BMJ Open Cardiovascular mortality of 40–69-yearolds in Sri Lanka from 1980 to 2010: a birth cohort analysis by age and sex

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ABSTRACT

Objectives To compare cardiovascular mortality

(ischaemic heart disease (IHD), hypertensive disease (HTN) and cerebrovascular disease (CeVD)) of 40-69-year-old Sri Lankans from 1980 to 2010 by age, birth cohort and sex. Design A comparative retrospective study. Setting Sri Lanka.

Participants 40-69-year-old Sri Lankans from 1980 to 2010.

Primary and secondary outcome

measures Cardiovascular deaths due to IHD, HTN and CeVD.

Results Mortality due to IHD increased with age but decreased with birth cohorts with time (range 3.7-390 per 100 000 population); there was a spike in the IHD mortality rates in both age groups and birth cohorts in 2000. Deaths due to HTN markedly increased after 55 years; however, mortality decreased in the younger cohorts (range

2.8-204.81 per 100 000 population). CeVD mortality linearly increased with age (range 3.3-153.3 per 100 000 population): birth cohorts of 1926-1930 and 1931-1935 had a spike in mortality among the 60-64 and 65-69 age groups, respectively. Changes were seen among both males and females; mortality rates were higher in males than in females.

Conclusions All cardiovascular mortality rates increased with age and are higher in males than in females. Agespecific cardiovascular mortality rates were lower in the younger birth cohorts as compared with the older birth cohorts. The increase in cardiovascular deaths in Sri Lanka is due to the ageing population.

INTRODUCTION

Cardiovascular diseases (CVDs) are the leading cause of death globally. They comprise a group of diseases of the heart and blood vessels including coronary heart disease (CHD), cerebrovascular disease (CeVD), peripheral artery disease, rheumatic heart disease, congenital heart disease, deep vein thrombosis and pulmonary embolism.

In 2019, 17.9 million deaths due to CVDs, accounting for 32% of all global deaths, were estimated: 85% were due to heart attack and stroke. More than 75% of all CVD deaths occur in low- and middle-income countries;

STRENGTHS AND LIMITATIONS OF THIS STUDY

- \Rightarrow Birth cohort analysis of mortality due to cardiovascular diseases by age and sex shows trends in mortality and allows comparisons over time.
- \Rightarrow Data used for the analysis have been published by the Registrar General's Department of Sri Lanka.
- \Rightarrow Inability to determine the relationship between mortality and economic indicators due to lack of data is a limitation of the study.

Protected by copyright, including for uses related CVD deaths comprised 38% of premature deaths (under the age of 70) due to noncommunicable diseases.¹

to Hypertension, a blood pressure above 140/90, is a leading risk factor for cardiotext vascular disease with a heavy public health burden worldwide.² Poorly controlled or uncontrolled blood pressure increases the data risk of hypertensive disease (HTN), giving a uncontrolled blood pressure increases the rise to microscopic and macroscopic cardiac remodelling and functional alterations.³ It is estimated that 1.28 billion adults aged 30-79 ≥ years worldwide have hypertension with twothirds living in low- and middle-income coun-tries.⁴ 46% of hypertensives are unaware they have the condition, and less than 42% are diagnosed and treated; only 21% of adults with hypertension have it controlled.⁴

simi The number of CVD deaths has been increasing over time; in 2000, around 14 million died of CVDs globally, while in 2019, close to 18 million died.⁵ CVD deaths are estimated to be >23 million by 2030.⁶ The decline in incidence and mortality rates for **g** developed countries was significantly higher **8** than those for developing countries from 1990 to 2017 (p<0.05); developing nations had a less steep decline.

Risk factors for cardiovascular diseases include age, sex and other modifiable lifestyle risk factors.⁷ The rising death toll is largely due to a growing and ageing global population. Death rates have been declining due to the implementation of preventive

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programmes; large declines in smoking; improvements in screening, diagnosis and monitoring; and advances in medical treatments, public health initiatives, emergency care and surgical procedures have all helped to reduce the impact of cardiovascular diseases on people's lives.⁷ The large disparities that still exist can be further reduced.

