Check for updates

OPEN ACCESS

EDITED BY Yashendra Sethi, PearResearch, India

REVIEWED BY Erik K. Arnesen, University of Oslo, Norway Abhinav Choubey, Baylor College of Medicine, United States

*CORRESPONDENCE Yuzhe Kong csuyuzhekong@foxmail.com

RECEIVED 24 August 2024 ACCEPTED 11 November 2024 PUBLISHED 07 January 2025

CITATION

Mao Q and Kong Y (2025) Effect of diet low in omega-6 polyunsaturated fatty acids on the global burden of cardiovascular diseases and future trends: evidence from the Global Burden of Disease 2021. *Front. Med.* 11:1485695. [doi: 10.3389/fmed.2024.1485695](https://doi.org/10.3389/fmed.2024.1485695)

COPYRIGHT

© 2025 Mao and Kong. This is an open-access article distributed under the terms of the [Creative Commons Attribution](http://creativecommons.org/licenses/by/4.0/) [License \(CC BY\)](http://creativecommons.org/licenses/by/4.0/). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

[Effect of diet low in omega-6](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full) [polyunsaturated fatty acids on the](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full) [global burden of cardiovascular](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full) [diseases and future trends:](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full) [evidence from the Global Burden](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full) [of Disease 2021](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full)

Qingsong Mao¹ and Yuzhe Kong^{2*}

1 Hepatobiliary Pancreatic Surgery, Banan Hospital Affiliated of Chongqing Medical University, Banan, China, 2Xiangya School of Medicine, Central South University, Changsha, China

Background: This research analyzes the worldwide impact of cardiovascular diseases (CVD) associated with low consumption of omega-6 polyunsaturated fatty acids, utilizing data from the 2021 Global Burden of Disease Study.

Method: The study explored the influence of diets deficient in omega-6 polyunsaturated fatty acids on CVD across global, regional, and national levels. It examined variations across different age groups and genders and analyzed the relationship between the disease burden and the socio-demographic index (SDI). Furthermore, it employed an ARIMA model to project the future prevalence of CVD linked to insufficient omega-6 intake until 2050.

Result: In 2021, insufficient omega-6 intake was linked to roughly 737.88 thousand deaths and 17.87 million disability-adjusted life years (DALYs) due to CVD, showing a decreasing trend in this health burden throughout the study period. The most significant effects were seen in individuals aged 75 and older, with a higher disease burden noted in males. Forecasts suggest likely declines in disease prevalence in regions with high SDI. On a national level, regions like Russia and various countries in North Africa and the Middle East might experience increasing challenges related to CVD due to low omega-6 intake by 2030 and 2050.

Conclusion: These results highlight the critical need for preventive strategies for CVD and stress the importance of managing dietary patterns to mitigate health risks.

KEYWORDS

cardiovascular disease, omega-6 polyunsaturated fatty acids, mortality forecasting, epidemiology, disease burden

1 Introduction

Cardiovascular diseases (CVD) are the primary cause of mortality worldwide [\(1\)](#page-11-0). Numerous risk factors including smoking, dyslipidemia, hypertension, obesity, lack of physical activity, and ethnicity are closely associated with CVD [\(1\)](#page-11-0). It is noteworthy that behavioral risk factors could prevent up to 90% of CVD incidents ([2\)](#page-11-1). Additionally, dietary habits and specific dietary components significantly influence CVD ([3\)](#page-11-2). LA, a crucial fatty acid that must be ingested

through diet sources like vegetable and nut oils, poultry, meat, eggs, milk, and margarine, cannot be synthesized by humans [\(4](#page-11-3), [5](#page-11-4)). AA, derived from LA, is also a key PUFA [\(6](#page-11-5), [7](#page-11-6)).

Growing research highlights the potential impact of omega-6 PUFAs on reducing CVD risk [\(8](#page-11-7)[–10\)](#page-12-0). For instance, a meta-analysis found that higher LA levels in the body were linked to a reduced risk of coronary heart disease (CHD) [\(11\)](#page-12-1). However, no significant association was found between AA levels and CHD risk ([8\)](#page-11-7). Another analysis demonstrated that increasing dietary LA by 5% of total energy, substituting saturated fats, correlated with a 9% reduction in CHD incidents [\(9\)](#page-12-2). A comprehensive analysis from 11 studies indicated that omega-6 PUFAs were associated with a decrease in both CHD events and mortality, showing hazard ratios of 0.87 and 0.74, respectively [\(8\)](#page-11-7). Another study showed replacing saturated fats with omega-6 PUFAs significantly cut total cholesterol by 19%, LDL-C by 22%, and increased HDL-C by 14% among nutrition students [\(9](#page-12-2)). Further, each 5% replacement of saturated fats with PUFAs reduced LDL-C by 10 mg/dL and the TC ratio by 0.16 ([12](#page-12-3)).

Despite these findings, some studies offer mixed results. The Sydney Diet Heart Study, an RCT with 458 middle-aged men with recent coronary events, reported increased mortality rates when replacing saturated fats with LA [\(13\)](#page-12-4). Additionally, comprehensive reviews have found no definitive links between omega-6 PUFA intake and CVD outcomes [\(14](#page-12-5), [15](#page-12-6)). It is suggested that diets high in omega-6 PUFAs might elevate proinflammatory mediators like prostaglandins and leukotrienes, which could adversely affect CVD risk ([4,](#page-11-3) [16](#page-12-7)).

