

AHA SCIENTIFIC STATEMENT

Forecasting the Burden of Cardiovascular Disease and Stroke in Women in the United States Through 2050: A Scientific Statement From the American Heart Association

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BACKGROUND: Forecasts for the future prevalence of cardiovascular disease and stroke are crucial to guide efforts to improve health outcomes across the life course for women.

METHODS: Using historical trends from the 2015 to 2020 National Health and Nutrition Examination Survey, 2015 to 2019 Medical Expenditure Panel Survey, and census estimates for population growth, we estimated trends in prevalence through 2050 for cardiovascular risk factors based on suboptimal levels of Life's Essential 8 and clinical cardiovascular disease and stroke, overall and by age and race and ethnicity.

RESULTS: Among adult women overall, the prevalence of hypertension is estimated to increase from 48.6% in 2020 to 59.1% in 2050. Diabetes (14.9% to 25.3%) and obesity (43.9% to 61.2%) will increase, whereas hypercholesterolemia will decline (42.1% to 22.3%). Prevalences of suboptimal diet, inadequate physical activity, and smoking will decline over time, and inadequate sleep will increase. Prevalences of coronary disease (6.85% to 8.21%), heart failure (2.45% to 3.60%), stroke (4.14% to 6.74%), atrial fibrillation (1.58% to 2.31%), and total cardiovascular disease and stroke (10.7% to 14.4%) will rise. Similar trends are projected in girls 2 to 19 years of age, with an increase from 19.6% to 32.0% projected in obesity. Most adverse trends are projected to be more pronounced among women and girls identifying as American Indian/Alaska Native or multiracial, Black, or Hispanic.

CONCLUSIONS: The prevalence of cardiovascular risk factors and disease in women and girls will increase over the next 30 years. Focused clinical and public health interventions are needed across the life course to address these adverse trends.

Key Words: AHA Scientific Statements ■ cardiovascular diseases ■ heart disease risk factors ■ population forecast ■ stroke

Cardiovascular disease (CVD) and stroke are the leading causes of morbidity and mortality for women in the United States¹ and globally.² In the United States between 2017 and 2020, >62 million women had some form of CVD (coronary heart disease [CHD], heart failure [HF], atrial fibrillation [AF], or stroke) or hypertension, with an overall prevalence of

almost 45%.³ In addition to the clinical burden of CVD, the economic burden is massive, with estimated direct and indirect spending in 2020 to 2021 of \$417.9 billion.⁴ Given that women make up just under half of the total incident and prevalent CVD cases in the United States and typically outlive their male counterparts,^{5,6} spending for women likely exceeds \$200 billion annually.

Supplemental Material is available at <https://www.ahajournals.org/doi/suppl/10.1161/CIR.0000000000001406>.

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Circulation is available at www.ahajournals.org/journal/circ

Although heart disease mortality declined precipitously from the 1970s to the early 2000s, it has since plateaued.⁷ Indeed, recent studies suggest worsening trends not only in premature mortality from cardiac disease⁸ but also in health factors and health behaviors that influence the development of CVD as contained in the American Heart Association's Life's Essential 8⁹: healthy blood pressure, body weight, blood sugar, and blood cholesterol; adequate sleep and physical activity; refraining from smoking; and a healthy diet.^{10,11} The biological and social underpinnings of CVD incidence and progression are different in women and men, and unique interventions may be needed to improve women's cardiovascular health.^{12–14} Furthermore, deep inequities associated with race and ethnicity remain in women's cardiovascular health outcomes. In the United States, Black, Hispanic, and American Indian/Alaska Native (AI/AN) women^{15–18} and subgroups of Asian Americans have higher rates of cardiovascular risk factors,¹⁹ adverse pregnancy outcomes, and incident and prevalent CVD and higher cardiovascular mortality.²⁰ Intersectionality, described as the compounding burden of co-occurring adverse social determinants of health (SDOH) such as racial and gender identity, poverty, low literacy, rural residence, and other psychosocial stressors,²¹ creates unique negative social experiences that contribute to poor cardiovascular outcomes.²² Intersectionality occurs more often in Black women, which contributes to the higher risk for incident CVD.^{23,24}

As the population continues to age and diversify,²⁵ projections for CVD in women will allow clinical leaders and policymakers to make strategic decisions about health care resources, future areas for research, and program development. Building on prior work by the American Heart Association projecting the future prevalence of disease^{5,6,26–29} and using national survey and epidemiological data, we aimed to estimate the prevalence and number of women with CVD risk factors and clinical disease from 2020 to 2050 for both women and girls and across different racial and ethnic populations. For the purposes of this document, we use “women” and “girls” to imply mostly sex-specific (rather than gender-specific) criteria.

METHODS

Baseline Prevalence

Detailed methods have been previously published.^{5,6} Briefly, we estimated the prevalence of suboptimal levels of each of the elements of American Heart Association's Life's Essential 8.⁹ We also estimated the prevalence of 4 diseases—CHD, HF, stroke, and AF—and a composite outcome henceforth referred to as total CVD that included any of these 4 diseases. We estimated the baseline prevalence of each outcome (except AF) using data from the 2015 to March 2020 prepandemic NHANES (National Health and Nutrition Examination Survey).³⁰ Lists of qualifying measures and questions used to define

each outcome for women and girls are given in [Supplemental Table 1a and 1b](#). We estimated the prevalence of AF using the 2015 to 2019 MEPS (Medical Expenditure Panel Survey).³¹

We estimated the prevalence of each outcome using logit regression models adjusted for demographics (age, sex, and race and ethnicity). We predicted the prevalence of each outcome in each age/sex/race and ethnicity cell using coefficients from the logit regressions for the 2017 to March 2020 NHANES cycle (or the 2019 MEPS for AF). Age groups for women were 20 to 44, 45 to 64, 65 to 79, and ≥80 years. Age groups for girls were 2 to 5, 6 to 11, and 12 to 19 years. Race and ethnicity categories were Hispanic, non-Hispanic Black, non-Hispanic White, non-Hispanic Asian, and combined AI/AN/multiracial based on NHANES categories. We generated bootstrapped SEs for the estimates of total CVD and stroke prevalence. We repeated the analyses of logit regression models, replacing the race and ethnicity stratifications with sex and age categories.

Prevalence Projections

We considered the baseline prevalence of each outcome from NHANES to represent prevalence for 2018. We used NHANES data from 2010 to 2020 to estimate growth or decline rates in age-adjusted prevalence of the risk factors and clinical CVD (except AF, see later discussion), and applied those rates to age/sex/race and ethnicity-specific prevalence for the years 2020 to 2050. The growth or decline rates were calculated overall rather than separately by sex. Growth and decline rates were estimated separately for adults and children given differences in variable definitions and cutoffs between the 2 populations. For adults, we did not estimate a growth or decline rate for inadequate sleep or poor diet because of insufficient historical data. For AF, we could not calculate a growth rate because of MEPS data limitations, so we rescaled the hypertension growth rate to the AF baseline prevalence given the relationship between these 2 clinical conditions and used that as the AF growth rate. For children, we did not estimate a growth or decline rate for hypertension, inadequate physical activity, tobacco use, or poor diet because of changes in definitions and limitations in data availability.

We combined estimates of prevalence by age/sex/race and ethnicity with the 2010 Census projections of population counts for 2020 through 2050 to generate the projected number of women and girls with each condition for 2020 through 2050. Projected population counts for 2020 through 2050 were obtained from the 2017 Population Projections of the US resident population by age, sex, race, and Hispanic origin generated by the US Census Bureau. Uncertainty was projected around the prevalence and counts estimates with a

bootstrapping approach. We ran 1000 iterations estimating the number of individuals with each condition in each year of the forecast, redrawing logit coefficients used to predict prevalence of the condition in each subgroup from a multivariate normal distribution, with the estimated coefficients as the mean and the estimated variance-covariance matrix as the variance. We also redrew linear regression coefficients that were used to calculate prevalence growth and decline rates using normal distributions. The lower and upper bounds of uncertainty intervals around estimated prevalence and count estimates were the 5th and 95th percentile, respectively, of the bootstrapped results. The 90% credible ranges for the results were generated with these boundaries. Statistical analyses were conducted with Stata version 17.0 (StataCorp LLC, 2021).

Analyses were conducted by the Research Triangle Institute International under a contract from the American Heart Association. The study was exempt from Institutional Review Board review because of the deidentified nature of the data.³²

RESULTS

Prevalence and Number of Adult Women With Cardiovascular Risk Factors and CVD

First, adverse levels of Life's Essential 8 were estimated among adult women. Figure 1A demonstrates adverse

levels of the health factors over time. According to current trends, the prevalence of hypertension will increase from 48.6% in 2020 to 59.1% in 2050. Increases were also projected for diabetes (14.9% to 25.3%) and obesity (43.9% to 61.2%). There was a projected decline in hypercholesterolemia (42.1% to 22.3%). Figure 1B shows levels of suboptimal health behaviors over time. Suboptimal diet (49.0% to 47.6%), inadequate physical activity (39.8% to 28.6%), and current smoking (13.8% to 7.2%) were estimated to improve over time, whereas inadequate sleep (40.3% to 42.2%) was projected to worsen. Figure 1C shows overall rates of CVD, with the prevalence of CHD (6.85% to 8.21%), HF (2.45% to 3.60%), stroke (4.14% to 6.74%), AF (1.58% to 2.31%), and total CVD (10.7% to 14.4%) expected to rise. Results for each factor and disease, along with the number of adult women affected and accompanying uncertainty intervals, are given in Supplemental Tables 2 and 3.

Prevalence and Number of Women With Cardiovascular Risk Factors and CVD, by Demographic Group

Figure 2 shows the projected prevalence for health behaviors by racial, ethnic, and age group. Black women (olive green bars) had the highest baseline and projected prevalence of inadequate sleep and poor diet; Asian women (pink bars) had the highest projected prevalence



Figure 1. Proportion of US adult women with adverse levels of cardiovascular health factors, health behaviors, and CVD and stroke, 2020 to 2050.

For adults, we did not estimate a growth or decline rate for inadequate sleep or poor diet because of insufficient historical data. CVD indicates cardiovascular disease.

of inadequate physical activity; and the aggregated group of AI/AN/multiracial women (blue bars) had the highest projected prevalence of smoking. The youngest women (20–44 years of age, pink bars) had the highest prevalence of poor diet and smoking but the lowest prevalence of inadequate physical activity. Poor sleep was particularly prevalent among the oldest women (≥80 years of age, purple bars), but smoking was the least prevalent.

Figure 3 shows the projected prevalence for health factors by racial, ethnic, and age group. Black women had the highest prevalence of hypertension, diabetes, and obesity at baseline but the lowest prevalence of hypercholesterolemia. Projected absolute increases in prevalence were generally greatest in Hispanic women (turquoise bars), although growth was also high in Asian women. The oldest women had the lowest baseline prevalence of obesity but the highest prevalence of hypertension and diabetes. The youngest women had the lowest prevalence of hypertension and diabetes but were projected to have the greatest increase in these conditions.

Figure 4 shows the projected prevalence for clinical CVD by racial, ethnic, and age group. Black women and AI/AN/multiracial women had the highest prevalence of total CVD plus hypertension, whereas all groups, particularly Hispanic and Asian women, had significant projected growth in this measure. Black women had the highest rates of HF and stroke and the greatest projected increases in prevalence; White women had the highest rates of AF; and AI/AN/multiracial women had the highest rate of CHD. In terms of age differences, the prevalence of CHD, stroke, HF, AF, and total CVD was markedly higher among the oldest women. The greatest projected increase, however, was among the youngest women, particularly for stroke and total CVD.

Full results for each factor and condition by age are given in Supplemental Table 4 (prevalence) and Supplemental Table 5 (number). Full results for each factor and condition by race and ethnicity are given in Supplemental Table 6 (prevalence) and Supplemental Table 7 (number).