South Asia has a disproportionately high burden of cardiovascular disease, with higher rates of CVD incidence, mortality and risk factor prevalences than many other regions.⁸ CVDs account for 3.9 million deaths in the WHO South-East Asia Region every year, comprising a quarter of all deaths from non-communicable diseases, with most of them being preventable.⁹ Even though South Asians comprise only 25% of the world's population, they account for more than 50% of the world's cardiovascular deaths.8

Sri Lanka, a country having one of the fastest ageing populations in Asia, is in the midst of an epidemiological and demographic transition. In 2022, WHO estimated that over 80% of the mortality in Sri Lanka is due to major non-communicable diseases, with CVDs contributing to over 34% of deaths, impacting both life expectancy and quality of life. The mortality from CVDs in Sri Lanka is estimated to be 524 deaths per 100000, which is higher than that observed in many high-income countries.¹⁰ Coronary artery disease is the leading cause of death in Sri Lanka, while stroke is the third highest cause of death.^{11 12}

Childhood risk factors such as obesity, exposure to indoor and outdoor tobacco smoke, dyslipidaemia and diabetes impact cardiovascular disease in adulthood¹³ and are now being targeted for prevention of cardiovascular disease.

Birth cohort analyses have been used to determine the causal relationship between potential risk factors during the prenatal and postnatal periods and the health status of the newborn up to childhood. Birth cohort analyses allow description of associations between early exposures and subsequent outcomes^{14 15}; in addition, they are able to identify the risk and environmental exposure factors shared by a given generation and are used to investigate disease trends and test a wide range of hypotheses.^{14–17} Birth cohort analyses of CVDs have not been conducted in Sri Lanka. The aim of this study was to find out whether there are variations in age-specific and sex-specific mortality rates of selected CVDs (ischaemic heart disease, hypertensive disease and cerebrovascular disease) among 40-69-year-old Sri Lankans from 1980 to 2010 by birth cohort.

METHODS

Study design and data sources

This comparative retrospective study was conducted from August 2022 to January 2024 using secondary data available in the public domain. Mortality data of Sri Lanka were extracted from the WHO mortality database from 1980 through 2010 (country code 3365).¹⁸ Mortality data are reported annually to the WHO from the civil registration

system of the country (Registrar General's department). Mortality data included the number of deaths for 5-year age groups and were coded as per the International Classification of Diseases (ICD) versions 9 (1979 up to 1992) and 10 (1993 to 2021). The 10th revision of the ICD was used in Sri Lanka from 1997 onwards.¹⁹ Separate ICD codes are given for different causes of deaths falling under cardiovascular diseases. The ICD codes used for deaths due to ischaemic heart disease, hypertensive disease and cerebrovascular diseases in the 9th 🕫 and 10th revisions used in this study are given in online supplemental table 1).

For each disease category, deaths of 5-year age groups Š (40-44, 45-49, 50-54, 55-59, 60-64 and 65-69) were extracted from the database for the codes and subcodes 8 given in online supplemental table 1) from 1980 to 2010.

The birth cohorts for different age groups corresponding to the selected years of data extraction are given in online supplemental table 2. For example, the age group 40-44 years in 1980 belonged to the 1936-1940 birth cohort. Likewise, the age group 65-69 years in 1980 belonged to the 1911-1915 birth cohort. For each birth cohort, the number of deaths for each group of ICD codes was summed for the age group studied. For example, for the birth cohort of 1936–1940, deaths coded as B27 (9th revision) were summed for the 40-44-year age group in 1980. đ

Mortality data from 1987 to 1995 were not available on the WHO website. Mortality data for 1990 was obtained by linear interpolation of the number of deaths separately for males and females in the different age groups considered. Mortality data by age and sex for 1995 were obtained data from the Registrar General's Department, Sri Lanka.