The limitations of individual studies, including small sample sizes, variations in research design, and diverse participant profiles, underline the need for broader investigations. This study utilized the 2021 Global Burden of Disease data to explore the effects of a diet low in omega-6 PUFAs on CVD. Moreover, future CVD trends were predicted using the ARIMA model, validated by previous research [\(17,](#page-12-8) [18](#page-12-9)).

2 Method

2.1 Study population

Our investigation drew upon the Global Burden of Disease Study 2021, sourced from [http://ghdx.healthdata.org/gbd-results-tool,](http://ghdx.healthdata.org/gbd-results-tool) encompassing assessments of 369 health conditions and the influence of 87 factors across 204 nations over the period from 1990 to 2021 [\(19\)](#page-12-10).

The study elaborates on methodologies applied to gauge the effects of cardiovascular diseases (CVD). The detailed methods are outlined in referenced studies [\(20,](#page-12-11) [21\)](#page-12-12). CVD mortality data were collected from various sources like vital registration systems and health surveys, among others, and were adjusted to account for inaccuracies and misclassifications ([22](#page-12-13)). These adjusted data were fed into the Cause of Death Ensemble model (CODEm) to produce annual CVD mortality estimates by age and sex for each region [\(21,](#page-12-12) [23\)](#page-12-14). Additionally, a comparative risk assessment pinpointed key factors contributing to CVD, with the population attributable fraction (PAF) being calculated to measure the impact of a diet low in omega-6 polyunsaturated fatty acids on CVD [\(19\)](#page-12-10). Both mortality and disability-adjusted life years (DALYs) associated with this dietary risk were assessed, where DALYs are comprised of years lost due to premature death (YLLs) and years lived with disability (YLDs). YLD calculations considered the prevalence of these infections, adjusted by disability severity [\(22\)](#page-12-13). The sociodemographic index (SDI) was computed considering factors such as fertility rates in individuals under 25 (TFU25), educational attainment

for those aged over 15 (EDU15+), and adjusted *per capita* income. Countries were categorized into five SDI levels: low, low-middle, middle, high-middle, and high [\(22](#page-12-13), [24\)](#page-12-15). In our studies, we used mortality and DALYs to estimate the risks.

2.2 Definition of diet low in omega-6 polyunsaturated fatty acids

A diet low in omega-6 polyunsaturated fatty acids is characterized by consuming less than 9–10% of total daily energy from sources such as linoleic acid, γ-linolenic acid, eicosadienoic acid, dihomo-γlinolenic acid, and arachidonic acid.¹

2.3 Statistical analysis

Our statistical analysis involved standardizing mortality and DALY rates for international comparisons using age-adjusted rates (AAR) to normalize for age distribution and demographic differences among countries. We employed a linear regression on the logarithmic values of these rates, expressed as $y = \alpha + \beta x + \varepsilon$, where *x* denotes the year. The estimated annual percentage change (EAPC) was derived using the equation 100 $*(e^{\beta} - 1)$, along with its 95% confidence interval (95% CI). Trends in AAR were categorized as increasing, decreasing, or stable based on the direction and magnitude of the EAPC and the limits of the 95% CI [\(25](#page-12-16), [26\)](#page-12-17). The relationship between AAR and SDI was explored using Gaussian process regression with Loess smoothing and analyzed through Spearman rank correlation tests [\(24,](#page-12-15) [26](#page-12-17), [27](#page-12-18)).

Additionally, the ARIMA model was applied to examine and forecast the influence of low omega-6 polyunsaturated fatty acid diets on CVD trends from 2020 to 2050. This model's configuration (p, d, q) includes "AR" for the autoregressive component, "MA" for the moving average terms, and "d" for the degree of differencing needed for data stability [\(28\)](#page-12-19). The model's efficacy was judged based on the Akaike information criterion (AIC) and the Bayesian information criterion (BIC).

All metrics were calculated with 95% uncertainty intervals and presented per 100,000 individuals. Data handling, analysis, and visualization were conducted using R software, version 4.3.2 ([29](#page-12-20)[–32](#page-12-21)).

3 Result

3.1 Spatiotemporal patterns of CVD attributable to diet low in polyunsaturated fatty acids

In 2021, a diet deficient in omega-6 polyunsaturated fatty acids contributed to approximately 737,880 deaths [percentage (95% UI): 3.78 (−11.29, 14.64)] and 17.87 million disability-adjusted life years (DALYs) [percentage (95% UI): 4.15 (−13.48, 15.59)] from cardiovascular diseases (CVDs). The age-standardized mortality rate

(ASMR) was 8.7966 per 100,000 population, with a 95% uncertainty interval (UI) ranging from −25.1372 to 33.0790, and the age-standardized DALY rate (ASDR) was 208.1459 per 100,000, with a 95% UI from −651.9420 to 768.8355. Over the last 30 years, there has been a significant reduction in the impact of diets low in omega-6 polyunsaturated fatty acids on CVDs ([Table 1](#page-3-0); [Supplementary Table S1\)](#page-11-8).

Regarding the socio-demographic index (SDI), higher SDI regions have seen a notable decline in CVD burden due to diets low in omega-6 fatty acids, with the largest decreases observed. Across all SDI categories, modest reductions were observed in both ASMR and ASDR for CVDs linked to these diets, with the exception of the high SDI regions [\(Table 1;](#page-3-0) [Supplementary Table S1](#page-11-8) and [Figure 1\)](#page-5-0).

