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GUIDELINES FOR MANAGEMENT OF HIGH-RISK AFRICAN AMERICANS WITH MULTIPLE CARDIOVASCULAR RISK FACTORS: RECOMMENDATIONS OF AN EXPERT CONSENSUS PANEL

African Americans have higher rates of cardiovascular disease (CVD) than do Caucasians, which contributes significantly to their reduced life expectancy. Most African American adults have at least one major risk factor for CVD. Nonetheless, African Americans are often underdiagnosed and undertreated, despite presenting to the healthcare system late in their course, often after a CVD event. Patients with multiple risk factors have a CVD risk far greater than the sum of their individual risks. Metabolic syndrome tends to be clustered to a greater degree in African American women. Aggressive management of African Americans is necessary.

In this report, we provide guidelines for the management of high-risk African Americans. For each individual risk factor, we address existing data and guidelines in the general population, existing data in African Americans, and proposed guidelines for African Americans based on evidence or extrapolation. In particular, for elevated cholesterol and blood pressure, evidence is emerging that lower is better, so aggressive management strategies are necessary. For dyslipidemia, statins alone will generally reach the goal, but for hypertension, multiple drugs are usually necessary. We conclude that further research in African Americans is necessary to complete the totality of evidence. (Ethn Dis. 2007;17:214-220)

Key Words: African American, Cardiovascular Disease, Multiple Risk Factors

From the Department of Medicine, David Geffen School of Medicine at UCLA and the Minority Health Institute, Los Angeles, California (RAW); Department of Internal Medicine, Wayne State University School of Medicine, Detroit, Michigan (JMF); Morehouse School of Medicine, Atlanta, Georgia (JRG); Department of Biomedical Science and Center of Excellence, Florida Atlantic University (WRS, CHH), Departments of Medicine & Epidemiology and Public Health, University of Miami Miller School of Medicine (CHH), Department of Preventive Medicine, Nova Southeastern University (CHH), Boca Raton, Florida.

Address correspondence and reprint requests to Charles H. Hennekens, MD, DrPH; 2800 South Ocean Blvd, PH-A; Boca Raton, FL 33432; 561-393-8845; profchhmd@ prodigy.net Richard Allen Williams, MD; John M. Flack, MD, MPH; James R. Gavin, III, MD, PhD; Wendy R. Schneider, RN, BSN, MSN; Charles H. Hennekens, MD, DrPH

INTRODUCTION

Cardiovascular disease (CVD) is and will remain the leading killer in the United States despite remarkable declines in mortality during the latter part of the 20th century.¹ Among the 35 million African Americans (12.3% of the US population), the average life expectancy is ≈ 69 years vs 76 in the general population (65.0 vs 73.2 in men and 73.9 vs 79.8 in women).² Cardiovascular disease (CVD) is a major contributor to this disparity, which has persisted unchanged during the 20th century.³ In 1995, age-standardized death rates from CVD were $\approx 40\%$ higher in African Americans (154 per 100,000) than in Caucasians (114 per 100,000).

African Americans have higher levels of individual risk factors, most notably smoking but also obesity, hypertension, and diabetes mellitus. African American multiple risk factor patients (AAMRP) often are underdiagnosed and undertreated, and they present to the healthcare system late in their course, often after a CVD event.

In this report, we provide guidelines for management of the AAMRP. The individual characteristics contributing to the diagnosis of the AAMRP are identified (Table 1), as well as other criteria (Table 2).⁴ For each individual risk factor, we address five issues: 1) existing data in the general population; 2) existing guidelines for the general population; 3) existing data on African Americans; and 4) proposed guidelines for African Americans, whether based on evidence or extrapolation. We conclude that further research in African In 1995, age-standardized death rates from CVD were $\approx 40\%$ higher in African Americans (154 per 100,000) than in Caucasians (114 per 100,000).