Population data by age group and sex were obtained **E** from the database displayed on the UN population website,²⁰ Sri Lanka being coded as 144. Population data ≥ are displayed as estimates for intercensal years as estitraining, mated by the Department of Census and Statistics of Sri Lanka and reported to the UN; during the census years, the actual figures are provided. Population data are reported for 5-year age groups as done for mortality data by sex.

Extracted mortality and population data were tabulated to yield six 5-year age groups from 40–44 to 65–69 years and seven quinquennial demographic profiles from 1980 to 2010 for the 12 birth cohorts. **Data analysis** Data were entered in Excel worksheets. Mortality rates for each of the three cardiovaccular discuss of

for each of the three cardiovascular diseases for each age category of both sexes were calculated based on the following formula:

Mortality rate for each age-sex category = (deaths / population) *100000

Using the calculated mortality rates for each age group and birth cohort, trends were determined to identify birth cohort and age effects by sex. Data are presented as tables and graphs. Mortality rates of ischaemic heart disease

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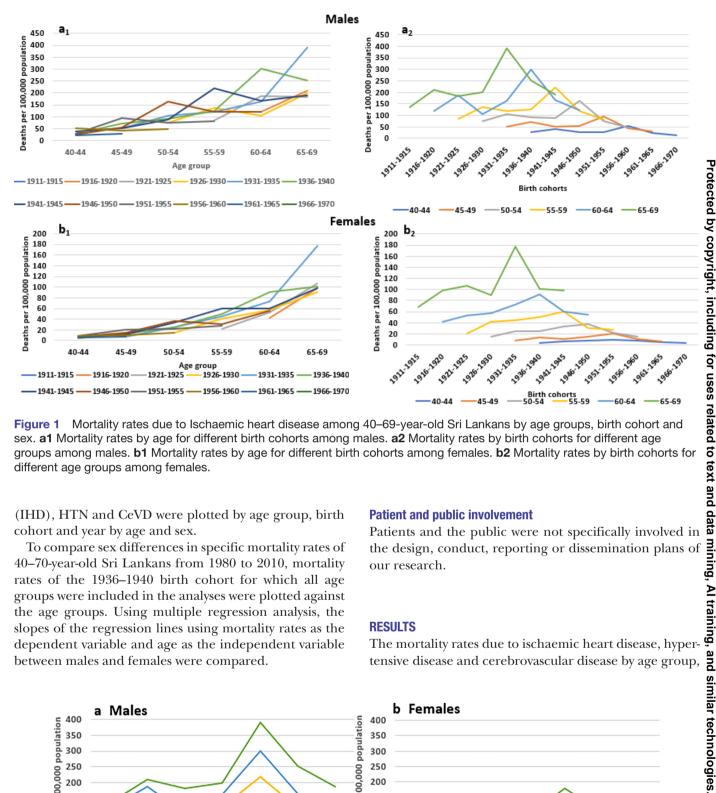


Figure 1 Mortality rates due to Ischaemic heart disease among 40–69-year-old Sri Lankans by age groups, birth cohort and sex. a1 Mortality rates by age for different birth cohorts among males. a2 Mortality rates by birth cohorts for different age groups among males. b1 Mortality rates by age for different birth cohorts among females. b2 Mortality rates by birth cohorts for different age groups among females.

(IHD), HTN and CeVD were plotted by age group, birth cohort and year by age and sex.

To compare sex differences in specific mortality rates of 40-70-year-old Sri Lankans from 1980 to 2010, mortality rates of the 1936-1940 birth cohort for which all age groups were included in the analyses were plotted against the age groups. Using multiple regression analysis, the slopes of the regression lines using mortality rates as the dependent variable and age as the independent variable between males and females were compared.