On a regional level, the greatest burdens from diets low in omega-6 polyunsaturated fatty acids were found in Asia and Europe. In contrast, Oceania and Australia reported the lowest burdens associated with such diets [\(Table 1](#page-3-0)).

Nationally, in 2021, there was considerable global variation in the ASDR and ASMR for CVDs associated with omega-6 fatty aciddeficient diets. The highest rates were seen in North Africa, the Middle East, and Russia [\(Figure 2\)](#page-6-0). From 1990 to 2019, a general trend of reductions in both ASMR and ASDR was noted across most countries, with the exception of those in North Africa and the Middle East ([Table 1](#page-3-0); [Supplementary Table S1](#page-11-8) and [Figure 3](#page-6-1)).

3.2 Age and gender pattern

[Figure 2](#page-6-0) depicts the age-specific global mortality and DALY rates for cardiovascular diseases (CVDs) in 2021, along with their trends from 1990 to 2021. The graph demonstrates a J-shaped curve, indicating rising mortality and DALY rates in individuals under 75 years, with a marked increase observed in the 75 to 95 and older age group. Across all age groups, males consistently registered higher mortality rates due to diets low in omega-6 polyunsaturated fatty acids compared to females. Likewise, DALY rates linked to such diets were elevated across all ages for males. Moreover, the reduction in mortality and DALY rates from 1990 to 2021 was less substantial among males in every age group.

Across all socio-demographic index (SDI) categories, male mortality and DALY rates were consistently higher, maintaining this gender disparity across the regions. However, in regions with highmiddle and high SDI, the differences between genders have lessened, whereas in other SDI regions, the disparities have either remained unchanged or increased ([Figure 4\)](#page-7-0).

3.3 Association with the socio-demographic index

[Figure 5](#page-8-0) displays a comparison between the actual and predicted age-standardized DALY (ASDR) and mortality rates (ASMR) for CVD related to diets low in omega-6 polyunsaturated fatty acids, aligning these with socio-demographic index (SDI) values at both regional and national levels from 1990 to 2021. A discernible negative correlation exists between ASDR and rising SDI, suggesting a decrease in disease burden as SDI increases. Regions such as Eastern Europe, Central Asia, North Africa, and the Middle East exhibited higher-thanexpected ASDR during this period. The pattern of observed versus forecasted ASMR based on SDI at regional levels mirrors the ASDR

¹ [https://www.healthdata.org/research-analysis/diseases-injuries-risks/](https://www.healthdata.org/research-analysis/diseases-injuries-risks/factsheets/2021-diet-low-omega-6-polyunsaturated-fatty)

[factsheets/2021-diet-low-omega-6-polyunsaturated-fatty](https://www.healthdata.org/research-analysis/diseases-injuries-risks/factsheets/2021-diet-low-omega-6-polyunsaturated-fatty)

TABLE 1 Global and regional deaths and DALYs of CVD attributable to diet low in omega-6 polyunsaturated fatty acids in 1990 and 2021 in 27 global regions.

(Continued)

TABLE 1 (Continued)

findings. [Figure 5](#page-8-0) also demonstrates the observed versus anticipated ASDR and ASMR at the national level for 2021, showing a similar negative correlation with SDI values, both regionally and nationally.

3.4 Forecasts for the mortality, DALYs rate, ASMR and ASDR of CVD attributable to diet low in polyunsaturated fatty acids in global (2022–2050)

[Figures 6](#page-9-0), [7](#page-10-0) outline the projected trends in mortality and DALY rates, including ASMR and ASDR for CVDs linked to diets deficient in omega-6 polyunsaturated fatty acids. At the regional scale, decreases in CVD burden are forecasted solely within the high and high-middle SDI regions, whereas other SDI regions are likely to maintain current rates regarding ASDR and ASMR. Middle and low-middle SDI

regions, on the other hand, are projected to experience increases in both mortality and DALY rates (see [Figure 8](#page-10-1)).

On a national scale, these patterns are expected to remain fairly consistent through the years 2030 and 2050. Nonetheless, the projected consequences of CVDs related to diets low in omega-6 polyunsaturated fatty acids are anticipated to be significantly greater in Russia and throughout the North Africa and Middle East region than in other regions for both 2030 and 2050.

4 Discussion

This investigation assessed the global consequences of cardiovascular disease (CVD) linked to diets deficient in omega-6 polyunsaturated fatty acids over the period from 1990 to 2021,

identifying a significant decline. The most impacted demographic was males over the age of 75, who displayed elevated incidence rates. Projections for the future indicate a likely reduction in mortality rates in regions with higher social development index (SDI) scores, although high mortality and age-standardized death rates from such diets are expected to continue in Russia and the North Africa and Middle East region. This study represents the first extensive review of CVD effects

stemming from insufficient intake of omega-6 polyunsaturated fatty acids.

Prior analyses, including a meta-analysis of 25 case-control studies, found a negative correlation between levels of linoleic acid (LA) in blood and tissues and coronary heart disease (CHD) risk [\(8\)](#page-11-7). While arachidonic acid (AA) was not significantly linked to CHD risk, lower levels of AA in adipose tissue were tied to increased CHD event

Age-specific rates of global deaths (a) and DALYs (b) of CVD attributable to diet low in omega-6 polyunsaturated fatty acids, by sex, in 2021 and the corresponding EAPC of global deaths (c) and DALYs (d) from 1990 to 2021.