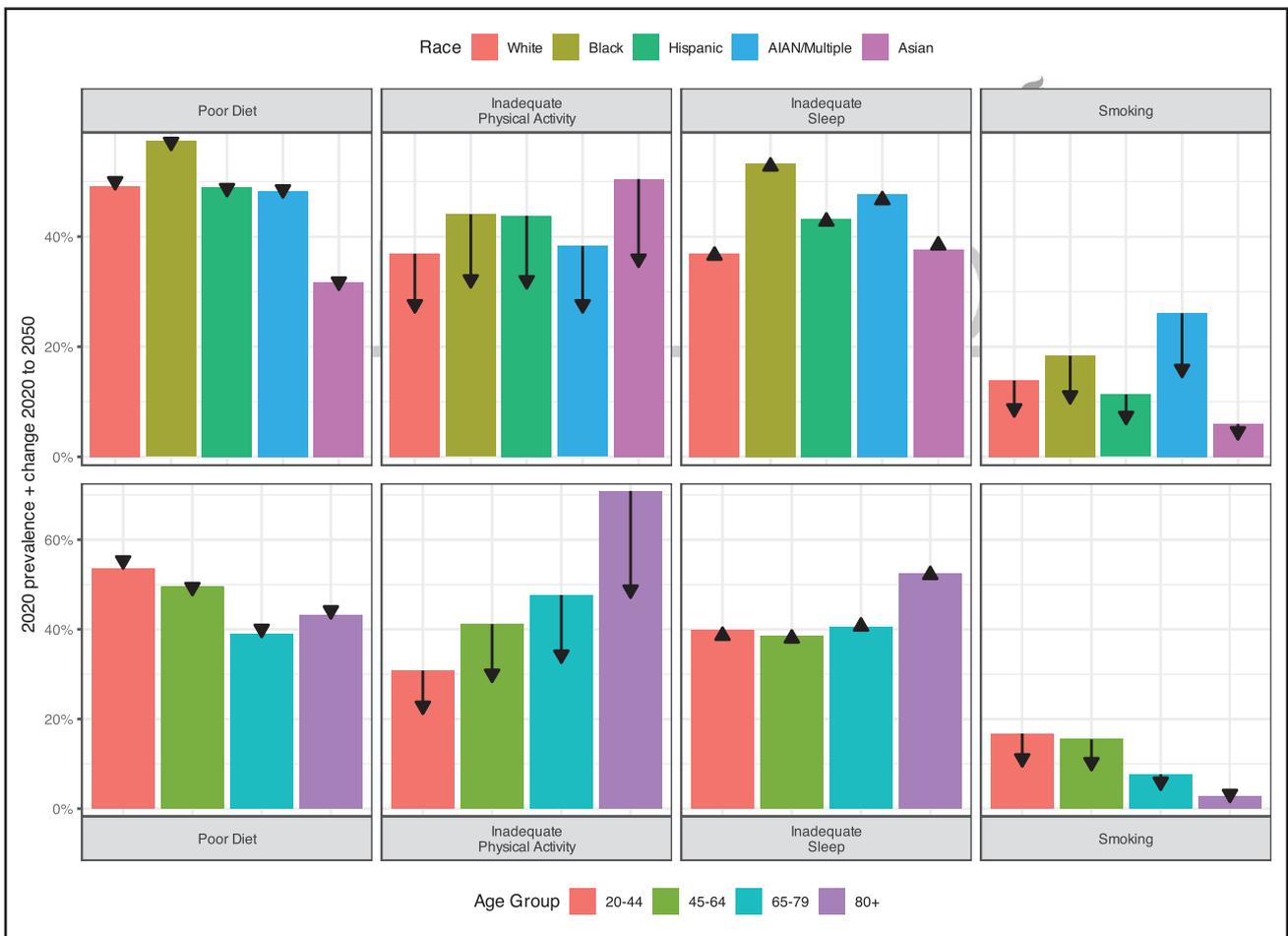


Figure 2. Trends in prevalence of adverse levels of cardiovascular health behaviors in US women, by race, ethnicity, and age. Bars represent the baseline estimate for 2020 for each factor; arrows represent the change from 2020 to 2050 and end at the 2050 estimate. For adults, we did not estimate a growth or decline rate for inadequate sleep or poor diet because of insufficient historical data, so the markers are centered around the 2020 baseline. AIAN indicates American Indian/Alaska Native.

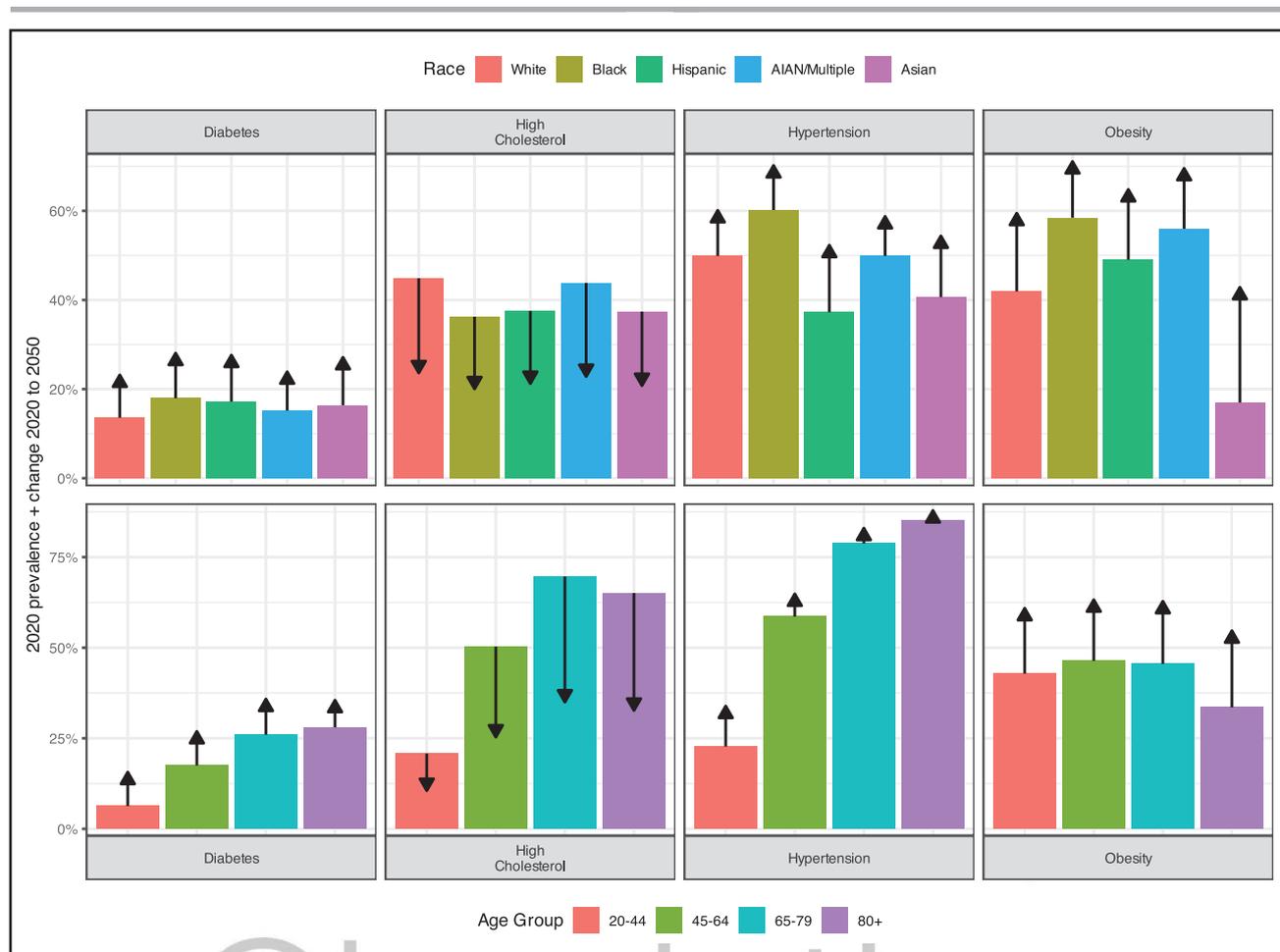


Figure 3. Trends in prevalence of adverse cardiovascular health factors in US women by race, ethnicity, and age. Bars represent the baseline estimate for 2020 for each factor; arrows represent the change from 2020 to 2050 and end at the 2050 estimate. AIAN indicates American Indian/Alaska Native.

Prevalence and Number of Girls With Cardiovascular Risk Factors, Overall and by Race and Ethnicity

Figure 5 shows projected prevalence for the adverse health factors and behaviors among girls. The prevalences of hypertension (3.3% to 3.2%) and diabetes (1.1% to 1.2%) are estimated to be stable from 2020 to 2050. A significant increase was projected for obesity (19.6% to 32.0%) and a decline for hypercholesterolemia (7.1% to 3.2%). The prevalences of poor diet (56.3% to 55.6%) and inadequate physical activity (63.4% to 63.9%) were high but stable; current smoking (7.3% to 7.0%) was also estimated to be stable. Sleep data are not available for girls. Full results for each factor are given in Supplemental Table 8 (prevalence) and Supplemental Table 9 (number).

Figure 6 disaggregates these projections by racial and ethnic group among girls. Black girls are projected to have the highest prevalence of hypertension and diabetes, whereas Hispanic girls are projected to have the

highest prevalence of obesity. Asian girls are projected to have the highest prevalence of hypercholesterolemia, although the prevalence in all groups is projected to decline over time. Inadequate physical activity is high for all racial and ethnic groups but is projected to be highest in Hispanic girls. Poor diet is projected to be highest in Black girls, whereas tobacco use is projected to be highest in AI/AN/multiracial girls. Full results for each factor are given in Supplemental Table 10 (prevalence) and Supplemental Table 11 (number).

DISCUSSION

The forecast for cardiovascular health in women and girls through 2050 supports a need for sustained action. The most efficient, most effective, and least costly way to reduce the prevalence and impact of CVD is through prevention.^{33,34} These projections signal that current prevention efforts are inadequate, particularly in women of color and younger women. Unique risk factors that pertain to women such as reproductive factors

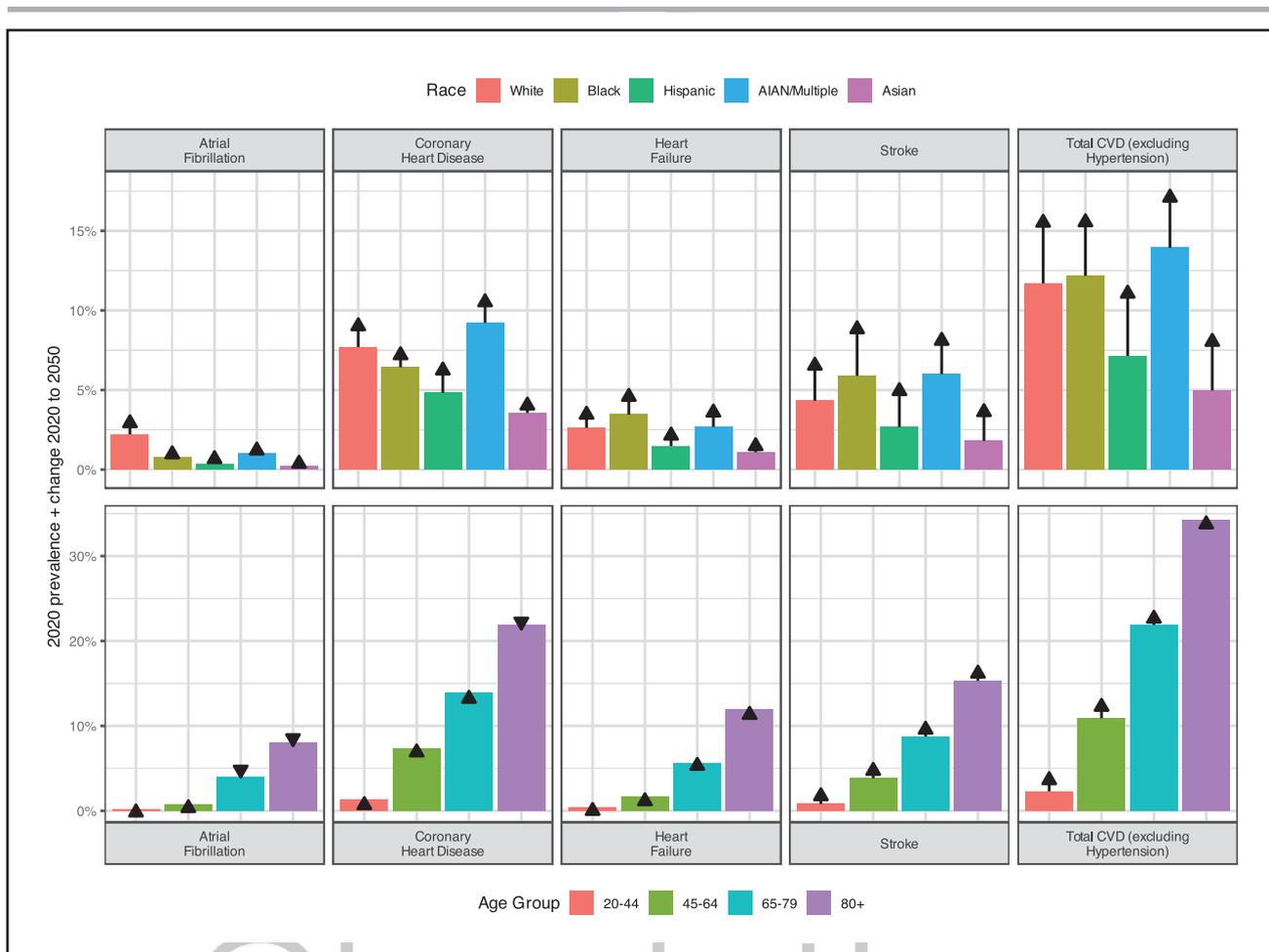


Figure 4. Trends in prevalence of clinical cardiovascular disease in US women by race, ethnicity, and age. Bars represent the baseline estimate for 2020 for each factor; arrows represent the change from 2020 to 2050 and end at the 2050 estimate. AIAN indicates American Indian/Alaska Native; and CVD, cardiovascular disease.

across the life span must be considered in prevention and care delivery.³⁵ Here, several considerations are outlined, encompassing prevention, treatment, and sustained care across the life course and for women of all racial and ethnic groups, with a focus on how these issues will affect women and girls in the coming 3 decades and beyond.