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ESTABLISHED RISK FACTORS

Hypertension

Hypertension affects >50 million people (20% of the US population) and increases risk of stroke, myocardial infarction, and CVD death. The costs are estimated to be >\$23 billion in medicines, health services, and loss of productivity.⁵ The Seventh Joint National Committee (JNC 7) Report on Prevention, Evaluation, and Treatment of High Blood Pressure⁶ estimates that 27% of treated hypertensives are at goal, and this ratio is lower in African Americans.

In the third National Health and Nutrition Evaluation Survey, the prevalences of hypertension in African American men and women were 35% and 34.2%, respectively, compared to 24.4% and 19.3% in Caucasians.⁷ Death rates from hypertension are >350% higher in African Americans than in Caucasians.⁸ African Americans have earlier onset, higher prevalence, and greater severity

Table 1. Common characteristics of African American multiple risk patients (AAMRP)

Systolic blood pressure ≥130 mm Hg and/or diastolic blood pressure ≥85 mm Hg
Left ventricular hypertrophy
Family history of type 2 diabetes mellitus
History of gestational diabetes mellitus
Birth of infant >9 lbs
Fasting plasma glucose concentration 110 to 125 mg/dL (6.1 to 6.9 mmol/L)
Serum triglyceride concentrations \geq 150 mg/dL (\geq 1.69 mmol/L)
Serum low-density lipoprotein cholesterol >140 mg/dL (>3.62 mmol/L)
Serum high-density lipoprotein cholesterol <40 mg/dL (<1.03 mmol/L) in men
Central (abdominal, visceral) obesity
Waist:hip ratio >.8 in women, >.95 in men
Body mass index >27 kg/m ²
Cigarette smoking
Serum homocysteine >15 µmol/L
Serum plasminogen activator inhibitor-1 >15 IU/L
History of clot formation
Polycystic ovary syndrome
Acanthosis nigricans
Gout or serum uric acid >8.5 mg/dL (>.51 mmol/L)

of hypertension, with double the risk of heart failure and CHD, and a five-fold increased risk of fatal stroke and end-stage renal disease (ESRD).^{9,10}

The Jackson Heart Study is a prospective cohort study of African Americans to identify risk factors for CVD, with emphasis on sequelae of hypertension (left ventricular hypertrophy [LVH], heart failure, CHD, stroke, and renovascular disease).¹¹

Left ventricular hypertrophy (LVH) is an independent risk factor for sudden

cardiac death¹² and usually a consequence of hypertension. In the Framingham Heart Study, patients with LVH had a five to six times higher risk of sudden death.¹² Left ventricular hypertrophy (LVH) is more common in African Americans and increases risks of arrhythmias and silent but lethal ischemic events.¹³

Diabetes Mellitus

Diabetes mellitus affects 10.3 million Americans, and another 5.4 million

Category	System	Abnormal If Any of These Criteria Are Met
1	Body weight/fat	Body weight \geq 120% of ideal adjusted for height, frame, sex
	distribution	Body mass index (BMI, kg/m ²) \geq 27.8 (men) or \geq 27.3 (women)
		Waist:hip ratio \geq .95 (men) or \geq .85 (women)
2	Blood pressure	Systolic ≥130 mm Hg, diastolic ≥85 mm Hg
3	Glucose tolerance†	Impaired fasting plasma glucose indicated by plasma glucose
		110–126 mg/dL (6.1 to 7.0 mmol/L in asymptomatic person
		(person could have diabetes if ≥126 mg/dL [≥7.0 mmol/L])
		Impaired glucose tolerance indicated by plasma glucose 2-hour
		post oral glucose challenge 140–199 mg/dL (7.8 to 11.0 mmol/L)
		in asymptomatic person (person could have diabetes if
		≥200 mg/dL [≥11.1 mmol/L])
4	Lipids	Fasting serum triglycerides >150 mg/dL (>1.69 mmol/L)
		High-density lipoprotein cholesterol <40 mg/dL (<1.03 mmol/L)
5	Family history	Positive history of type 2 diabetes mellitus or premature
	. ,	cardiovascular disease in first-degree relative

Table 2. Suggested approach to identification* of the insulin resistance syndrome

* A patient who satisfies criteria for abnormalities in at least two of the five categories is likely to be insulin resistant and may have insulin resistance syndrome.