Global distribution of ASMR (a) and ASDR (c) of CVD attributable to diet low in omega-6 polyunsaturated fatty acids for both sexes in 2021 in 204 countries and territories. EAPC of ASMR (b) and ASDR (d) of CVD attributable to diet low in omega-6 polyunsaturated fatty acids from 1990 to 2021 in 204 countries and territories.

risk in longitudinal studies, yet were associated with a reduced risk in cross-sectional analyses [\(8\)](#page-11-7). A different meta-analysis covering 13 cohort studies involving 30,602 subjects (with 12,479 CHD events, including 5,882 fatalities) highlighted the protective effects of increased LA intake against CHD occurrences and mortality, indicating that substituting saturated fats with LA could lower CHD event and death risks by 9 and 13%, respectively ([9\)](#page-12-2). Earlier analyses, including 60 clinical trials ([33](#page-12-22)), and another encompassing 8 randomized control trials (RCTs) with 13,614 participants experiencing 1,042 CHD events [\(12\)](#page-12-3), noted that omega-6 PUFAs positively influenced blood lipids and reduced CHD events. Moreover, a Cochrane review of dietary fat modifications through RCTs indicated that substituting saturated fats with monounsaturated or polyunsaturated fatty acids modestly but effectively decreased CVD risk over a minimum of 6 months ([34](#page-12-23)).

A study of 1,551 middle-aged men explored the impact of dietary LA and total PUFA consumption on CVD mortality, revealing an inverse relationship with CVD deaths [\(35\)](#page-12-24). A systematic review of 19 studies including 6,461 adults (average age: 50 years) suggested minimal effects on mortality or CVD events over 1–8 years by increasing omega-6 PUFA intake, though it could potentially lower heart attack risks [\(36\)](#page-12-25). Omega-6 PUFAs were found to reduce serum

total cholesterol (TC) by 6%, without significant changes to triglycerides (TG), high-density lipoprotein cholesterol (HDL-C), and low-density lipoprotein cholesterol (LDL-C). It is important to note that limited event numbers and the underrepresentation of participants from developing countries and women may have affected these findings ([13\)](#page-12-4).

Additional cohort research indicates that a diet low in saturated fatty acids and enriched with omega-6 PUFAs correlates with decreased CHD risk and lower levels of LDL-C [\(10,](#page-12-0) [11](#page-12-1)). Jakobsen et al. [\(11\)](#page-12-1) demonstrated that substituting carbohydrates with PUFAs (omega-6) can reduce levels of TC, HDL-C, LDL-C, and TG. However, the validity of these findings is limited due to the short duration of interventions and the inclusion of non-randomized studies, which may undermine their clinical relevance [\(34\)](#page-12-23).

The impact of omega-6 PUFAs on CVD risk remains debated. While some studies support the positive effects of omega-6 PUFA supplementation on cardiovascular outcomes, contrasting studies present divergent results. A potential explanation for the complex relationship between omega-6 PUFAs and CVD risk involves the enhanced synthesis of 2-series prostaglandins and 4-series leukotrienes, which are proinflammatory and may adversely influence residual CVD risk [\(4](#page-11-3), [16](#page-12-7)). Furthermore, compounds like AA, a highly unsaturated

fatty acid, could increase LDL-C and VLDL oxidation, elevating their atherogenic potential [\(4\)](#page-11-3). Conversely, certain investigations show no correlation between omega-6 PUFA consumption and CVD risk, including a 14-year longitudinal study involving 43,732 men which found no link between overall fat consumption, cholesterol, and stroke risk [\(37\)](#page-12-26). Additionally, an analysis of prospective observational studies and RCTs indicated that high PUFA intake coupled with low saturated fat intake did not significantly alter CVD outcomes [\(15](#page-12-6)). Likewise, a Cochrane review involving 660 participants detected no significant connections between varying omega-6 PUFA intake levels and CVD risk factors [\(14\)](#page-12-5). Furthermore, another systematic review concluded that replacing saturated fats with cis-PUFAs, akin to omega-6 PUFAs, markedly reduced TC, supporting findings of our current study [\(38](#page-12-27)–[46\)](#page-12-28).

Previously, another study analyzes the global impact of nonoptimal dietary fat intake on coronary heart disease (CHD) mortality, highlighting the health burden linked to low ω-6 polyunsaturated fat (PUFA) intake and high saturated (SFA) and trans fats (TFA) consumption. It reveals significant disparities in CHD mortality, particularly in countries with poor dietary management and lower socioeconomic status. Targeted public health interventions, such as promoting ω-6-rich foods and reducing SFA and TFA intake, could substantially lower CHD rates, especially in high-risk regions. Additionally, regional variations indicate that global dietary guidelines should be tailored to local nutritional contexts. The study underscores the effectiveness of past interventions while revealing ongoing challenges in food policy. Ultimately, it provides a vital evidence base for future public health strategies to combat diet-related CHD worldwide. Future research should continue to track these trends and evaluate the impact of targeted dietary interventions ([33](#page-12-22)).

Firstly, the variability in results could be attributed to the heterogeneity in dietary sources and the composition of omega-6 PUFAs in different studies. For example, arachidonic acid (AA), a type of omega-6 PUFA, has been shown in some studies to promote inflammatory processes that could exacerbate CVD risks. In contrast, linoleic acid (LA), another omega-6 PUFA, is often associated with protective effects against CVD. This discrepancy suggests that not all omega-6 PUFAs uniformly affect CVD risk, and their impact could be context-dependent based on their specific biochemical pathways and interactions within the body.