Patient and public involvement

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

RESULTS

The mortality rates due to ischaemic heart disease, hypertensive disease and cerebrovascular disease by age group,

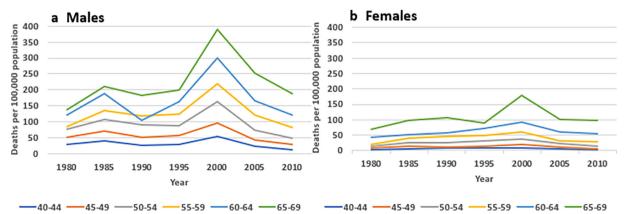


Figure 2 Mortality rates due to ischaemic heart disease among 40-69-year-old Sri Lankans by year by age groups and sex. a For males. b For females.

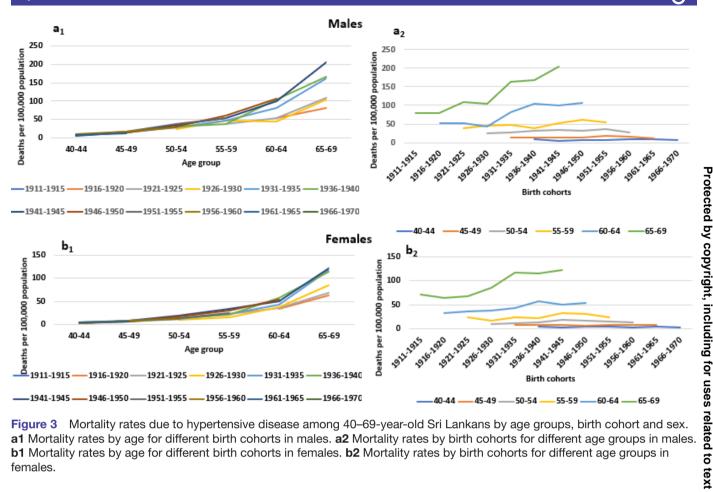


Figure 3 Mortality rates due to hypertensive disease among 40–69-year-old Sri Lankans by age groups, birth cohort and sex. a1 Mortality rates by age for different birth cohorts in males. a2 Mortality rates by birth cohorts for different age groups in males. b1 Mortality rates by age for different birth cohorts in females. b2 Mortality rates by birth cohorts for different age groups in females.

birth cohort and sex are given in figures 1-6 and online supplemental tables 3-5, respectively.

Mortality rates due to ischaemic heart disease increased with age among both males and females in each birth cohort (figures 1 and 2 and online supplemental table 3), the rates being higher in males than in females for each age group in each birth cohort each year. The mortality rates due to ischaemic heart disease for each age group were higher among the early birth cohorts as compared with the younger birth cohorts among both males and females. In each age group, the mortality rates of each

birth cohort peaked corresponding to the year 2000 in both males and females, after which there was a progressive decline in each subsequent birth cohort.

Similar to mortality rates due to ischaemic heart disease, mortality rates due to hypertensive disease increased with age among both males and females (figures 3 and 4 and online supplemental table 4); the mortality rates were training, and similar technologies higher in males than in females for each birth cohort and age group. The highest mortality rates due to hypertensive disease for each age group were higher among the early birth cohorts as compared with the later birth cohorts

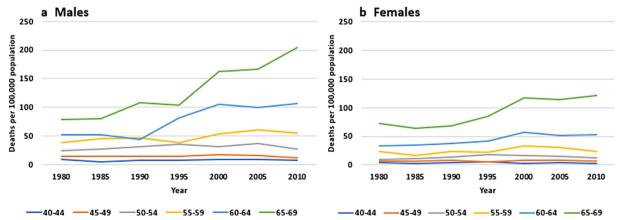


Figure 4 Mortality rates due to hypertensive disease (HTN) among 40–69-year-old Sri Lankans by year by age groups and sex. a For males. b For females.