† See reference 4.

are undiagnosed. It is the fourth leading cause of death in African Americans, compared with seventh for the general population. The American Diabetes Association estimates that 2.8 million African Americans (13%) have diabetes, predominantly type 2.¹⁴ In middle age (45–64 years), African Americans have a 51% higher prevalence of type 2 diabetes than do Caucasians. Age-adjusted death rates in diabetes were higher for African American men (117%) and women (167%) than for Caucasians.

Impaired fasting glucose is a precursor to diabetes, characterized by levels between 110 to 126 mg/dL (6.1 to 7.0 mmol/L) (Table 1). The prevalence in African Americans is 7.0%.¹⁵ Approximately 50% to 60% of individuals with impaired fasting glucose also have impaired glucose tolerance.

Dyslipidemia

Statins significantly lower total cholesterol, low-density lipoprotein (LDL) cholesterol, and triglycerides as well as increase high-density lipoprotein (HDL) cholesterol. In large-scale randomized trials, statins significantly reduce risks of myocardial infarction, stroke, cardiovascular mortality, and total mortality.¹⁶⁻²¹ In the National Cholesterol Education Program III, the primary target is LDL cholesterol. Secondary targets include HDL cholesterol and triglycerides. Recently, these guidelines have been updated to lower targets on the basis of randomized trials that show greater benefits of higher dose of statins compared with usual dose.²²

Obesity

Approximately 97 million adults in the United States are obese or overweight.²³ Obesity leads to diabetes, hypertension, and dyslipidemia. Abdominal (central) obesity is particularly associated with insulin resistance and is more atherogenic than gluteofemoral obesity.

African American women have double the rate of abdominal obesity than

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Caucasians (Figure 1).²⁴ Abdominal obesity is assessed by the waist:hip ratio or waist circumference (Table 3).²³ In randomized trials of weight loss of short duration, psychological benefits include improved quality of life, feeling better, looking better, and having more energy. Further research is needed in African American women, including randomized trials of longer duration testing the benefits and risks of newer pharmacologic therapies for obesity.

Physical Inactivity

Physical inactivity is an independent risk factor for CVD. Individuals who increase their level of physical activity to modest frequency and low intensity experience significant reductions in risks of CVD.²⁵

Cigarette Smoking

Cigarette smoking is the leading avoidable cause of premature cancer and total deaths in the United States.²⁶ Current smokers have double the risk of CVD mortality compared with nonsmokers, and cessation yields benefits that equate to those of the nonsmoker within a few years. Beneficial changes begin within months, even in the elderly.²⁷

African American men smoke more than Caucasians but consume fewer cigarettes per day. African American and Caucasian women smoke at approximately equal rates.^{28,29}

RECOMMENDATIONS FOR MANAGEMENT OF THE AAMRP

Therapeutic Lifestyle Changes

Therapeutic lifestyle changes confer large and more than additive CVD benefits. As regards dietary changes, recommendations include saturated fat <7% of total calories, cholesterol <200 mg/d, plant stanols and sterols, and foods with viscous (soluble) fiber.³⁰ Dietary advice includes distribution of meals evenly throughout the day, avoid-

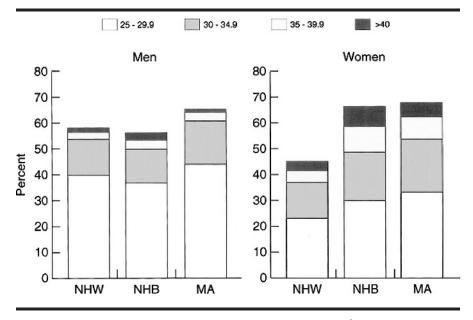


Fig 1. Prevalence of overweight body mass index $(BMI/kg/m^2)$ in US by sex and ethnicity, 1988–1994. MA=Mexican American; NHB=non-Hispanic Black; NHW=non-Hispanic White. Three levels of obesity: Class I BMI=30–34.9; Class II BMI=35–39.9; Class III BMI \geq 40 (Data derived from Reference 24, p 44, Table 6)

ance of "unconscious" eating, and reduction of daily intake by 200 to 500 calories to achieve modest weight loss. Patients should be followed for compliance and weight loss.