Secondly, the methodological differences across studies, such as the duration of follow-up, the age and baseline health conditions of participants, and how dietary intake was assessed, can also lead to conflicting results. Longitudinal studies might capture more reliable effects of long-term dietary patterns, whereas cross-sectional studies may not adequately reflect the chronic nature of dietary impacts on cardiovascular health.

Furthermore, genetic factors could play a significant role in modulating the effects of omega-6 PUFAs on CVD risk. Individual genetic variability in fatty acid metabolism might influence how these fats affect the individual's risk profile. This aspect is often not fully accounted for in many studies, leading to an incomplete picture.

To enhance the credibility and depth of our discussion on this topic, future research should aim to standardize the types of omega-6 PUFAs studied and consider a more detailed analysis of the genetic and metabolic contexts of study populations. Additionally, more rigorous control of dietary assessment methods and a comprehensive evaluation of inflammatory markers could provide clearer insights into the

FIGURE 7 Estimated trends of mortality rate (a,b) and ASMR (c,d) in 2030 (a,c) and 2050 (b,d) at the national level.

pathways through which omega-6 PUFAs influence CVD risk. Addressing these complexities in a detailed manner would not only resolve the existing controversies but also significantly advance our understanding of the nutritional management of cardiovascular health.

This study has several limitations. First, the time frame of the data used primarily comes from the Global Burden of Disease (GBD) database, which provides epidemiological information for the selected period but does not adequately reflect the latest trends in dietary habits and public health interventions. This may impact the generalizability of the findings. Second, the study lacks data on genetic susceptibility and lifestyle factors (such as physical activity, smoking, and psychological stress), limiting our ability to explore the potential effects of these variables on cardiovascular disease risk. Finally, the analysis does not delve into the molecular mechanisms by which omega-6 polyunsaturated fatty acids (PUFAs) influence cardiovascular disease, restricting our understanding of the mechanisms of action and preventing the formulation of new intervention strategies.

Besides, a notable limitation is the inconsistent data quality and accessibility from less developed areas, particularly Sub-Saharan Africa. Relying on mathematical models to estimate CVD burdens in these regions introduces considerable uncertainty. Moreover, while the study spans from 1990 to 2021, offering a long-term overview, it may not capture the latest dietary shifts or the impact of recent public health initiatives, potentially skewing the study's findings. This temporal limitation indicates that the latest changes in dietary habits and health policies might not be accurately reflected in the study's outcomes.

5 Conclusion

This study thoroughly analyzes the global influence of cardiovascular disease (CVD) connected to insufficient intake of omega-6 polyunsaturated fatty acids from 1990 to 2021, highlighting a significant decrease over the period. The group most impacted included seniors above the age of 75, with men experiencing a particularly high incidence. Predictions suggest that areas with high socio-demographic index (SDI) values could see decreases in mortality, DALY rates, ASMR, and ASDR in the future. At the national level, places like Russia and regions in North Africa and the Middle East are expected to continue experiencing substantial CVD burdens linked to diets deficient in omega-6 polyunsaturated fatty acids through 2030 and 2050.

These insights are vital for developing CVD prevention tactics and underline the need to regulate dietary patterns.

References

1. Roth GA, Mensah GA, Johnson CO, Addolorato G, Ammirati E, Baddour LM, et al. Global burden of cardiovascular diseases and risk factors, 1990–2019. *J Am Coll Cardiol*. (2020) 76:2982–3021. doi: [10.1016/j.jacc.2020.11.010](https://doi.org/10.1016/j.jacc.2020.11.010)

2. Booth JN III, Colantonio LD, Howard G, Safford MM, Banach M, Reynolds K, et al. Healthy lifestyle factors and incident heart disease and mortality in candidates for primary prevention with statin therapy. *Int J Cardiol*. (2016) 207:196–202. doi: [10.1016/j.](https://doi.org/10.1016/j.ijcard.2016.01.001) [ijcard.2016.01.001](https://doi.org/10.1016/j.ijcard.2016.01.001)

3. Australian Institute of Health and Welfare. Burden of disease and injury in Australia. Canberra: Australian Institute of Health and Welfare (2003).

4. Russo GL. Dietary *n* − 6 and *n* − 3 polyunsaturated fatty acids: from biochemistry to clinical implications in cardiovascular prevention. *Biochem Pharmacol*. (2009) 77:937–46. doi: [10.1016/j.bcp.2008.10.020](https://doi.org/10.1016/j.bcp.2008.10.020)

Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Author contributions

QM: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Validation, Visualization, Writing – original draft. YK: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Project administration, Resources, Software, Supervision, Validation, Visualization, Writing – original draft, Writing – review & editing.