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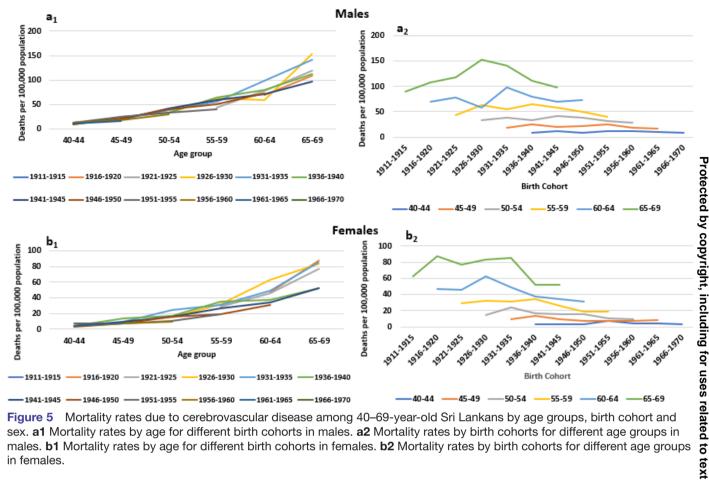


Figure 5 Mortality rates due to cerebrovascular disease among 40–69-year-old Sri Lankans by age groups, birth cohort and sex. a1 Mortality rates by age for different birth cohorts in males. a2 Mortality rates by birth cohorts for different age groups in males. b1 Mortality rates by age for different birth cohorts in females. b2 Mortality rates by birth cohorts for different age groups in females.

among both males and females, the rates being higher in males. Among both males and females, the mortality rates due to hypertensive disease of the 65-69 age group increased from the 1911-1915 to the 1941-1945 birth cohorts; spikes are seen in the mortality rates corresponding to the year 2000. In the 1936–1940 birth cohort, the mortality rate due to hypertensive disease increased from 8.4 per 100000 population in the 40-44-year age group to 167.1 per 100000 population in the 65-69year age group. Similar to the trends in ischaemic heart disease, a spike in the mortality rates due to hypertensive

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and disease is seen in all birth cohorts in different age groups corresponding to the year 2000.

data mining Similar to ischaemic heart disease and hypertensive disease, mortality due to cerebrovascular disease (CeVD) increased with age among both males and females (figures 5 and 6 and online supplemental table 5), with the mortality rates being higher in males than in females for each birth cohort. The mortality rates due to CeVD for each age group were higher among the early birth cohorts as compared with the later birth cohorts among both males and females. Unlike the distinct spike in

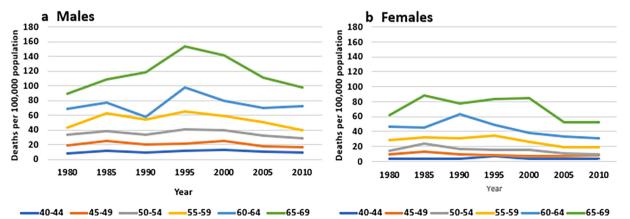


Figure 6 Mortality rates due to cerebrovascular disease (CeVD) among 40–69-year-old Sri Lankans by year by age groups and sex. a For males. b For females.

mortality rates due to ischaemic heart disease and hypertensive disease corresponding to year 2000 data, the spike for CeVD mortality rates was less marked. The mortality rates due to cerebrovascular diseases among males of the earlier cohorts for the older age groups (the 65–69-year age group of the 1926–1930 cohort and the 60–64-year age group of the 1931–1935 cohort) peaked corresponding to the year 1995 (figure 6); among females, this peak was less distinct for the same year. In subsequent birth cohorts, the mortality rates have been declining for each age group.

Mortality rates due to IHD, HTN and CeVD among 40–69-year-old Sri Lankans of birth cohort 1936–1940 by sex are given in online supplemental figures 1–3, respectively. Mortality rates due to all three causes show an increasing trend in both sexes, with the mortality rates of males being higher than that of females. There is a significant difference in the mortality rates for all three causes with age between males and females (p<0.05).