Health Behaviors

The prevention of risk factors and disease before they occur is crucial for any strategy to reverse worsening CVD trends, and health behaviors are key to this approach.³⁶ Women who reach midlife with optimal physical activity, dietary, and sleep patterns who have never smoked will have low rates of CVD in the decades that follow.³⁷⁻³⁹ Evidence-based public health efforts to promote and sustain optimal cardiovascular health behaviors in women are thus essential.⁴⁰ Environmental and physiological drivers⁴¹ of worsening sleep trends in women such as screen time, stress, and hormonal changes must be identified and miti-

gated. Early efforts to identify and support behavior change in girls and women with suboptimal cardiovascular health behaviors should occur in schools,⁴² community organizations,⁴⁰ and pediatric⁴³ and gynecology⁴⁴ offices. Digital engagement strategies will increasingly be part of the landscape for primordial and primary prevention.⁴⁵

Considerations: (1), Focus on supporting healthy behaviors in schools, community organizations, and pediatric and gynecology offices. (2), Encourage digital engagement strategies when appropriate to support positive behavior change.

Health Factor Management: Hypertension, Diabetes, Obesity, and Cholesterol

Tailored and sustained interventions for chronic disease are needed among high-risk populations earlier in the course of disease.⁴⁶⁻⁵⁰ Guidelines already recommend blood pressure screening for adults and children,⁵¹ and observational studies suggest the potential for initiating

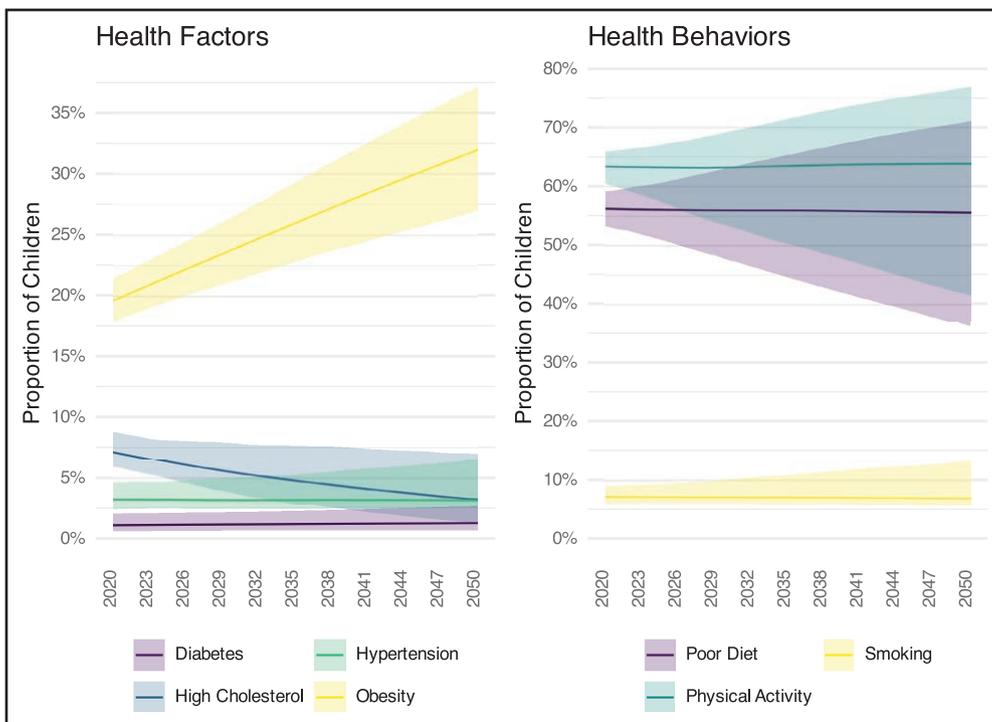


Figure 5. Prevalence of adverse levels of cardiovascular health behaviors and health factors in US girls, 2020 and 2050, overall. For girls, we did not estimate clinical cardiovascular disease because of low prevalence. Girls within the specified age group were included in the corresponding categories: inadequate physical activity, obesity, and poor diet, 2 to 19 years; high cholesterol, 6 to 19 years; hypertension, 8 to 19 years; and diabetes and current tobacco use, 12 to 19 years.



therapy for stage 1 hypertension in appropriate individuals to disproportionately benefit women for stroke and HF risk reduction.^{52–54} However, there was no interaction between sex and treatment effect in blood pressure-lowering trials such as SPRINT (Systolic Blood Pressure Intervention Trial)⁵⁵ and ACCORD (Action to Control Cardiovascular Risk in Diabetes).⁵⁶ Given that only a minority of patients have their blood pressure controlled, there is massive opportunity for improving outcomes simply through better management of this chronic disease. Team-based care and digitally enabled monitoring and medication titration have shown good evidence of success and could be pursued much more broadly.

There is also a need for better chronic disease management in diabetes. Diabetes is a particularly potent risk factor for women, conferring excess risk compared with men for a range of cardiovascular conditions, including CHD, HF, and stroke.^{57–59} Women also experience greater cardiovascular risk associated with undiagnosed diabetes and prediabetes than men and derive similar benefit from treatment as men.^{60,61} The recently developed Predicting Risk of Cardiovascular Disease Events calculator highlights the growing focus on cardio-kidney-metabolic health, which includes diabetes, obesity, and chronic kidney disease, and its role in the progression of CVD.^{62,63} Similar to hypertension, team-based, digitally enabled care has great potential to improve disease management.

Obesity is a chronic disease that affects women to a greater degree than men, particularly at the most severe levels,⁶⁴ and the treatment landscape for obesity is rapidly changing. Given the recent development of highly effective medications to treat obesity, in particular the glucagon-like peptide receptor agonists, and a growing understanding of the hormonal and genetic underpinnings of obesity, it is possible that the coming decades could see meaningful changes in the prevalence of obesity. However, much remains to be learned about the long-term effects of these medications, about whether their safety or efficacy differs by sex,⁶⁵ about the most effective ways to use them in concert with changes in diet and physical activity, and about their use in childhood and adolescence. Furthermore, issues of cost, access, and tolerability may affect the population impact of these medications and should be studied.

The projected decline in the prevalence of hypercholesterolemia in women may be surprising but has been demonstrated in multiple prior studies.^{3,66–69} Increased screening for hypercholesterolemia and a shift toward risk-based approaches instead of targeting specific lipid levels may account for continued decline in cholesterol levels nationally.⁷⁰ Despite these declines, ongoing attention should be paid to ensuring that women are equitably treated with statins and other lipid-lowering therapies, particularly because treatment benefits women and men

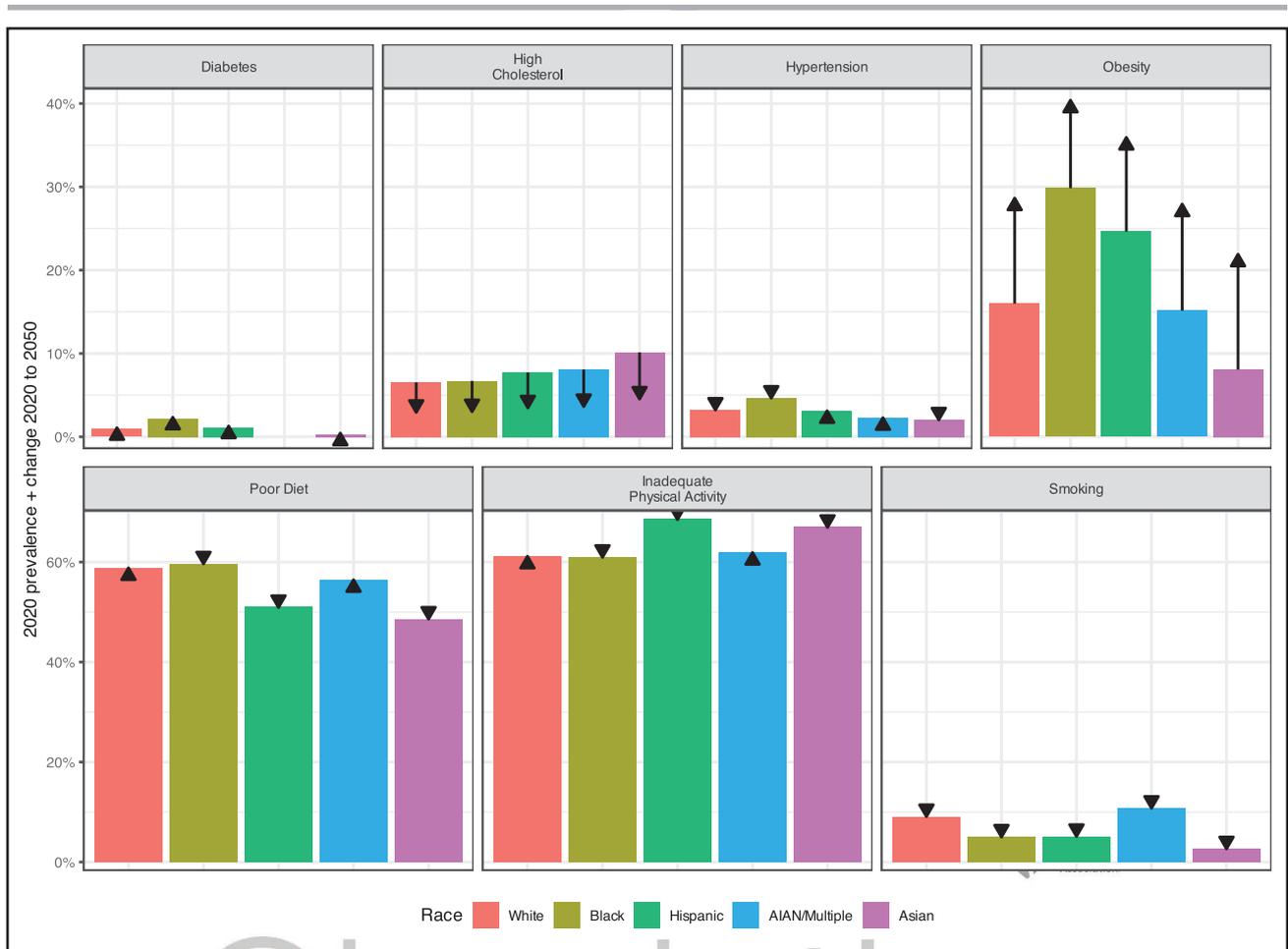


Figure 6. Prevalence of adverse levels of cardiovascular health behaviors and health factors in U.S. girls, 2020 and 2050, by racial or ethnic group.

Bars represent the baseline estimate for 2020 for each factor; arrows represent the change from 2020 to 2050 and end at the 2050 estimate. For girls, we did not estimate clinical cardiovascular disease because of low prevalence. Girls within the specified age group were included in the corresponding categories: inadequate physical activity, obesity, and poor diet, 2 to 19 years; high cholesterol, 6 to 19 years; hypertension, 8 to 19 years; and diabetes and current tobacco use, 12 to 19 years. AIAN indicates American Indian/Alaska Native.

to a similar degree.⁷¹ Furthermore, the decline in hypercholesterolemia in concert with a rise in obesity, other CVD risk factors, and CVD overall may relate to a shift in American diets toward highly processed, high-sugar foods and suggests a need for more attention in this area.

Considerations: (1), Chronic disease management should be a focus of ongoing clinical and policy efforts toward improving women's health, including supporting early intervention, team-based care, and digitally enabled care. (2), The sex-specific effects and safety of emerging obesity therapies should be studied.

Clinical CVD

Coronary Heart Disease

In association with adverse trends in risk factors, the prevalence of CHD is projected to rise in every age cat-

egory and race and ethnicity group through 2050. Historically, women have had lower rates of revascularization and received less evidence-based care for CHD events than men.^{72,73} Although these gaps have narrowed, they remain particularly salient for younger women.⁸ However, attention needs to remain on optimizing care for women experiencing acute myocardial infarction and other forms of CHD to ensure high-quality treatment for all.