Funding

The author(s) declare that no financial support was received for the research, authorship, and/or publication of this article.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Supplementary material

The Supplementary material for this article can be found online at: [https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full#supplementary-material) [full#supplementary-material](https://www.frontiersin.org/articles/10.3389/fmed.2024.1485695/full#supplementary-material)

5. Bibus D, Lands B. Balancing proportions of competing omega-3 and omega-6 highly unsaturated fatty acids (HUFA) in tissue lipids. *Prostaglandins Leukot Essent Fatty Acids*. (2015) 99:19–23. doi: [10.1016/j.plefa.2015.04.005](https://doi.org/10.1016/j.plefa.2015.04.005)

6. Mann CJ, Kaduce TL, Figard PH, Spector AA. Docosatetraenoic acid in endothelial cells: formation, retroconversion to arachidonic acid, and effect on prostacyclin production. *Arch Biochem Biophys*. (1986) 244:813–23. doi: [10.1016/0003-9861\(86\)90650-8](https://doi.org/10.1016/0003-9861(86)90650-8)

7. Rosenthal MD, Whitehurst MC. Fatty acyl Δ6 desaturation activity of cultured human endothelial cells modulation by fetal bovine serum. *Biochim Biophys Acta*. (1983) 750:490–6. doi: [10.1016/0005-2760\(83\)90189-3](https://doi.org/10.1016/0005-2760(83)90189-3)

8. Harris WS, Poston WC, Haddock CK. Tissue *n* − 3 and *n* − 6 fatty acids and risk for coronary heart disease events. *Atherosclerosis*. (2007) 193:1. doi: [10.1016/j.](https://doi.org/10.1016/j.atherosclerosis.2007.03.018) [atherosclerosis.2007.03.018](https://doi.org/10.1016/j.atherosclerosis.2007.03.018)

9. Farvid MS, Ding M, Pan A, Sun Q, Chiuve SE, Steffen LM, et al. Dietary linoleic acid and risk of coronary heart disease: a systematic review and meta-analysis of prospective cohort studies. *Circulation*. (2014) 130:1568–78. doi: [10.1161/](https://doi.org/10.1161/CIRCULATIONAHA.114.010236) [CIRCULATIONAHA.114.010236](https://doi.org/10.1161/CIRCULATIONAHA.114.010236)

10. Hodson L, Skeaff C, Chisholm WA. The effect of replacing dietary saturated fat with polyunsaturated or monounsaturated fat on plasma lipids in free-living young adults. *Eur J Clin Nutr*. (2001) 55:908–15. doi: [10.1038/sj.ejcn.1601234](https://doi.org/10.1038/sj.ejcn.1601234)

11. Jakobsen MU, O'Reilly EJ, Heitmann BL, Pereira MA, Bälter K, Fraser GE, et al. Major types of dietary fat and risk of coronary heart disease: a pooled analysis of 11 cohort studies. *Am J Clin Nutr*. (2009) 89:1425–32. doi: [10.3945/ajcn.2008.27124](https://doi.org/10.3945/ajcn.2008.27124)

12. Mozaffarian D, Micha R, Wallace S. Effects on coronary heart disease of increasing polyunsaturated fat in place of saturated fat: a systematic review and meta-analysis of randomized controlled trials. *PLoS Med*. (2010) 7:e1000252. doi: [10.1371/journal.](https://doi.org/10.1371/journal.pmed.1000252) [pmed.1000252](https://doi.org/10.1371/journal.pmed.1000252)

13. Ramsden CE, Zamora D, Leelarthaepin B, Majchrzak-Hong SF, Faurot KR, Suchindran CM, et al. Use of dietary linoleic acid for secondary prevention of coronary heart disease and death: evaluation of recovered data from the Sydney Diet Heart Study and updated meta-analysis. *BMJ*. (2013) 346:e8707. doi: [10.1136/bmj.e8707](https://doi.org/10.1136/bmj.e8707)

14. Al-Khudairy L, Hartley L, Clar C, Flowers N, Hooper L, Rees K. Omega 6 fatty acids for the primary prevention of cardiovascular disease. *Cochrane Database Syst Rev*. (2015) 11:CD011094. doi: [10.1002/14651858.CD011094.pub2](https://doi.org/10.1002/14651858.CD011094.pub2)

15. Chowdhury R, Warnakula S, Kunutsor S, Crowe F, Ward HA, Johnson L, et al. Association of dietary, circulating, and supplement fatty acids with coronary risk. *Ann Int Med*. (2014) 160:398–406. doi: [10.7326/M13-1788](https://doi.org/10.7326/M13-1788)

16. Siriwardhana N, Kalupahana NS, Fletcher S, Xin W, Claycombe KJ, Quignard-Boulange A, et al. *n* − 3 and *n* − 6 polyunsaturated fatty acids differentially regulate adipose angiotensinogen and other inflammatory adipokines in part via NF-κBdependent mechanisms. *J Nutr Biochem*. (2012) 23:1661–7. doi: [10.1016/j.](https://doi.org/10.1016/j.jnutbio.2011.11.009) [jnutbio.2011.11.009](https://doi.org/10.1016/j.jnutbio.2011.11.009)

17. Wafa HA, Marshall I, Wolfe CDA, Xie W, Johnson CO, Veltkamp R, et al. Burden of intracerebral haemorrhage in Europe: forecasting incidence and mortality between 2019 and 2050. *Lancet Reg Health Eur*. (2024) 38:100842. doi: [10.1016/j.](https://doi.org/10.1016/j.lanepe.2024.100842) [lanepe.2024.100842](https://doi.org/10.1016/j.lanepe.2024.100842)

18. Gong Y, Jiang Q, Zhai M, Tang T, Liu S. Thyroid cancer trends in China and its comparative analysis with G20 countries: projections for 2020–2040. *J Glob Health*. (2024) 14:04131. doi: [10.7189/jogh.14.04131](https://doi.org/10.7189/jogh.14.04131)

19. GBD 2019 Risk Factors Collaborators. Global burden of 87 risk factors in 204 countries and territories, 1990–2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet*. (2020) 396:1223–49. doi: [10.1016/S0140-6736\(20\)30752-2](https://doi.org/10.1016/S0140-6736(20)30752-2)

