelly M, *et al*. Building capacity for mortality statistics programs: perspectives from the Indonesian experience. *J Epidemiol Glob Health* 2019;9:98–102.
- 45 Rao C. Medical certification of cause of death for COVID-19. *Bull World Health Organ* 2020;98:298-A.