Heart Failure

The rising prevalence of HF in women reflects the combined impact of an increasing burden of risk factors and the rising prevalences of CHD and AF. Risk factors such as diabetes, hypertension, and obesity are more likely to lead to HF among women than men.⁷⁴ Women with HF are more likely than men with HF to have preserved left ventricular ejection fraction, a form of HF with fewer treatment options.⁷⁵ New, possibly sex-specific approaches to prevention, diagnosis, and

management will be needed to limit the burden of HF for women, particularly among those with preserved ejection fraction.⁷⁴ Clinical trials in women of medical therapies for the management of HF for preserved ejection fraction are sorely needed and should include attention to potential heterogeneity of treatment effect for women compared with men and to different rates of adverse drug events and medication discontinuation for women compared with men.

Stroke

Women have a higher burden of stroke than men, are less likely to receive treatments such as thrombolysis and thrombectomy, and are more likely to have poor outcomes than men.^{76,77} In addition, stroke in midlife has profound economic implications with regard to health care use and lowered quality of life, as well as implications for caregiving, particularly among women.^{27,78} Drivers of increased stroke incidence in younger women may include the concomitant increased prevalence of hypertension in this age group and increases in hypertensive disorders of pregnancy,⁷⁹ which are associated with short- and long-term risk of stroke.⁸⁰ In addition to prevention, as discussed previously, it is important to ensure that women experiencing strokes have adequate access to thrombolysis, thrombectomy, and other interventions.

Atrial Fibrillation

Women are at higher risk of stroke with AF than men⁸¹ yet receive less anticoagulation⁸² and less rhythm control⁸³ than men, which can lead to poor outcomes, including both stroke and HF. The increasing prevalence of AF is related in part to increases in hypertension and obesity; thus, management of chronic disease is important for preventing AF. Further interventions are needed to improve rates of use of anticoagulation when indicated, and shared decision-making should incorporate sex-specific risk factors.

Brain Health

The increasing prevalence of cardiovascular risk factors and clinical CVD, particularly hypertension and stroke, has major implications for brain health in women. Strokes cause cognitive impairment and dementia directly but can also signal underlying cerebral small-vessel disease, a contributor to both vascular dementia and Alzheimer disease.⁸⁴ Conversely, optimizing cardiovascular health through hypertension control, exercise, and diet effectively supports brain health by reducing the risk of mild cognitive impairment.^{85–88} Sex differences exist for dementia risk,⁸⁹ and tailored strategies for maintaining optimal brain health in women should be investigated.

Considerations: (1), For the acute presentation of illness for myocardial infarction and stroke, quality improvement programs are needed to ensure that women receive high-quality care and adequate treatment. (2), For chronic disease management in AF and HF, new approaches to management may be needed that incorporate sex-specific factors into shared decision-making and treatment. (3), Brain health, particularly dementia, should be considered an important part of CVD prevention and treatment in women.

Cardiovascular Risk Throughout the Reproductive Life Span: Opportunities for Intervention

Puberty, Menarche, and Adolescence

Understanding risk factor development in girls is crucial, and markers of differential risk appear early. Menarche before 12 years of age is associated with increased risk for CVD⁹⁰ and is mediated by higher risks for obesity, hypertension, and impaired glucose metabolism as women age.^{91,92} The median age at menarche continues to decrease in the United States and at 11.9 years⁹³ may portend worsening CVD trends beyond 2050. Women with irregular or absent menstrual cycles in adolescence and young adulthood also have higher risks of CVD, even after adjustment for other CVD risk factors.⁹⁴ These findings reinforce the importance of menstrual history assessment for cardiovascular health promotion.

Women of Reproductive Age and Pregnancy

An especially concerning finding of our study is the projected rise in many adverse health behaviors and factors in women of reproductive age. This is a life stage during which women are more likely than men to attend primary care visits, leaving ample opportunity for risk factor detection and control.⁹⁵ Intervention in the reproductive years through multidisciplinary collaboration between obstetrician-gynecologists, internists, and subspecialists should improve the screening and treatment of diabetes, lipid abnormalities, hypertension, mental health issues, and subclinical CVD.^{96,97} Coronary calcium scoring could further risk stratify women in this age group, who are often overlooked for cardiovascular screening but who have particularly poor outcomes when cardiac events do occur.^{98,99}

Prepregnancy cardiovascular health affects pregnancy and obstetric outcomes.^{100–102} In addition, suboptimal maternal cardiovascular health, including excessive weight gain during pregnancy, has intergenerational impact with short-term and long-term effects on the offspring such as macrosomia, congenital abnormalities, obesity and excessive weight gain, hypertension, impaired glucose tolerance, and increased risk of future

CVD.^{103–105} Hypertension screening, diagnosis, and control during pregnancy is an increasingly important issue, with a need for continued codification of postpartum hypertension management as more evidence emerges about the short- and long-term consequences of the hypertensive diseases of pregnancy, including coronary calcification and HF.^{106,107} There is a need to focus on addressing adverse SDOH, improving health literacy, reducing barriers to attaining adequate physical activity, providing access to health care and food, and reducing psychological stressors in girls and women of reproductive age to mitigate these concerns.¹⁰⁸

Menopause

Individuals entering the menopause transition (perimenopause and menopause, 45–64 years of age) also demonstrated adverse trends in disease prevalence. Over the past 2 decades, longitudinal studies of women going through menopause^{109–113} have helped differentiate the influence of chronological aging from ovarian aging on cardiovascular health and identified adverse shifts in body composition, lipids and lipoproteins, and vascular function accompanying menopause.^{110–119} In addition to the pronounced sex hormone changes, vasomotor symptoms, sleep problems, and adverse changes in mental health have implications for women's cardiovascular health.^{120–123} Although multiple adverse changes in cardiometabolic health parameters occur simultaneously during menopause, clinical trials incorporating multiple interventions focused on symptoms or lifestyle measures to reverse risk of CVD are lacking, and the best timing of lifestyle interventions remains unclear. The impact of hormone therapy on risk also remains unclear, with newer hormone formulations requiring investigation, especially in women with vasomotor symptoms and low risk.¹⁰⁹

Considerations: (1), Pediatricians should be aware that early menarche may carry adverse prognostic information, and menstrual history should be considered part of a cardiovascular risk assessment. (2), Multidisciplinary collaborations in women's health should allow integrated care for prepregnancy, peripartum, and postpartum care across disciplines. (3), Studies should assess lifestyle interventions and hormone therapy in terms of their effect on CVD risk and outcomes around the time of menopause.

Considerations for Demographic Determinants of Health and SDOH

Race and Ethnicity

Specific attention should be paid to differences in cardiovascular risk factors and outcomes in different racial and ethnic groups. Black girls and Black women have more cardiovascular risk factors; are less likely to have access to care, to be diagnosed, and to receive treatment; and

have higher rates of morbidity and mortality compared with their White counterparts.^{124,125} Despite decades of knowledge of these health inequities, major gaps remain, suggesting that we need broad interventions across the life span to address inequities in cardiovascular risk factors in girls and women.

Social Determinants of Health

SDOH are recognized as important drivers of health outcomes and represent crucial targets for cardiovascular health interventions.^{126–128} Women disproportionately bear the burden of poverty in the United States and are more likely to delay or forego health care because of costs.¹²⁹ Independent contributors to increased HF risk in Black women include a lower household income,¹³⁰ whereas residing in a rural area is a risk factor for both Black women and White women.¹³¹ Among adults <75 years of age, greater SDOH burden is associated with higher stroke risk.¹³² To mitigate the future risk of CVD in women, it will be essential to implement targeted interventions that address both traditional risk factors and SDOH.¹²⁷

Considerations: (1), Specific programs should be developed that acknowledge and address the elevated risk of CVD in Black women. (2), Attention should be paid to the intersection of SDOH and medical risk factors, and setting specific interventions should be developed to improve cardiovascular health.

Broader Policy and Public Health Implications

This sobering forecast of CVD risk factors and disease in women is consistent with other forecasting efforts,^{133,134} albeit not sex-specific ones, and has enormous public health and societal implications. These findings are combined with an overall decreased awareness that CHD is the number 1 killer of women, especially among the women at highest risk.¹³⁵ It is important to note that these trends related to modifiable risk factors are not set in stone: Simulation studies suggest that incident CVD and stroke events, including death, could be reduced by 17% to 23% by reducing the prevalence of health factors such as hypertension, hypercholesterolemia, diabetes, and obesity by 10%.⁵ Strategies for screening in women across the life span, including attention to both traditional and female-specific risk factors, can alert clinicians to these enhanced risks, increase awareness in both women and their clinicians, and allow early implementation of interventions to improve the health factors and behaviors that are identified.⁸⁰ The *Lancet* Women and CVD Commission recommendations for reducing the global burden of CVD by 2030 support targeting public health strategies that increase awareness of CVD in women; targeting established, sex-specific, and

underrecognized risk factors; and strengthening health-care systems and engaging health-care professionals about these risks.¹³⁶ Reversing the trends of CVD in women would positively affect families, the workplace, and society as a whole.

Limitations and Knowledge Gaps

In addition to methodological limitations as published,⁵ we acknowledge that this analysis does not include prevalence estimates for other cardiovascular conditions with known sex differences in prevalence and outcomes, including valvular heart disease and specific cardiomyopathies. Cardiac arrhythmias (excluding AF) and conduction disorders are not addressed here. Growth and decline factors were not calculated separately by sex. Thus, differences in projections between males and females reflect differences in prevalence and demographics alone; trend differences by sex were not examined in this study. We were not able to incorporate sex-specific risk factors into this analysis. Race and ethnicity are social constructs and do not necessarily imply biological differences. These broad categories, however, offer valuable insights into social factors such as racial identity and lived experiences that influence cardiovascular health. Results presented here do not account for other SDOH at the individual (educational attainment, income, employment), interpersonal, community, or societal level.¹²⁸ The interplay between gender and sex and issues specific to the transgender community were not included in our modeling because of data limitations and represent areas of important future work. Our projections did not account for the COVID-19 pandemic or the potential impact of new pharmaceuticals or procedures. These projections should be updated at regular intervals to capture evolving changes in the treatment landscape.

Conclusions

The high burden of CVD in women in the United States is a major problem that will continue to grow over the

next 25 years, fueled by an increasing prevalence of multiple risk factors in women across the life span in every demographic group. Sex-specific studies, including clinical trials, would be beneficial to improve the understanding of the causes, manifestations, treatments, and outcomes in women. Public health interventions are needed to limit the impact of CVD on women, along with broad innovations in data collection, screening, diagnosis, and treatment that can be effectively applied to women in all areas of the country, across life stages, and with a diversity of life experiences.

ARTICLE INFORMATION

The American Heart Association makes every effort to avoid any actual or potential conflicts of interest that may arise as a result of an outside relationship or a personal, professional, or business interest of a member of the writing panel. Specifically, all members of the writing group are required to complete and submit a Disclosure Questionnaire showing all such relationships that might be perceived as real or potential conflicts of interest.

This statement was approved by the American Heart Association Science Advisory and Coordinating Committee on October 13, 2025, and the American Heart Association Executive Committee on December 8, 2025. A copy of the document is available at <https://professional.heart.org/statements> by using either "Search for Guidelines & Statements" or the "Browse by Topic" area. To purchase additional reprints, call 215-356-2721 or email Meredith.Edelman@wolterskluwer.com

The American Heart Association requests that this document be cited as follows: Joynt Maddox KE, Reynolds HR, Adedinsowo D, Bushnell C, DeVon HA, Gooding HC, Howard VJ, Mauricio R, Miller EC, Sharma G, Waken RJ; on behalf of the American Heart Association Women's Health Science Committee of the Council on Clinical Cardiology and Stroke Council; Council on Basic Cardiovascular Sciences; Council on Cardiovascular and Stroke Nursing; Council on Lifelong Congenital Heart Disease and Heart Health in the Young; and Council on Peripheral Vascular Disease. Forecasting the burden of cardiovascular disease and stroke in the United States through 2050 in women: a scientific statement from the American Heart Association. *Circulation*. 2026;153:e00000000001406. doi: 10.1161/CIR.0000000000001406

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Disclosures

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This table represents the relationships of writing group members that may be perceived as actual or reasonably perceived conflicts of interest as reported on the Disclosure Questionnaire, which all members of the writing group are required to complete and submit. A relationship is considered to be "significant" if (a) the person receives \$5000 or more during any 12-month period, or 5% or more of the person's gross income; or (b) the person owns 5% or more of the voting stock or share of the entity, or owns \$5000 or more of the fair market value of the entity. A relationship is considered to be "modest" if it is less than "significant" under the preceding definition. †Significant.