20. GBD 2019 LRI Collaborators. Age-sex differences in the global burden of lower respiratory infections and risk factors, 1990–2019: results from the Global Burden of Disease Study 2019. *Lancet Infect Dis*. (2022) 22:1626–47. doi: [10.1016/](https://doi.org/10.1016/S1473-3099(22)00510-2) [S1473-3099\(22\)00510-2](https://doi.org/10.1016/S1473-3099(22)00510-2)

21. Safiri S, Mahmoodpoor A, Kolahi AA, Nejadghaderi SA, Sullman MJM, Mansournia MA, et al. Global burden of lower respiratory infections during the last three decades. *Front Public Health*. (2023) 10:1028525. doi: [10.3389/fpubh.2022.1028525](https://doi.org/10.3389/fpubh.2022.1028525)

22. GBD 2019 Diseases and Injuries Collaborators. Global burden of 369 diseases and injuries in 204 countries and territories, 1990–2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet*. (2020) 396:1204–22. doi: [10.1016/](https://doi.org/10.1016/S0140-6736(20)30925-9) [S0140-6736\(20\)30925-9,](https://doi.org/10.1016/S0140-6736(20)30925-9) Erratum in: Lancet. 2020; 396 (10262): 1562. doi:10.1016/ S0140-6736(20)32226-1

23. Foreman KJ, Lozano R, Lopez AD, Murray CJ. Modeling causes of death: an integrated approach using CODEm. *Popul Health Metr*. (2012) 10:1. doi: [10.1186/1478-7954-10-1](https://doi.org/10.1186/1478-7954-10-1)

24. Mao Q, Zhu X, Zhang X, Kong Y. Effect of air pollution on the global burden of cardiovascular diseases and forecasting future trends of the related metrics: a systematic analysis from the Global Burden of Disease Study 2021. *Front Med*. (2024) 11:1472996. doi: [10.3389/fmed.2024.1472996](https://doi.org/10.3389/fmed.2024.1472996)

25. Lu H, Tan Z, Liu Z, Wang L, Wang Y, Suo C, et al. Spatiotemporal trends in stroke burden and mortality attributable to household air pollution from solid fuels in 204 countries and territories from 1990 to 2019. *Sci Total Environ*. (2021) 775:145839. doi: [10.1016/j.scitotenv.2021.145839](https://doi.org/10.1016/j.scitotenv.2021.145839)

26. Safiri S, Kolahi AA, Hoy D, Buchbinder R, Mansournia MA, Bettampadi D, et al. Global, regional, and national burden of neck pain in the general population, 1990–2017: systematic analysis of the Global Burden of Disease Study 2017. *BMJ*. (2020) 368:m791. doi: [10.1136/bmj.m791](https://doi.org/10.1136/bmj.m791)

27. Hu J, Ke R, Teixeira W, Dong Y, Ding R, Yang J, et al. Global, regional, and national burden of CKD due to glomerulonephritis from 1990 to 2019: a systematic analysis from the Global Burden of Disease Study 2019. *Clin J Am Soc Nephrol*. (2023) 18:60–71. doi: [10.2215/CJN.0000000000000017](https://doi.org/10.2215/CJN.0000000000000017)

28. Zhu B, Wang Y, Zhou W, Jin S, Shen Z, Zhang H, et al. Trend dynamics of gout prevalence among the Chinese population, 1990–2019: a joinpoint and age-periodcohort analysis. *Front Public Health*. (2022) 10:1008598. doi: [10.3389/](https://doi.org/10.3389/fpubh.2022.1008598) [fpubh.2022.1008598](https://doi.org/10.3389/fpubh.2022.1008598)

29. Cheng Z, Kong Y, Yang W, Xu H, Tang D, Zuo Y. Association between serum copper and blood glucose: a mediation analysis of inflammation indicators in the NHANES (2011–2016). *Front Public Health*. (2024) 12:1401347. doi: [10.3389/](https://doi.org/10.3389/fpubh.2024.1401347) [fpubh.2024.1401347](https://doi.org/10.3389/fpubh.2024.1401347)

30. Kong YZ, Xu HT, Li CY, Yang Y, Zhu XY, Zuo Y. Construction of PARI public health education programs for Chinese undergraduates: a Delphi study. *Front Public Health*. (2024) 12:1390011. doi: [10.3389/fpubh.2024.1390011](https://doi.org/10.3389/fpubh.2024.1390011)

31. Kong Y, Zhu X, Yang Y, Xu H, Ma L, Zuo Y. Knowledge, attitudes, practice, and public health education demand regarding PARI prevention: a cross-sectional study among Chinese undergraduates. *Front Public Health*. (2024) 12:1387789. doi: [10.3389/](https://doi.org/10.3389/fpubh.2024.1387789) [fpubh.2024.1387789](https://doi.org/10.3389/fpubh.2024.1387789)

32. Mao Q, Zhu X, Kong Y. Sleep duration mediates the association between heavy metals and the prevalence of depression: an integrated approach from the NHANES (2005–2020). *Front Psychiatry*. (2024) 15:1455896. doi: [10.3389/fpsyt.2024.1455896](https://doi.org/10.3389/fpsyt.2024.1455896)