DISCUSSION

The findings of this study show age, sex and 5-year birth cohort effects of selected cardiovascular deaths in 40-69-year-old Sri Lankans from 1980 to 2010. As expected, cardiovascular mortality rates due to IHD, HTN and CeVD increased with age; in each age group, mortality rates were higher among men. In the older age groups, the mortality rates due to HTN increased in the early cohorts until about the year 2000; in the subsequent cohorts, age group mortality continued to decrease over time. CeVD mortality rates in males peaked in the older age groups (65-69 and 60-64) corresponding to the year 1995 with a decline in younger cohorts over time; among females, the trend was less marked. Although the number of deaths increased with time for different age groups due to an ageing population, the mortality rates decreased with each birth cohort.

An age-period-cohort study of CVD in Japan using data from 1995 to 2018 showed that mortality rates of both sexes increased with age,²¹ similar to what is reported here. The Japanese study also revealed an association between other comorbidities with advancing age, such as diabetes mellitus, and atherosclerosis and age-related changes in the cardiovascular system, such as reduced elasticity in blood vessels, increased arterial stiffness and hypertrophy of the heart, etc which could be probable causes for the higher mortality in older age.²¹ Another Japanese study on mortality due to IHD from 1955 to 2000 showed nonlinear birth cohort effects with an initial increase followed by a decreasing trend in both sexes.¹⁶

The findings of this study suggest that mortality rates due to HTN start to rise after about 50 years of age. Again, there are birth cohort effects, with younger cohorts having lower mortality rates and males having higher rates.²² In Mexico, mortality due to hypertension affected more women than men; in the recent cohorts, the risk of dying from hypertension is two times higher in men compared BMJ Open: first published as 10.1136/bmjopen-2024-094083 on 19 May 2025. Downloaded from http://bmjopen.bmj.com/ on May 20, 2025 at Department GEZ-LTA Erasmushogeschool

with women. Hypertensive kidney disease was the main underlying cause, with an average increase throughout the study period.²²

Similar to the trends of HTN and CeVDs, mortality rates are relatively constant in younger birth cohorts up to about 50 years. After 1985, there is a declining trend in mortality across the birth cohorts. Similar findings were reported from China, where age-standardised stroke mortality rates started declining in every age group.²³ In Japan, age-standardised stroke mortality rates per 100 000 population have declined from 98 in 1990, to 74 in 2000, to 50 in 2010 and to 33 in 2019 in males; among females, the decline was from 69 in 1990, to 46 in 2000, to 27 in 2010 and to 18 in 2019.^{24 25}

8 In this study, the overall mortality rates of males are higher than those of females for all three cardiovascular diseases. Similar results were reported from southern Spain²⁶ and China.²⁷ The differences in mortality between males and females are partly due to biological differences including the protective effect of oestrogen in females; after menopause, this effect gradually declines, but the prevalences of other risk factors are lower than that of males. Other factors such as healthcare-seeking behaviour and social determinants also influence these differences. In Sri Lanka, females are subjected to routine screening at different ages during their lifetime from birth including antenatal and postnatal care clinics and well-woman clinics; female attendance at healthy lifestyle to text centres where screening for CVD risk factors is done is much higher than that of males.

An interesting feature in this analysis is the spikes seen in mortality rates of all three cardiovascular deaths. For IHD and HTN, the spikes correspond to data of the year 2000; for CeVD, the spike was less conspicuous. A possible reason for this observation is the change of the ICD coding system used from version 9 to 10 in 1997, despite ICD-10 being released in 1995¹⁹; ICD version 10 ≥ had more specific cause of death codes and had more causes of death for each disease, which is likely to have ĝ increased the mortality rates for each cause of death. The subsequent decrease in the mortality rates may be due to the natural decrease in the mortality rates seen with S successive birth cohorts.

Mortality rates of all three cardiovascular diseases considered show similar trends; like all non-communicable diseases, these disease entities share distinct common characteristics that are influenced by a broader range of lifestyle factors such as smoking, diet, physical activity and obesity, which have been targeted in preventive programmes.