Hypertension

African American patients present at younger ages, with higher blood pressure levels and greater target organ damage (eg, LVH, reduced renal function). Current JNC 7 guidelines⁶ are based largely on Caucasian populations, although racial and ethnic minorities are discussed. Recently, guidelines for African Americans³¹ include:

- Combination therapy
- Target blood pressure of 130/ 80 mm Hg for those with heart disease, diabetes, or kidney disorders
- Include angiotensin-converting enzyme (ACE) inhibitors or angiotensin receptor blockers in combination therapy for patients with diabetes or renal disease

Randomized trials indicate that pharmacologic treatment of systolic blood pressures >140 mm Hg or diastolic pressures >90 mm Hg yield 4to 6 mm Hg reductions. Over three to five years, these reductions produce

Table 3. Disease risk relative to normal weight and waist size²³

			Disease Risk by Waist Size			
			Men \leq 102 cm (\leq 40 in.)	Men >102 cm (>40 ir		
Weight	BMI (kg/m ²)	Obesity Class	Women ≤88 cm (≤35 in.)	Women >88 cm (>35 in.)		
Underweight	<18.5					
Normal	18.5-24.9					
Overweight	25.0-29.9		Increased	High		
Obesity	30.0-34.9	I	High	Very high		
,	35.0-39.9	II	Very high	Very high		
Extreme obesity	>40	111	Extremely high	Extremely high		

Table 4.	Strategies	for	reducing	insulin	resistance

Nonpharmacologic	American Heart Association Step 1 or 2 diet
	Weight loss through hypocaloric diet (\downarrow initial body weight by 10% to 15% and maintain)
	Regular exercise (30 minutes of modest aerobic exercise at least 3 or 4 days, week, aim for 70% maximum heart rate)
	Smoking cessation
	Counseling
Pharmacologic	Hyperlipidemia (aim for total cholesterol ≤200 mg/dL [≤5.17 mmol/L], LDL- C ≤ 130 mg/dL [≤3.36 mmol/L]*, and HDL-C >35 mg/dL [>0.91 mmol/L] in men or >45 mg/dL [>1.16 mmol/L] in women) Elevated LDL-C: statin, resin, niacin
	Elevated triglycerides: gemfibrozil, fenofibrate, niacin
	Elevated LDL-C and triglycerides: statin and fibric acid derivative, resin, niacin, or high-dose statin
	Hypertension (aim for blood pressure ≤130/85 mm Hg)
	ACE inhibitors, α_1 -receptor blockers, calcium channel blockers, angiotensin II receptor antagonists
	Consider aspirin 81–325 mg/d
	Consider vitamin E 400 IU/d
	Insulin-sensitizing drugs: theoretical but unproven benefit as primary therapy for insulin resistance, except to treat hyperglycemia

 \ast LDL-C goal is ${<}100$ mg/dL (${<}2.59$ mmol/L) in patients with coronary artery disease, peripheral vascular disease, or stroke.

 $\label{eq:ACE} ACE=angiotensin-converting enzyme; \ \mbox{HDL-C}=\mbox{high-density lipoprotein cholesterol; \ \mbox{LDL-C}=\mbox{low-density lipoprotein cholesterol.}$

lower risks of stroke (42%), vascular death (20%), and CHD (14%).³² Drug selection should be governed by compelling indications, contraindications, and clinical judgment. Nonpharmacologic treatment (eg, weight loss, salt restriction) and drug therapy of patients' high-normal elevated diastolic blood pressure (85 to 89 mm Hg) reduces the rate of development of hypertension.