33. Wang Q, Afshin A, Yakoob MY, Singh GM, Rehm CD, Khatibzadeh S, et al. Global burden of diseases nutrition and chronic diseases expert group (NutriCoDE). Impact of nonoptimal intakes of saturated, polyunsaturated, and trans fat on global burdens of coronary heart disease. *J Am Heart Assoc*. (2016) 5:e002891. doi: [10.1161/](https://doi.org/10.1161/JAHA.115.002891) [JAHA.115.002891,](https://doi.org/10.1161/JAHA.115.002891) Erratum in: J Am Heart Assoc. 2016; 5(1): e002076. doi:10.1161/ JAHA.115.002076. Erratum in: J Am Heart Assoc. 2017; 6(11): e002181

34. Hooper L, Summerbell CD, Thompson R, Sills D, Roberts FG, Moore H, et al. Reduced or modified dietary fat for preventing cardiovascular disease. *Cochrane Database Syst Rev*. (2011) 6:CD002137. doi: [10.1002/14651858.CD002137.pub2](https://doi.org/10.1002/14651858.CD002137.pub2)

35. Laaksonen DE, Nyyssönen K, Niskanen L, Rissanen TH, Salonen JT. Prediction of cardiovascular mortality in middle-aged men by dietary and serum linoleic and polyunsaturated fatty acids. *Arch Intern Med*. (2005) 165:193–9. doi: [10.1001/](https://doi.org/10.1001/archinte.165.2.193) [archinte.165.2.193](https://doi.org/10.1001/archinte.165.2.193)

36. Abdelhamid AS, Brown TJ, Brainard JS, Biswas P, Thorpe GC, Moore HJ, et al. Omega-3 fatty acids for the primary and secondary prevention of cardiovascular disease. *Cochrane Database Syst Rev*. (2018) 7:CD003177. doi: [10.1002/14651858.](https://doi.org/10.1002/14651858.CD003177.pub3) [CD003177.pub3](https://doi.org/10.1002/14651858.CD003177.pub3)

37. He K, Merchant A, Rimm EB, Rosner BA, Stampfer MJ, Willett WC, et al. Dietary fat intake and risk of stroke in male US healthcare professionals: 14 year prospective cohort study. *BMJ*. (2003) 327:777–82. doi: [10.1136/bmj.327.7418.777](https://doi.org/10.1136/bmj.327.7418.777)

38. Mensink RPWorld Health Organization. Effects of saturated fatty acids on serum lipids and lipoproteins: a systematic review and regression analysis. Geneva: World Health Organization (2016).

39. Nuesch E, Trelle S, Reichenbach S, Rutjes AWS, Tschannen B. Small study effects in meta-analyses of osteoarthritis trials: meta-epidemiological study. *BMJ*. (2010) 341:c3515. doi: [10.1136/bmj.c3515](https://doi.org/10.1136/bmj.c3515)

40. Sterne JA, Gavaghan D, Egger M. Publication and related bias in meta-analysis: power of statistical tests and prevalence in the literature. *J Clin Epidemiol*. (2000) 53:1119–29. doi: [10.1016/s0895-4356\(00\)00242-0](https://doi.org/10.1016/s0895-4356(00)00242-0)

41. Sahebkar A, Simental-Mendía LE, Mikhailidis DP, Pirro M, Banach M, Sirtori CR, et al. Effect of omega-3 supplements on plasma apolipoprotein C-III concentrations: a systematic review and meta-analysis of randomized controlled trials. *Ann Med*. (2018) 50:565–75. doi: [10.1080/07853890.2018.1511919](https://doi.org/10.1080/07853890.2018.1511919)

42. Patti AM, Al-Rasadi K, Giglio RV, Nikolic D, Mannina C, Castellino G, et al. Natural approaches in metabolic syndrome management. *Arch Med Sci*. (2018) 14:422–41. doi: [10.5114/aoms.2017.68717](https://doi.org/10.5114/aoms.2017.68717)

43. Mazidi M, Mikhailidis DP, Banach M. Omega-3 fatty acids and risk of cardiovascular disease: systematic review and meta-analysis of randomized controlled trials with 127,447 individuals and a Mendelian randomization study. *Circulation*. (2019) 140:e965–e1011. doi: [10.1001/2012.jama.11374](https://doi.org/10.1001/2012.jama.11374)

44. Dobrzyńska MA, Przysławski J. The effect of camelina oil (α-linolenic acid) and canola oil (oleic acid) on lipid profile, blood pressure, and anthropometric parameters in postmenopausal women. *Arch Med Sci*. (2020) 17:1566–74. doi: [10.5114/](https://doi.org/10.5114/aoms.2020.94033) [aoms.2020.94033](https://doi.org/10.5114/aoms.2020.94033)

45. Ruscica M, Penson PE, Ferri N, Sirtori CR, Pirro M, Mancini GBJ, et al. Impact of nutraceuticals on markers of systemic inflammation: potential relevance to cardiovascular diseases—a position paper from the international lipid expert panel (ILEP). *Prog Cardiovasc Dis*. (2021) 67:40–52. doi: [10.1016/j.pcad.2021.06.010](https://doi.org/10.1016/j.pcad.2021.06.010)

46. Banach M, Burchardt P, Chlebus K, Dobrowolski P, Dudek D, Dyrbuś K, et al. PoLA/CFPiP/PCS/PSLD/PSD/PSH guidelines on diagnosis and therapy of lipid disorders in Poland 2021. *Arch Med Sci*. (2021) 17:1447–547. doi: [10.5114/](https://doi.org/10.5114/aoms/141941) [aoms/141941](https://doi.org/10.5114/aoms/141941)