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This table represents the relationships of reviewers that may be perceived as actual or reasonably perceived conflicts of interest as reported on the Disclosure Questionnaire, which all reviewers are required to complete and submit. A relationship is considered to be "significant" if (a) the person receives \$5000 or more during any 12-month period, or 5% or more of the person's gross income; or (b) the person owns 5% or more of the voting stock or share of the entity, or owns \$5000 or more of the fair market value of the entity. A relationship is considered to be "modest" if it is less than "significant" under the preceding definition.

*Modest.

†Significant.

REFERENCES

- Ahmad FB, Cisewski JA, Anderson RN. Leading causes of death in the US, 2019–2023. *JAMA*. 2024;332:957–958. doi: 10.1001/jama.2024.15563
- World Health Organization. Global health estimates: life expectancy and leading causes of death and disability. Accessed June 27, 2025. <https://who.int/data/gho/data/themes/mortality-and-global-health-estimates>
- Martin SS, Aday AW, Allen NB, Almarzoq ZI, Anderson CAM, Arora P, Avery CL, Baker-Smith CM, Bansal N, Beaton AZ, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention Statistics Committee and Stroke Statistics Committee. 2025 Heart disease and stroke statistics: a report of US and global data from the American Heart Association [published correction appears in *Circulation*. 2025;151:e1096]. *Circulation*. 2025;151:e41–e660. doi: 10.1161/CIR.0000000000001303
- Martin SS, Aday AW, Almarzoq ZI, Anderson CAM, Arora P, Avery CL, Baker-Smith CM, Barone Gibbs B, Beaton AZ, Boehme AK, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention Statistics Committee and Stroke Statistics Subcommittee. 2024 Heart disease and stroke statistics: a report of US and global data from the American Heart Association [published corrections appear in *Circulation*. 2024;149:e1164 and *Circulation*. 2025;151:e1095]. *Circulation*. 2024;149:e347–e913. doi: 10.1161/CIR.0000000000001209
- Joynt Maddox KE, Elkind MSV, Aparicio HJ, Commodore-Mensah Y, de Ferranti SD, Dowd WN, Hernandez AF, Khavjou O, Michos ED, Palaniappan L; on behalf of the American Heart Association. Forecasting the burden of cardiovascular disease and stroke in the United States through 2050: prevalence of risk factors and disease: a presidential advisory from the American Heart Association. *Circulation*. 2024;150:e65–e88. doi: 10.1161/cir.0000000000001256
- Kazi DS, Elkind MSV, Deutsch A, Dowd WN, Heidenreich P, Khavjou O, Mark D, Mussolino ME, Ovbiagele B, Patel SS, et al; on behalf of the American Heart Association. Forecasting the economic burden of cardiovascular disease and stroke in the United States through 2050: a presidential advisory from the American Heart Association. *Circulation*. 2024;150:e89–e101. doi: 10.1161/CIR.0000000000001258
- King SJ, Wangdak Yuthok TY, Bacong AM, Khandelwal A, Kazi DS, Mussolino ME, Wong SS, Martin SS, Lewis EF, Rodriguez F, et al. Heart disease mortality in the United States, 1970 to 2022. *J Am Heart Assoc*. 2025;14:e038644. doi: 10.1161/JAHA.124.038644
- Gonuguntla K, Badu I, Duhan S, Sandhyavenu H, Chobufo MD, Taha A, Thyagaturu H, Sattar Y, Keisham B, Ali S, et al. Sex and racial disparities in proportionate mortality of premature myocardial infarction in the United States: 1999 to 2020. *J Am Heart Assoc*. 2024;13:e033515. doi: 10.1161/jaha.123.033515
- Lloyd-Jones DM, Allen NB, Anderson CAM, Black T, Brewer LC, Foraker RE, Grandner MA, Lavretsky H, Perak AM, Sharma G, et al; on behalf of the American Heart Association. Life's Essential 8: updating and enhancing the American Heart Association's construct of cardiovascular health: a presidential advisory from the American Heart Association. *Circulation*. 2022;146:e18–e43. doi: 10.1161/CIR.0000000000001078
- Liu M, Aggarwal R, Zheng Z, Yeh RW, Kazi DS, Joynt Maddox KE, Wadhwa RK. Cardiovascular health of middle-aged U.S. adults by income level, 1999 to March 2020: a serial cross-sectional study. *Ann Intern Med*. 2023;176:1595–1605. doi: 10.7326/m23-2109
- Aggarwal R, Yeh RW, Joynt Maddox KE, Wadhwa RK. Cardiovascular risk factor prevalence, treatment, and control in US adults aged 20 to 44 years, 2009 to March 2020. *JAMA*. 2023;329:899–909. doi: 10.1001/jama.2023.2307
- Bairey Merz CN, Shaw LJ, Reis SE, Bittner V, Kelsey SF, Olson M, Johnson BD, Pepine CJ, Mankad S, Sharaf BL, et al; WISE Investigators. Insights from the NHLBI-sponsored Women's Ischemia Syndrome Evaluation (WISE) study, part II: gender differences in presentation, diagnosis, and outcome with regard to gender-based pathophysiology of atherosclerosis and macrovascular and microvascular coronary disease. *J Am Coll Cardiol*. 2006;47:S21–S29. doi: 10.1016/j.jacc.2004.12.084
- Shaw LJ, Bairey Merz CN, Pepine CJ, Reis SE, Bittner V, Kelsey SF, Olson M, Johnson BD, Mankad S, Sharaf BL, et al; WISE Investigators. Insights from the NHLBI-sponsored Women's Ischemia Syndrome Evaluation (WISE) study, part I: gender differences in traditional and novel risk factors, symptom evaluation, and gender-optimized diagnostic strategies. *J Am Coll Cardiol*. 2006;47:S4–S20. doi: 10.1016/j.jacc.2005.01.072
- Mehta LS, Velarde GP, Lewey J, Sharma G, Bond RM, Navas-Acien A, Fretts AM, Magwood GS, Yang E, Blumenthal RS, et al; on behalf of the American Heart Association Cardiovascular Disease and Stroke in Women and Underrepresented Populations Committee of the Council on Clinical Cardiology; Council on Cardiovascular and Stroke Nursing; Council on Hypertension; Council on Lifelong Congenital Heart Disease and Heart Health in the Young; Council on Lifestyle and Cardiometabolic Health; Council on Peripheral Vascular Disease; and Stroke Council. Cardiovascular disease risk factors in women: the impact of race and ethnicity: a scientific statement from the American Heart Association. *Circulation*. 2023;147:1471–1487. doi: 10.1161/CIR.0000000000001139
- Rodriguez CJ, Allison M, Daviglius ML, Isasi CR, Keller C, Leira EC, Palaniappan L, Piña IL, Ramirez SM, Rodriguez B, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention. Status of cardiovascular disease and stroke in Hispanics/Latinos in the United States: a science advisory from the American Heart Association. *Circulation*. 2014;130:593–625. doi: 10.1161/CIR.0000000000000701
- Carnethon MR, Fu J, Howard G, Albert MA, Anderson CAM, Bertoni AG, Mujahid MS, Palaniappan L, Taylor HA, Willis M, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention; Council on Cardiovascular Disease in the Young; Council on Cardiovascular and Stroke Nursing; Council on Clinical Cardiology; Council on Functional Genomics and Translational Biology; and Stroke Council. Cardiovascular health in African Americans: a scientific statement from the American Heart Association. *Circulation*. 2017;136:e393–e423. doi: 10.1161/CIR.0000000000000534
- Breathett K, Sims M, Gross M, Jackson EA, Jones EJ, Navas-Acien A, Taylor H, Thomas KL, Howard BV; on behalf of the American Heart Association Council on Epidemiology and Prevention; Council on Quality of Care and Outcomes Research; Council on Cardiovascular and Stroke Nursing; Council on Clinical Cardiology; and Council on Lifestyle and Cardiometabolic Health. Cardiovascular health in American Indians and Alaska Natives: a scientific statement from the American Heart Association. *Circulation*. 2020;141:e948–e959. doi: 10.1161/CIR.0000000000000773
- Kwan TW, Wong SS, Hong Y, Kanaya AM, Khan SS, Hayman LL, Shah SH, Welty FK, Deedwania PC, Khaliq A, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention; Council on Lifestyle and Cardiometabolic Health; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Clinical Cardiology; Council on Cardiovascular and Stroke Nursing; and Council on Genomic and Precision Medicine. Epidemiology of diabetes and atherosclerotic cardiovascular disease among Asian American adults: implications, management, and future directions: a scientific statement from the American Heart Association. *Circulation*. 2023;148:74–94. doi: 10.1161/CIR.0000000000001145
- Havranek EP, Mujahid MS, Barr DA, Blair IV, Cohen MS, Cruz-Flores S, Davey-Smith G, Dennison-Himmelfarb CR, Lauer MS, Lockwood DW, et al. Social determinants of risk and outcomes for cardiovascular disease. *Circulation*. 2015;132:873–898. doi: 10.1161/CIR.0000000000000228
- Kyalwazi AN, Loccoh EC, Brewer LC, Ofili EO, Xu J, Song Y, Joynt Maddox KE, Yeh RW, Wadhwa RK. Disparities in cardiovascular mortality between Black and White adults in the United States, 1999 to 2019. *Circulation*. 2022;146:211–228. doi: 10.1161/CIRCULATIONAHA.122.060199
- Powell-Wiley TM, Baumer Y, Baah FO, Baez AS, Farmer N, Mahlobo CT, Pita MA, Potharaju KA, Tamura K, Wallen GR. Social determinants of cardiovascular disease. *Circ Res*. 2022;130:782–799. doi: 10.1161/CIRCRESAHA.121.319811
- Allana S, Ski CF, Thompson DR, Clark AM. Bringing intersectionality to cardiovascular health research in Canada. *CJC Open*. 2021;3:S4–S8. doi: 10.1016/j.cjco.2021.08.016
- Baiden D, Parry M, Nerenberg K, Hillan EM, Dogba MJ. Connecting the dots: structural racism, intersectionality, and cardiovascular health outcomes for African, Caribbean, and Black mothers. *Health Equity*. 2022;6:402–405. doi: 10.1089/heq.2021.0077
- Ogunniyi MO, Mahmoud Z, Commodore-Mensah Y, Fleg JL, Fatade YA, Quesada O, Aggarwal NR, Mattina DJ, Oliveira GMM, Lindley KJ, et al. Eliminating disparities in cardiovascular disease for Black women. *J Am Coll Cardiol*. 2022;80:1762–1771. doi: 10.1016/j.jacc.2022.08.769
- Vespa J, Medina L, Armstrong DM. Demographic turning points for the United States: population projections for 2020 to 2060. US Census Bureau. 2020. Accessed April 12, 2025. <https://census.gov/content/dam/Census/library/publications/2020/demo/p25-1144.pdf>
- Ovbiagele B, Goldstein LB, Higashida RT, Haddock VJ, Johnston SC, Khavjou OA, Lackland DT, Lichtman JH, Mohl S, Sacco RL, et al. Forecasting the future of stroke in the United States. *Stroke*. 2013;44:2361–2375. doi: 10.1161/str.0b013e31829734f2
- Dunbar SB, Khavjou OA, Bakas T, Hunt G, Kirch RA, Leib AR, Morrison RS, Poehler DC, Roger VL, Whitsett LP; on behalf of the American Heart Association. Projected costs of informal caregiving for cardiovascular