Racial differences practically disappear in combination therapies that most patients need to achieve blood pressure control. ACE inhibitors are the first choice in patients with metabolic syndrome or diabetes.33 Indeed, ACE inhibitors and angiotensin receptor blockers affect the renin-angiotensin system and provide target-organ protection superior to other agents, even when blood pressure lowering is similar.^{34–38} Some data suggest that African Americans may have a less beneficial response to ACE inhibition. Nonetheless, ACE inhibitors have beneficial effects on kidney function and CHF. In multidrug regimens, virtually no difference in blood pressure lowering is seen in initial therapy with an ACE inhibitor or calcium blocker.³⁸ In the African American Study of Kidney Disease and Hypertension trial, ramipril produced a 36% slower mean decline in glomerular filtration rates over three years and a 38% reduction in the risk of renal clinical endpoints among African Americans with hypertensive renal disease.³⁸

Glucose Intolerance/ Diabetes Mellitus

The American Diabetes Association Standards of Care³⁹ outlines management of glucose intolerance and addresses criteria for screening obese African Americans. Randomized trials in patients with type 1^{40} or type 2^{41} diabetes indicate beneficial effects of lifestyle modifications, such as weight loss and exercise, as well as aspirin, ACE inhibitors, and a variety of hypoglycemic agents (Table 4). In patients with type 2 diabetes, sulfonylurea or insulin therapy significantly reduced the risk of microvascular complications (eg, retinopathy, nephropathy) by 25%.⁴¹

Dyslipidemia

Lowering LDL cholesterol to <100 mg/dL (<2.59 mmol/L), with an optimal goal of <70 mg/dL, is the target for patients with prior myocardial infarction or stroke, peripheral vascular disease, 10-year CHD risk ≥20% or higher, or diabetes. For patients between 10% and 19%, the LDL cholesterol goal is <130 mg/dL (<3.36 mmol/L) with an optimal goal of <100 mg/dL. For individuals whose 10-year CHD risk is <10%, the LDL cholesterol goal is <160 mg/dL (<4.14 mmol/L). These 10-year risks are based on Framingham risk scores from a primarily Caucasian population. African Americans have higher absolute risks for any given Framingham risk score.

Obesity

The US National Heart, Lung and Blood Institute guidelines for clinicians include dietary therapy, physical activity, drug therapy, and if needed, obesity surgery (Figure 2).²³ All individuals should have body mass index (BMI) and obesity-related disease risks assessed and routinely monitored. For overweight patients (Table 3), weight maintenance may suffice if the patient has no additional risk factors and is resistant to weight reduction. Weight reduction is indicated if patients are obese (BMI $>30 \text{ kg/m}^2$) and especially with risk factors and increasing severity of obesity. Recommended dietary changes for all patients whose BMI is $>25 \text{ kg/m}^2$ include caloric restriction and adherence to the US Department of Agriculture's food pyramid guidelines that call for increased intake of fruits, vegetables, and grains.

Physical Inactivity

Low to moderate physical activity, defined as a 20-minute walk every other day, is associated with a 35% to 55% decrease in CVD in women, predominantly Caucasian.²⁵ Regular physical activity well into old age can be achieved if focused on an enjoyable activity, such

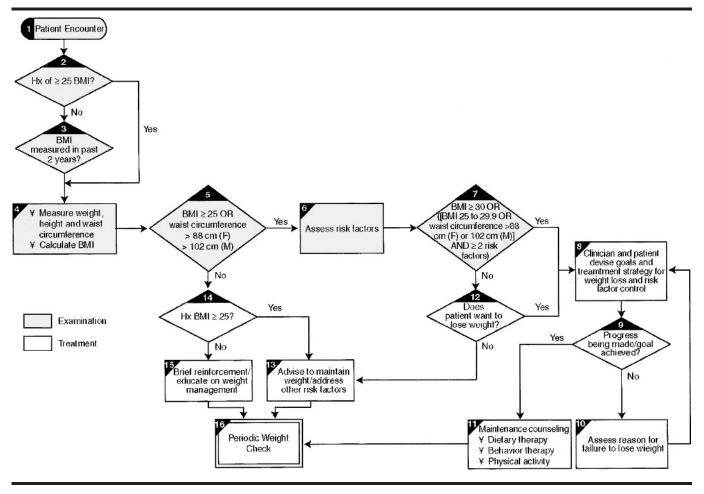


Fig 2. Treatment algorithm developed by the National Heart, Lung, and Blood Institute (1998) for overweight and obesity. Initial overall assessments for other conditions and diseases are not reflected. Boxes 1–7, examination; Boxes 8–15 treatment²³

as dancing, gardening, or dog walking, performed for sufficient duration on a regular basis.³¹