Decline in mortality rates in successive birth cohorts may be due to many factors. First, it may be due to differences in exposures in utero, childhood and even adulthood. Implementation of screening programmes and advances in medical and surgical procedures have contributed largely to early diagnosis and secondary prevention increasing the life expectancy of those diagnosed with cardiovascular diseases.⁷ In addition, large declines in smoking due to increased public awareness and regulation, including heavy taxation, are now considered a 'best buy' for reduction of cardiovascular disease.' Ma et al suggested that lower mortality in younger birth cohorts in Japan was probably a result of improvements in lifestyle factors, including the national hypertension control prevention programme and improved nutrition in Japan during the previous few decades.¹⁶ It is also likely to be due to improved economic status over time. The per capita GNI in Sri Lanka was current dollars 2510 in 1990; in 2010 it was 8150.

Over several decades, Sri Lanka's health system has improved tremendously in all aspects, achieving impressive health indicators comparable to those of developed countries but at a much lower cost. This achievement is attributable to an excellent preventive health service originally initiated in 1926; though initially concentrating on maternal and child health, the service has been extended to cover environmental health and later to non-communicable diseases after the inclusion of prevention and control of non-communicable diseases in the first national health policy based on primary healthcare and subsequently revised with a focus on universal health coverage.²⁸ Simultaneously, clinical services were improved and expanded with an increase in the number of hospital beds, advancements in technology and other related services. The number of doctors per 10000 population has increased from 1401 in 1991 to 11924 in 2021.²⁹ Initially, general physicians in medical wards treated cardiology patients; subsequently, since the early 1970s, cardiology as a specialty emerged in Sri Lanka. Specialisation in cardiology and cardiothoracic surgery followed, and special units were created; in the public sector in 2020, there were 644 inward beds for cardiology compared with 470 in 2008.^{30 31} The improvement and expansion of healthcare services and the emergence of cardiology as a specialty beginning in the 1970s probably explain the high mortality rates in older age groups of early cohorts.

In Sri Lanka, preventative activities carried out by the Ministry of Health at the national level are likely to have contributed to improvements in cardiovascular health among the younger generations: these include tobacco control measures (both policy and legislation); nutrition and food policies to address obesity and high lipid levels; and health promotion to raise awareness about the importance of cardiovascular health and the prevention of CVDs.³²

Limitations

Data for 1990 and 2008 were not available, and data were interpolated separately for males and females. The fact that the validity of our results is dependent on the accuracy and reliability of mortality data and the vital statistics registration system of the country is acknowledged. Studies have reported that cause of death certification is not 100% accurate, as expected.^{33 34} However, given the CVD causes of death that were investigated, it is unlikely to have adversely impacted our results; the rates were calculated based on large denominators that would have little or no effect on the overall mortality rates.

Relating the differences in mortality rates to economic indicators was not attempted. The GNI was available from 1990 onwards and requires a more detailed analysis that should be conducted in the future.

CONCLUSIONS

Protected There are age, period and birth cohort effects on mortality due to IHD, HTN and CeVD among 40-69-year-old male d by and female Sri Lankans between 1980 and 2010. Mortality increased with age and declined in younger cohorts over r copyrigh time as compared with older cohorts. Males had higher mortality rates than females for each age group in each birth cohort.

It is likely that age-specific and sex-specific mortality rates due to cardiovascular diseases will decline further, but at a much lower rate, provided that prevention and control measures for CVD risk factors are enhanced and a sustained; however, the number of deaths due to cardiouses related to text and data mining, Al training, and similar technologies vascular diseases will increase due to the increasing ageing population for which adequate care facilities should be improved, expanded and provided.

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Contributors DTHDeS, EDSMDeA, DMDeM, AHDDeS and ARW contributed to the conceptualisation and developing the methodology. ARW supervised data collection. All authors were involved in data curation, data analysis, writing the original draft, and reviewing and editing the final draft. ARW is the guarantor and accepts full responsibility for the work and conduct of the study.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval The Ethics Review Committee of the Faculty of Medicine, University of Kelaniya, Sri Lanka, exempted this study from ethics review (Ref No. P/149/11/2022).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as supplementary information. All data extracted are given in the supplementary tables and in the tables.

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