- disease: 2015 to 2035: a policy statement from the American Heart Association. *Circulation*. 2018;137:e558–e577. doi: 10.1161/cir.0000000000000570
28. Heidenreich PA, Albert NM, Allen LA, Bluemke DA, Butler J, Fonarow GC, Ikonomicis JS, Khavjou O, Konstam MA, Maddox TM, et al; on behalf of the American Heart Association Advocacy Coordinating Committee; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Cardiovascular Radiology and Intervention; Council on Clinical Cardiology; Council on Epidemiology and Prevention; Stroke Council. Forecasting the impact of heart failure in the United States: a policy statement from the American Heart Association. *Circ Heart Fail*. 2013;6:606–619. doi: 10.1161/HHF.0b013e318291329a
 29. Heidenreich PA, Trogdon JG, Khavjou OA, Butler J, Dracup K, Ezekowitz MD, Finkelstein EA, Hong Y, Johnston SC, Khera A, et al; on behalf of the American Heart Association Advocacy Coordinating Committee; Stroke Council; Council on Cardiovascular Radiology and Intervention; Council on Clinical Cardiology; Council on Epidemiology and Prevention; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Cardiology; Council on Cardiovascular Surgery and Anesthesia; and Interdisciplinary Council on Quality of Care and Outcomes Research. Forecasting the future of cardiovascular disease in the United States: a policy statement from the American Heart Association. *Circulation*. 2011;123:933–944. doi: 10.1161/CIR.0b013e31820a55f5
 30. Chen T-C, Clark J, Riddles MK, Mohadjer LK, Fakhouri THI. National Health and Nutrition Examination Survey, 2015–2018: sample design and estimation procedures. 2020. Accessed April 12, 2025. https://cdc.gov/nchs/data/series/sr_02/sr02-184-508.pdf
 31. Agency for Healthcare Research and Quality. Medical Expenditure Panel Survey (MEPS). Accessed October 8, 2025. <https://meps.ahrq.gov/mepsweb/>
 32. US Department of Health and Human Services. *45 CFR Part 46: Protection of Human Subjects*. Federal Register; 2023.
 33. Liu E, Bigeh A, Ledingham L, Mehta L. Prevention of coronary artery disease in women. *Curr Cardiol Rep*. 2022;24:1041–1048. doi: 10.1007/s11886-022-01721-5
 34. Arnett DK, Blumenthal RS, Albert MA, Buroker AB, Goldberger ZD, Hahn EJ, Himmelfarb CD, Khera A, Lloyd-Jones D, McEvoy JW, et al. 2019 ACC/AHA guideline on the primary prevention of cardiovascular disease: a report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation*. 2019;140:e596–e646. doi: 10.1161/CIR.0000000000000678
 35. Cho L, Davis M, Elgendy I, Epps K, Lindley KJ, Mehta PK, Michos ED, Minissian M, Pepine C, Vaccarino V, et al; ACC CVD Women's Committee Members. Summary of updated recommendations for primary prevention of cardiovascular disease in women: JACC state-of-the-art review. *J Am Coll Cardiol*. 2020;75:2602–2618. doi: 10.1016/j.jacc.2020.03.060
 36. Weintraub WS, Daniels SR, Burke LE, Franklin BA, Goff DC Jr, Hayman LL, Lloyd-Jones D, Pandey DK, Sanchez EJ, Schram AP, et al; on behalf of the American Heart Association Advocacy Coordinating Committee; Council on Cardiovascular Disease in the Young; Council on the Kidney in Cardiovascular Disease; Council on Epidemiology and Prevention; Council on Cardiovascular Nursing; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Clinical Cardiology, and Stroke Council. Value of primordial and primary prevention for cardiovascular disease: a policy statement from the American Heart Association. *Circulation*. 2011;124:967–990. doi: 10.1161/CIR.0b013e3182285a81
 37. Chomistek AK, Chiuvè SE, Eliassen AH, Mukamal KJ, Willett WC, Rimm EB. Healthy lifestyle in the primordial prevention of cardiovascular disease among young women. *J Am Coll Cardiol*. 2015;65:43–51. doi: 10.1016/j.jacc.2014.10.024
 38. Sotos-Prieto M, Mattei J, Hu FB, Chomistek AK, Rimm EB, Willett WC, Eliassen AH, Chiuvè SE. Association between a healthy heart score and the development of clinical cardiovascular risk factors among women: potential role for primordial prevention. *Circ Cardiovasc Qual Outcomes*. 2016;9:S77–S85. doi: 10.1161/circoutcomes.115.002372
 39. Colpani V, Baena CP, Jaspers L, van Dijk GM, Farajzadegan Z, Dhana K, Tielemans MJ, Voortman T, Freak-Poli R, Veloso GGV, et al. Lifestyle factors, cardiovascular disease and all-cause mortality in middle-aged and elderly women: a systematic review and meta-analysis. *Eur J Epidemiol*. 2018;33:831–845. doi: 10.1007/s10654-018-0374-z
 40. Seguin-Fowler RA, Eldridge GD, Rethorst CD, Graham ML, Demment M, Strogatz D, Folta SC, Maddock JE, Nelson ME, Ha S. Improvements and maintenance of clinical and functional measures among rural women: strong hearts, Healthy Communities-2. 0 cluster randomized trial. *Circ Cardiovasc Qual Outcomes*. 2022;15:e009333. doi: 10.1161/CIRCOUTCOMES.122.009333
 41. Pengo MF, Won CH, Bourjeily G. Sleep in women across the life span. *Chest*. 2018;154:196–206. doi: 10.1016/j.chest.2018.04.005
 42. Gooding HC, Aguayo L. Hard lessons from a school-based intervention to improve cardiovascular health in adolescence. *JAMA Cardiol*. 2023;8:825–826. doi: 10.1001/jamacardio.2023.2244
 43. Muth ND, Bolling C, Hannon T, Sharifi M; Section on Obesity; Committee on Nutrition. The role of the pediatrician in the promotion of healthy, active living. *Pediatrics*. 2024;153:e2023065480. doi: 10.1542/peds.2023-065480
 44. Brown HL, Warner JJ, Gianos E, Gulati M, Hill AJ, Hollier LM, Rosen SE, Rosser ML, Wenger NK; on behalf of the American Heart Association and the American College of Obstetricians and Gynecologists. Promoting risk identification and reduction of cardiovascular disease in women through collaboration with obstetricians and gynecologists: a presidential advisory from the American Heart Association and the American College of Obstetricians and Gynecologists. *Circulation*. 2018;137:e843–e852. doi: 10.1161/CIR.0000000000000582
 45. Petkovic J, Duench S, Travin J, Dewidar O, Pardo Pardo J, Simeon R, DesMeules M, Gagnon D, Hatcher Roberts J, Hossain A, et al. Behavioural interventions delivered through interactive social media for health behaviour change, health outcomes, and health equity in the adult population. *Cochrane Database Syst Rev*. 2021;5:CD012932. doi: 10.1002/14651858.CD012932.pub2
 46. Kazi DS. State of the heart (and brain) in 2025. *Circulation*. 2025;151:515–517. doi: 10.1161/CIRCULATIONAHA.125.073826
 47. Meng Y, Sharman JE, Iiskala F, Wu F, Juonala M, Pahkala K, Rovio SP, Fraser BJ, Kelly RK, Hutri N, et al. Tracking and transition probability of blood pressure from childhood to midadulthood. *JAMA Pediatr*. 2025;179:34–45. doi: 10.1001/jamapediatrics.2024.4368
 48. Aris IM, Rifas-Shiman SL, Peng W, Yi L, de Ferranti SD, Hivert MF, Oken E. Trajectory of cardiovascular health across childhood and adolescence. *JAMA Cardiol*. 2024;10:417. doi: 10.1001/jamacardio.2024.4022
 49. Flynn JT, Urbina EM, Brady TM, Baker-Smith C, Daniels SR, Hayman LL, Mitsnefes M, Tran A, Zachariah JP; on behalf of the Atherosclerosis, Hypertension, and Obesity in the Young Committee of the American Heart Association Council on Lifelong Congenital Heart Disease and Heart Health in the Young; Council on Cardiovascular Radiology and Intervention; Council on Epidemiology and Prevention; Council on Hypertension; and Council on Lifestyle and Cardiometabolic Health. Ambulatory blood pressure monitoring in children and adolescents: 2022 update: a scientific statement from the American Heart Association. *Hypertension*. 2022;79:e114–e124. doi: 10.1161/HYP.0000000000000215
 50. Jones DW, Whelton PK, Allen N, Clark D 3rd, Gidding SS, Muntner P, Nesbitt S, Mitchell NS, Townsend R, Falkner B; on behalf of the American Heart Association Council on Hypertension; Council on the Kidney in Cardiovascular Disease; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Cardiovascular Radiology and Intervention; Council on Lifelong Congenital Heart Disease and Heart Health in the Young; and Stroke Council. Management of stage 1 hypertension in adults with a low 10-year risk for cardiovascular disease: filling a guidance gap: a scientific statement from the American Heart Association. *Hypertension*. 2021;77:e58–e67. doi: 10.1161/HYP.0000000000000195
 51. Flynn JT, Kaelber DC, Baker-Smith CM, Blowey D, Carroll AE, Daniels SR, de Ferranti SD, Dionne JM, Falkner B, Flinn SK, et al; Subcommittee on Screening and Management of High Blood Pressure in Children. Clinical practice guideline for screening and management of high blood pressure in children and adolescents. *Pediatrics*. 2017;140:e20171904. doi: 10.1542/peds.2017-1904
 52. Lam CSP, Arnott C, Beale AL, Chandramouli C, Hilfiker-Kleiner D, Kaye DM, Ky B, Santema BT, Sliwa K, Voors AA. Sex differences in heart failure. *Eur Heart J*. 2019;40:3859–3868. doi: 10.1093/eurheartj/ehz835
 53. Wang X, Carcel C, Woodward M, Schutte AE. Blood pressure and stroke: a review of sex- and ethnic/racial-specific attributes to the epidemiology, pathophysiology, and management of raised blood pressure. *Stroke*. 2022;53:1114–1133. doi: 10.1161/STROKEAHA.121.035852
 54. Rexrode KM, Madsen TE, Yu AYY, Carcel C, Lichtman JH, Miller EC. The impact of sex and gender on stroke. *Circ Res*. 2022;130:512–528. doi: 10.1161/CIRCRESAHA.121.319915
 55. Wright JT Jr, Williamson JD, Whelton PK, Snyder JK, Sink KM, Rocco MV, Reboussin DM, Rahman M, Oparil S, Lewis CE, et al; SPRINT Research Group. A randomized trial of intensive versus standard blood-pressure control. *N Engl J Med*. 2015;373:2103–2116. doi: 10.1056/NEJMoa1511939
 56. Cushman WC, Evans GW, Byington RP, Goff DC Jr, Grimm RH Jr, Cutler JA, Simons-Morton DG, Basile JN, Corson MA, Probstfield JL, et al; ACCORD Study Group. Effects of intensive blood-pressure control in

- type 2 diabetes mellitus. *N Engl J Med*. 2010;362:1575–1585. doi: 10.1056/NEJMoa1001286
57. Yaow CYL, Chong B, Chin YH, Kueh MTW, Ng CH, Chan KE, Tang ASP, Chung C, Goh R, Kong G, et al. Higher risk of adverse cardiovascular outcomes in females with type 2 diabetes mellitus: an umbrella review of systematic reviews. *Eur J Prev Cardiol*. 2023;30:1227–1235. doi: 10.1093/eurjpc/zwad133
 58. Peters SA, Huxley RR, Woodward M. Diabetes as a risk factor for stroke in women compared with men: a systematic review and meta-analysis of 64 cohorts, including 775,385 individuals and 12,539 strokes. *Lancet*. 2014;383:1973–1980. doi: 10.1016/S0140-6736(14)60040-4
 59. Sillars A, Ho FK, Pell GP, Gill JMR, Sattar N, Gray S, Celis-Morales C. Sex differences in the association of risk factors for heart failure incidence and mortality. *Heart*. 2020;106:203–212. doi: 10.1136/heartjnl-2019-314878
 60. Yoshida Y, Chen Z, Fonseca VA, Mauvais-Jarvis F. Sex differences in cardiovascular risk associated with prediabetes and undiagnosed diabetes. *Am J Prev Med*. 2023;65:854–862. doi: 10.1016/j.amepre.2023.05.011
 61. Gerstein HC, Miller ME, Byington RP, Goff DC Jr, Bigger JT, Buse JB, Cushman WC, Genuth S, Ismail-Beigi F, Grimm RH Jr, et al; Action to Control Cardiovascular Risk in Diabetes Study Group. Effects of intensive glucose lowering in type 2 diabetes. *N Engl J Med*. 2008;358:2545–2559. doi: 10.1056/NEJMoa0802743
 62. Khan SS, Matsushita K, Sang Y, Ballew SH, Grams ME, Surapaneni A, Blaha MJ, Carson AP, Chang AR, Ciemiens E, et al. Development and validation of the American Heart Association Predicting Risk of Cardiovascular Disease EVENTS (PREVENT) equations. *Circulation*. 2024;149:430–449. doi: 10.1161/circulationaha.123.067626
 63. Ndumele CE, Rangaswami J, Chow SL, Neeland IJ, Tuttle KR, Khan SS, Coresh J, Mathew RO, Baker-Smith CM, Carnethon MR, et al; on behalf of the American Heart Association. Cardiovascular-kidney-metabolic health: a presidential advisory from the American Heart Association [published correction appears in *Circulation*. 2024;149:e1023]. *Circulation*. 2023;148:1606–1635. doi: 10.1161/cir.0000000000001184
 64. Emmerich SD, Fryar CD, Stierman B, Ogden CL. Obesity and severe obesity prevalence in adults: United States, August 2021–August 2023. *NCHS Data Brief*. 2024;10.15620/cdc/159281. doi: 10.15620/cdc/159281
 65. Börchers S, Skibicka KP. GLP-1 and its analogs: does sex matter? *Endocrinology*. 2025;166:bqae165. doi: 10.1210/endo/bqae165
 66. Perak AM, Ning H, Kit BK, de Ferranti SD, Van Horn LV, Wilkins JT, Lloyd-Jones DM. Trends in levels of lipids and apolipoprotein B in US youths aged 6 to 19 years, 1999–2016. *JAMA*. 2019;321:1895–1905. doi: 10.1001/jama.2019.4984
 67. Hayes DK, Robbins CL, Ko JY. Trends in selected chronic conditions and related risk factors among women of reproductive age: Behavioral Risk Factor Surveillance System, 2011–2017. *J Womens Health (Larchmt)*. 2020;29:1576–1585. doi: 10.1089/jwh.2019.8275
 68. Casagrande SS, Menke A, Cowie CC. Cardiovascular risk factors of adults age 20–49 years in the United States, 1971–2012: a series of cross-sectional studies. *PLoS One*. 2016;11:e0161770. doi: 10.1371/journal.pone.0161770
 69. Patel N, Bhargava A, Kalra R, Parcha V, Arora G, Muntner P, Arora P. Trends in Lipid, Lipoproteins, and statin use among U.S. adults: impact of 2013 cholesterol guidelines. *J Am Coll Cardiol*. 2019;74:2525–2528. doi: 10.1016/j.jacc.2019.09.026
 70. Wu B, Fan B, Qu Y, Li C, Chen J, Liu Y, Wang J, Zhang T, Chen Y. Trajectories of Blood lipids profile in midlife women: does menopause matter? *J Am Heart Assoc*. 2023;12:e030388. doi: 10.1161/jaha.123.030388
 71. Mora S, Glynn RJ, Hsia J, MacFadyen JG, Genest J, Ridker PM. Statins for the primary prevention of cardiovascular events in women with elevated high-sensitivity C-reactive protein or dyslipidemia: results from the Justification for the Use of Statins in Prevention: An Intervention Trial Evaluating Rosuvastatin (JUPITER) and Meta-Analysis of Women From Primary Prevention trials. *Circulation*. 2010;121:1069–1077. doi: 10.1161/CIRCULATIONAHA.109.906479
 72. Lee MT, Mahitta D, Ramsey DJ, Liu J, Misra A, Nasir K, Samad Z, Itchhaporia D, Khan SU, Schofield RS, et al. Sex-related disparities in cardiovascular health care among patients with premature atherosclerotic cardiovascular disease. *JAMA Cardiol*. 2021;6:782–790. doi: 10.1001/jamacardio.2021.0683
 73. Udell JA, Fonarow GC, Maddox TM, Cannon CP, Frank Peacock W, Laskey WK, Grau-Sepulveda MV, Smith EE, Hernandez AF, Peterson ED, et al. Sustained sex-based treatment differences in acute coronary syndrome care: insights from the American Heart Association Get With The Guidelines Coronary Artery Disease Registry. *Clin Cardiol*. 2018;41:758–768. doi: 10.1002/clc.22938
 74. American College of Cardiology. Focus on heart failure: heart failure in women: understanding the differences to change the paradigm. 2024. Accessed May 22, 2025. <https://acc.org/Latest-in-Cardiology/Articles/2024/02/01/01/42/focus-on-heart-failure-heart-failure-in-women-understanding-the-differences-to-change-the-paradigm>
 75. Walli-Attai M, Joseph P, Johansson I, Siwa K, Lonn E, Maggioni AP, Mielniczuk L, Ross H, Karaye K, Hage C, et al; G-CHF Investigators. Characteristics, management, and outcomes in women and men with congestive heart failure in 40 countries at different economic levels: an analysis from the Global Congestive Heart Failure (G-CHF) registry. *Lancet Glob Health*. 2024;12:e396–e405. doi: 10.1016/S2214-109X(23)00557-0
 76. Strong B, Lisabeth LD, Reeves M. Sex differences in IV thrombolysis treatment for acute ischemic stroke: a systematic review and meta-analysis. *Neurology*. 2020;95:e11–e22. doi: 10.1212/WNL.0000000000009733
 77. Otite FO, Morris N. Race, ethnicity, and gender disparities in the management and outcomes of critically ill adults with acute stroke. *Crit Care Clin*. 2024;40:709–740. doi: 10.1016/j.ccc.2024.05.006
 78. Khan SU, Khan MZ, Khan MU, Khan MS, Mamas MA, Rashid M, Blankstein R, Virani SS, Johansen MC, Shapiro MD, et al. Clinical and economic burden of stroke among young, midlife, and older adults in the United States, 2002–2017. *Mayo Clin Proc Innov Qual Outcomes*. 2021;5:431–441. doi: 10.1016/j.mayocpiqo.2021.01.015
 79. Ford ND, Cox S, Ko JY, Ouyang L, Romero L, Colarusso T, Ferre CD, Kroelinger CD, Hayes DK, Barfield WD. Hypertensive disorders in pregnancy and mortality at delivery hospitalization—United States, 2017–2019. *MMWR Morb Mortal Wkly Rep*. 2022;71:585–591. doi: 10.15585/mmwr.mm7117a1
 80. Bushnell K, Kernan WN, Sharrief AZ, Chaturvedi S, Cole JW, Cornwall WK 3rd, Cosby-Gaither C, Doyle S, Goldstein LB, Lennon O, et al. 2024 Guideline for the primary prevention of stroke: a guideline from the American Heart Association/American Stroke Association [published corrections appear in *Stroke*. 2024;55:e439 and *Stroke*. 2025;56:e98]. *Stroke*. 2024;55:e344–e424. doi: 10.1161/STR.00000000000000475
 81. Ko D, Rahman F, Schnabel RB, Yin X, Benjamin EJ, Christophersen IE. Atrial fibrillation in women: epidemiology, pathophysiology, presentation, and prognosis. *Nat Rev Cardiol*. 2016;13:321–332. doi: 10.1038/nrcardio.2016.45
 82. Yong CM, Tremmel JA, Lansberg MG, Fan J, Askari M, Turakhia MP. Sex differences in oral anticoagulation and outcomes of stroke and intracranial bleeding in newly diagnosed atrial fibrillation. *J Am Heart Assoc*. 2020;9:e015689. doi: 10.1161/JAHA.120.015689
 83. Subramanya V, Claxton JS, Lutsey PL, MacLehose RF, Chen LY, Chamberlain AM, Norby FL, Alonso A. Sex differences in treatment strategy and adverse outcomes among patients 75 and older with atrial fibrillation in the MarketScan database. *BMC Cardiovasc Disord*. 2021;21:598. doi: 10.1186/s12872-021-02419-2
 84. Iadecola C, Duering M, Hachinski V, Joutel A, Pendlebury ST, Schneider JA, Dichgans M. Vascular cognitive impairment and dementia: JACC Scientific Expert Panel. *J Am Coll Cardiol*. 2019;73:3326–3344. doi: 10.1016/j.jacc.2019.04.034
 85. SPRINT MIND Investigators for the SPRINT Research Group; Williamson JD, Pajewski NM, Auchus AP, Bryan RN, Chelune G, Cheung AK, Cleveland ML, Coker LH, Crowe MG, Cushman WC, et al; Effect of intensive vs standard blood pressure control on probable dementia: a randomized clinical trial. *JAMA*. 2019;321:553–561. doi: 10.1001/jama.2018.21442
 86. Lazar RM, Howard VJ, Kernan WN, Aparicio HJ, Levine DA, Viera AJ, Jordan LC, Nyenhuis DL, Possin KL, Sorond FA, et al; on behalf of the American Heart Association Stroke Council. A primary care agenda for brain health: a scientific statement from the American Heart Association. *Stroke*. 2021;52:e295–e308. doi: 10.1161/STR.0000000000000367
 87. Hachinski V, Einhaupl K, Ganten D, Alladi S, Brayne C, Stephan BCM, Sweeney MD, Zlokovic B, Itturria-Medina Y, Iadecola C, et al. Preventing dementia by preventing stroke: the Berlin Manifesto. *Alzheimers Dement*. 2019;15:961–984. doi: 10.1016/j.jalz.2019.06.001
 88. Morris HR, Spillantini MG, Sue CM, Williams-Gray CH. The pathogenesis of Parkinson's disease. *Lancet*. 2024;403:293–304. doi: 10.1016/S0140-6736(23)01478-2
 89. Kolahchi Z, Henkel N, Eladawi MA, Villarreal EC, Kandimalla P, Lundh A, McCullumsmith RE, Cuevas E. Sex and gender differences in Alzheimer's disease: genetic, hormonal, and inflammation impacts. *Int J Mol Sci*. 2024;25:8485. doi: 10.3390/ijms25158485
 90. Behboudi-Gandevan S, Moe CF, Skjesol I, Arntzen EC, Bidhendi-Yarandi R. The J shaped association of age at menarche and cardiovascular events: systematic review and meta-analysis. *Sci Rep*. 2024;14:2695. doi: 10.1038/s41598-024-53011-5

91. Lakshman R, Forouhi NG, Sharp SJ, Luben R, Bingham SA, Khaw KT, Wareham NJ, Ong KK. Early age at menarche associated with cardiovascular disease and mortality. *J Clin Endocrinol Metab*. 2009;94:4953–4960. doi: 10.1210/jc.2009-1789
92. Dreyfus J, Jacobs DR Jr, Mueller N, Schreiner PJ, Moran A, Carnethon MR, Demerath EW. Age at menarche and cardiometabolic risk in adulthood: the Coronary Artery Risk Development in Young Adults study. *J Pediatr*. 2015;167:344–352.e1. doi: 10.1016/j.jpeds.2015.04.032
93. Martinez GM. Trends and patterns in menarche in the United States: 1995 through 2013–2017. 2020. Accessed May 7, 2025. <https://stacks.cdc.gov/view/cdc/93643>
94. Wang YX, Stuart JJ, Rich-Edwards JW, Missmer SA, Rexrode KM, Farland LV, Mukamal KJ, Nelson SM, Solomon CG, Fraser A, et al. Menstrual cycle regularity and length across the reproductive lifespan and risk of cardiovascular disease. *JAMA Netw Open*. 2022;5:e2238513. doi: 10.1001/jamanetworkopen.2022.38513
95. Hing E, Albert M. State variation in preventive care visits, by patient characteristics, 2012. Accessed May 23, 2025. 2012. [https://cdc.gov/nchs/products/databriefs/db234.htm#:~:text=The%20percentage%20of%20preventive%20care%20visits%20to,those%20for%20women%20\(21.8%%20and%2044.1%%2C%20respectively\)](https://cdc.gov/nchs/products/databriefs/db234.htm#:~:text=The%20percentage%20of%20preventive%20care%20visits%20to,those%20for%20women%20(21.8%%20and%2044.1%%2C%20respectively))
96. Sharma G, Gaffey AE, Hameed A, Kasparian NA, Mauricio R, Marsh EB, Beck D, Skowronski J, Wolfe D, Levine GN; on behalf of the American Heart Association Women's Health Science Committee of the Council on Clinical Cardiology and the Stroke Council; Council on Cardiovascular and Stroke Nursing; Council on Cardiovascular Radiology and Intervention; Council on Cardiovascular Surgery and Anesthesia; Council on Lifelong Congenital Heart Disease and Heart Health in the Young; Council on Lifestyle and Cardiometabolic Health; and Council on Quality of Care and Outcomes Research. Optimizing psychological health across the perinatal period: an update on maternal cardiovascular health: a scientific statement from the American Heart Association [published correction appears in *J Am Heart Assoc*. 2025;14:e10898]. *J Am Heart Assoc*. 2025;14:e041369. doi: 10.1161/JAHA.125.041369
97. Lewey J, Beckie TM, Brown HL, Brown SD, Garovic VD, Khan SS, Miller EC, Sharma G, Mehta LS; on behalf of the American Heart Association Cardiovascular Disease and Stroke in Women and Underrepresented Populations Committee of the Council on Clinical Cardiology; Council on Cardiopulmonary, Critical Care, Perioperative and Resuscitation; and Council on Cardiovascular and Stroke Nursing. Opportunities in the postpartum period to reduce cardiovascular disease risk after adverse pregnancy outcomes: a scientific statement from the American Heart Association. *Circulation*. 2024;149:e330–e346. doi: 10.1161/CIR.0000000000001212
98. Leifheit-Limson EC, D'Onofrio G, Daneshvar M, Geda M, Bueno H, Spertus JA, Krumholz HM, Lichtman JH. Sex differences in cardiac risk factors, perceived risk, and health care provider discussion of risk and risk modification among young patients with acute myocardial infarction: the VIRGO study. *J Am Coll Cardiol*. 2015;66:1949–1957. doi: 10.1016/j.jacc.2015.08.859
99. D'Onofrio G, Safdar B, Lichtman JH, Strait KM, Dreyer RP, Geda M, Spertus JA, Krumholz HM. Sex differences in reperfusion in young patients with ST-segment-elevation myocardial infarction: results from the VIRGO study. *Circulation*. 2015;131:1324–1332. doi: 10.1161/CIRCULATIONAHA.114.012293
100. Sheidu MO, Agarwala A, Lakshmanan S, Honigberg MC, Spitz JA, Sharma G. Management of pregnancy-related disorders to prevent future risk of coronary artery disease. *Heart*. 2024;111:83–92. doi: 10.1136/heartjnl-2022-321606
101. Parikh NI, Gonzalez JM, Anderson CAM, Judd SE, Rexrode KM, Hlatky MA, Gunderson EP, Stuart JJ, Vaidya D; on behalf of the American Heart Association Council on Epidemiology and Prevention; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Cardiovascular and Stroke Nursing; and the Stroke Council. Adverse pregnancy outcomes and cardiovascular disease risk: unique opportunities for cardiovascular disease prevention in women: a scientific statement from the American Heart Association. *Circulation*. 2021;143:e902–e916. doi: 10.1161/CIR.0000000000000961
102. Khan SS, Brewer LC, Canobbio MM, Cipolla MJ, Grobman WA, Lewey J, Michos ED, Miller EC, Perak AM, Wei GS, et al; on behalf of the American Heart Association Council on Epidemiology and Prevention; Council on Clinical Cardiology; Council on Cardiovascular and Stroke Nursing; Council on Arteriosclerosis, Thrombosis and Vascular Biology; Council on Hypertension; Council on Lifestyle and Cardiometabolic Health; Council on Peripheral Vascular Disease; and Stroke Council. Optimizing prepregnancy cardiovascular health to improve outcomes in pregnant and postpartum individuals and offspring: a scientific statement from the American Heart Association. *Circulation*. 2023;147:e76–e91. doi: 10.1161/CIR.0000000000001124
103. Gaillard R, Jadot V. Maternal cardiovascular disorders before and during pregnancy and offspring cardiovascular risk across the life course. *Nat Rev Cardiol*. 2023;20:617–630. doi: 10.1038/s41569-023-00869-z
104. Palinski W. Effect of maternal cardiovascular conditions and risk factors on offspring cardiovascular disease. *Circulation*. 2014;129:2066–2077. doi: 10.1161/CIRCULATIONAHA.113.001805
105. Russell MW. JAHA spotlight on pregnancy and its impact on maternal and offspring cardiovascular health. *J Am Heart Assoc*. 2022;11:e025167. doi: 10.1161/JAHA.121.025167
106. Benschop L, Brouwers L, Zoet GA, Meun C, Boersma E, Budde RPJ, Fauser B, de Groot CMJ, van der Schouw YT, Maas A, et al. Early onset of coronary artery calcification in women with previous preeclampsia. *Circ Cardiovasc Imaging*. 2020;13:e010340. doi: 10.1161/circimaging.119.010340
107. Williams D, Stout MJ, Rosenbloom JI, Olsen MA, Joynt Maddox KE, Deych E, Davila-Roman VG, Lindley KJ. Preeclampsia predicts risk of hospitalization for heart failure with preserved ejection fraction. *J Am Coll Cardiol*. 2021;78:2281–2290. doi: 10.1016/j.jacc.2021.09.1360
108. Mehta A, Spitz J, Sharma S, Bonomo J, Brewer LC, Mehta LS, Sharma G. Addressing social determinants of health in maternal cardiovascular health. *Can J Cardiol*. 2024;40:1031–1042. doi: 10.1016/j.cjca.2024.02.010
109. El Khoudary SR, Aggarwal B, Beckie TM, Hodis HN, Johnson AE, Langer RD, Limacher MC, Manson JE, Stefanick ML, Allison MA; on behalf of the American Heart Association Prevention Science Committee of the Council on Epidemiology and Prevention; and Council on Cardiovascular and Stroke Nursing. Menopause transition and cardiovascular disease risk: implications for timing of early prevention: a scientific statement from the American Heart Association. *Circulation*. 2020;142:e506–e532. doi: 10.1161/CIR.0000000000000912
110. Kravitz HM, Janssen I, Bromberger JT, Matthews KA, Hall MH, Ruppert K, Joffe H. Sleep trajectories before and after the final menstrual period in the Study of Women's Health Across the Nation (SWAN). *Curr Sleep Med Rep*. 2017;3:235–250. doi: 10.1007/s40675-017-0084-1
111. Gold EB, Crawford SL, Avis NE, Crandall CJ, Matthews KA, Waetjen LE, Lee JS, Thurston R, Vuga M, Harlow SD. Factors related to age at natural menopause: longitudinal analyses from SWAN. *Am J Epidemiol*. 2013;178:70–83. doi: 10.1093/aje/kws421
112. Dorjgochoo T, Kallianpur A, Gao YT, Cai H, Yang G, Li H, Zheng W, Shu XO. Dietary and lifestyle predictors of age at natural menopause and reproductive span in the Shanghai Women's Health Study. *Menopause*. 2008;15:924–933. doi: 10.1097/gme.0b013e3181786ad6
113. Kok HS, van Asselt KM, van der Schouw YT, van der Tweel I, Peeters PH, Wilson PW, Pearson PL, Grobbee DE. Heart disease risk determines menopausal age rather than the reverse. *J Am Coll Cardiol*. 2006;47:1976–1983. doi: 10.1016/j.jacc.2005.12.066
114. Nagata C, Takatsuka N, Kawakami N, Shimizu H. Association of diet with the onset of menopause in Japanese women. *Am J Epidemiol*. 2000;152:863–867. doi: 10.1093/aje/152.9.863
115. Parente RC, Faerstein E, Celeste RK, Werneck GL. The relationship between smoking and age at the menopause: a systematic review. *Maturitas*. 2008;61:287–298. doi: 10.1016/j.maturitas.2008.09.021
116. Matthews KA, Crawford SL, Chae CU, Everson-Rose SA, Sowers MF, Sternfeld B, Sutton-Tyrrell K. Are changes in cardiovascular disease risk factors in midlife women due to chronological aging or to the menopausal transition? *J Am Coll Cardiol*. 2009;54:2366–2373. doi: 10.1016/j.jacc.2009.10.009
117. Thurston RC, Karvonen-Gutierrez CA, Derby CA, El Khoudary SR, Kravitz HM, Manson JE. Menopause versus chronologic aging: their roles in women's health. *Menopause*. 2018;25:849–854. doi: 10.1097/GME.0000000000001143
118. Rosenson RS, Brewer HB Jr, Chapman MJ, Fazio S, Hussain MM, Kontush A, Krauss RM, Otvos JD, Remaley AT, Schaefer EJ. HDL measures, particle heterogeneity, proposed nomenclature, and relation to atherosclerotic cardiovascular events. *Clin Chem*. 2011;57:392–410. doi: 10.1373/clinchem.2010.155333
119. Lejskova M, Alusik S, Valenta Z, Adamkova S, Pitha J. Natural postmenopause is associated with an increase in combined cardiovascular risk factors. *Physiol Res*. 2012;61:587–596. doi: 10.33549/physiolres.932313
120. Tepper PG, Randolph JF Jr, McConnell DS, Crawford SL, El Khoudary SR, Joffe H, Gold EB, Zheng H, Bromberger JT, Sutton-Tyrrell K. Trajectory clustering of estradiol and follicle-stimulating hormone during the

- menopausal transition among women in the Study of Women's Health across the Nation (SWAN). *J Clin Endocrinol Metab.* 2012;97:2872–2880. doi: 10.1210/jc.2012-1422
121. El Khoudary SR, Greendale G, Crawford SL, Avis NE, Brooks MM, Thurston RC, Karvonen-Gutierrez C, Waetjen LE, Matthews K. The menopause transition and women's health at midlife: a progress report from the Study of Women's Health Across the Nation (SWAN). *Menopause.* 2019;26:1213–1227. doi: 10.1097/GME.0000000000001424
 122. Matthews KA, Everson-Rose SA, Kravitz HM, Lee L, Janssen I, Sutton-Tyrrell K. Do reports of sleep disturbance relate to coronary and aortic calcification in healthy middle-aged women? Study of Women's Health Across the Nation. *Sleep Med.* 2013;14:282–287. doi: 10.1016/j.sleep.2012.11.016
 123. Zhou Y, Yang R, Li C, Tao M. Sleep disorder, an independent risk associated with arterial stiffness in menopause. *Sci Rep.* 2017;7:1904. doi: 10.1038/s41598-017-01489-7
 124. Leigh JA, Alvarez M, Rodriguez CJ. Ethnic minorities and coronary heart disease: an update and future directions. *Curr Atheroscler Rep.* 2016;18:9. doi: 10.1007/s11883-016-0559-4
 125. Pearson-Stuttard J, Guzman-Castillo M, Penalvo JL, Rehm CD, Afshin A, Danaei G, Kyridemos C, Gaziano T, Mozaffarian D, Capewell S, et al. Modeling future cardiovascular disease mortality in the United States: national trends and racial and ethnic disparities. *Circulation.* 2016;133:967–978. doi: 10.1161/CIRCULATIONAHA.115.019904
 126. Lindley KJ, Aggarwal NR, Briller JE, Davis MB, Douglass P, Epps KC, Fleg JL, Hayes S, Itchhaporia D, Mahmoud Z, et al; American College of Cardiology Cardiovascular Disease in Women Committee and the American College of Cardiology Health Equity Taskforce. Socioeconomic determinants of health and cardiovascular outcomes in women: JACC review topic of the week. *J Am Coll Cardiol.* 2021;78:1919–1929. doi: 10.1016/j.jacc.2021.09.011
 127. Kaur G, Masket D, Reddy T, Revankar S, Satish P, Paquin A, Mulvagh S, O'Donoghue ML, Zieroth S, Farkouh M, et al. Socioeconomic disparities in women's cardiovascular health in the United States and Canada. *Can J Cardiol.* 2024;40:1056–1068. doi: 10.1016/j.cjca.2024.04.001
 128. Morris AA, Masoudi FA, Abdullah AR, Banerjee A, Brewer LC, Commodore-Mensah Y, Cram P, DeSilvey SC, Hines AL, Ibrahim NE, et al. 2024 ACC/AHA key data elements and definitions for social determinants of health in cardiology: a report of the American College of Cardiology/American Heart Association Joint Committee on Clinical Data Standards. *Circ Cardiovasc Qual Outcomes.* 2024;17:e000133. doi: 10.1161/hcq.0000000000000133
 129. Kaiser Family Foundation. Americans' challenges with health care costs. Accessed August 25, 2025. <https://kff.org/health-costs/americans-challenges-with-health-care-costs/>
 130. Piña IL, Jimenez S, Lewis EF, Morris AA, Onwuanyi A, Tam E, Ventura HO. Race and ethnicity in heart failure. *J Am Coll Cardiol.* 2021;78:2589–2598. doi: 10.1016/j.jacc.2021.06.058
 131. Turecamo SE, Xu M, Dixon D, Powell-Wiley TM, Mumma MT, Joo J, Gupta DK, Lipworth L, Roger VL. Association of rurality with risk of heart failure. *JAMA Cardiol.* 2023;8:231–239. doi: 10.1001/jamacardio.2022.5211
 132. Reshetnyak E, Ntamungiro M, Pinheiro LC, Howard VJ, Carson AP, Martin KD, Safford MM. Impact of multiple social determinants of health on incident stroke. *Stroke.* 2020;51:2445–2453. doi: 10.1161/STROKEAHA.120.028530
 133. National-level and state-level prevalence of overweight and obesity among children, adolescents, and adults in the USA, 1990–2021, and forecasts up to 2050. *Lancet.* 2024;404:2278–2298. doi: 10.1016/s0140-6736(24)01548-4
 134. Mohebi R, Chen C, Ibrahim NE, McCarthy CP, Gaggin HK, Singer DE, Hyle EP, Wasfy JH, Januzzi JL Jr. Cardiovascular disease projections in the United States based on the 2020 Census estimates. *J Am Coll Cardiol.* 2022;80:565–578. doi: 10.1016/j.jacc.2022.05.033
 135. Cushman M, Shay CM, Howard VJ, Jiménez MC, Lewey J, McSweeney JC, Newby LK, Poudel R, Reynolds HR, Rexrode KM, et al; on behalf of the American Heart Association. Ten-year differences in women's awareness related to coronary heart disease: results of the 2019 American Heart Association National Survey: a special report from the American Heart Association. *Circulation.* 2021;143:e239–e248. doi: 10.1161/CIR.0000000000000907
 136. Vogel B, Acevedo M, Appelman Y, Bairey Merz CN, Chieffo A, Figtree GA, Guerrero M, Kunadian V, Lam CSP, Maas A, et al; The Lancet Women and Cardiovascular Disease Commission: reducing the global burden by 2030. *Lancet.* 2021;397:2385–2438. doi: 10.1016/s0140-6736(21)00684-x

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