Cigarette Smoking

Smoking cessation has a 50% to 70% success rate for patients hospitalized with CHD. African Americans, have lower rates of quitting.⁴² Further research is needed to determine whether nicotine intake, biological factors, or possibly the preference for mentholated cigarettes affects nicotine dependence and ability to abstain.⁴³

Smoking in African Americans should be considered an addictive behavior, and younger men and women, especially pregnant women, should be targeted. Outreach through community or church-based programs that offer individual counseling and culturally appropriate self-help materials may be effective.

CONCLUSION

In the United States, African Americans have significantly higher risks of CVD than do Caucasians. Most middle-aged African Americans have at least one risk factor, many of which contribute to metabolic syndrome. Most are identifiable and modifiable even in early life.

Innovative approaches are necessary to improve compliance with therapeutic lifestyle changes.⁴⁴ For the general population, dropout rates for most physical activity programs are 50% within three to six months. Even cardiac rehabilitation patients, regardless of severity of the illness, dropout at a rate of 50% within a year. Group cohesion and social support are important to decrease dropout rates.⁴⁵

Informal care in nontraditional settings, as well as alternative and complementary medicine, may be more a culturally acceptable and cost-effective means to maintain and promote health for African Americans...

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Informal care in nontraditional settings, as well as alternative and complementary medicine, may be a more culturally acceptable and cost-effective means to maintain and promote health for African Americans, effectively overcoming barriers to accessing traditional care (financial, transportation, organizational and linguistic barriers) and offering ongoing education and support.⁴⁶

Cultural differences between patients and providers may affect adherence and outcomes. Cultural competence requires provider sensitivity to African American communication methods, belief systems, and the impact of their culture on healthcare relationships.⁴⁷

In addition to therapeutic lifestyle changes, drug therapies of proven benefit should also be used. For dyslipidemia and hypertension, lower is better, so aggressive management strategies for high-risk AAMRP would be beneficial. While for dyslipidemia, statins alone will generally reach the goal, for hypertension multidrug therapies are usually necessary.

Efforts are needed to increase the number of trained minority health professionals and to improve crosscultural interaction skills of all healthcare providers. Most African Americans are not treated by African American healthcare professionals. The following four steps outline ethnic-sensitive services⁴⁶:

- 1) Develop access to services (bilingual, bicultural staff)
- 2) Learn others' culture to tailor interventions
- Modify services (integrate traditional medicine with formal Westernized medicine)
- Initiate an appropriate organization development model/specialized program model (example: faithbased health care delivery model)

The current reimbursement environment does not facilitate issues of diagnosis, treatment, and overall access to health care for many African Americans. This is likely to be a factor in the disparities between African Americans and Caucasians. Healthcare providers have an urgent challenge to diagnose and aggressively manage the AAMRP, on the basis of the best clinical information available. Further research is needed to acquire a sufficient totality of evidence to achieve evidence-based changes in public health policy.

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REFERENCES

- National Vital Statistics Report. Number of deaths from 113 selected causes by age: United States, 1999 (Table 9). September 21, 2001, Vol 49, No 8. Available at: www.cdc.gov/ nchs/fastats/heart.htm. Accessed on April 18, 2002.
- National Heart, Lung, and Blood Institute. Morbidity and Mortality Chartbook 1996.
 Washington, DC: US Public Health Service; May 1996.
- Levine RS, Foster JE, Fullilove RE, et al. Black-White inequalities in mortality and life expectancy, 1933–1999: implications for Healthy People 2010. *Public Health Rep.* 2002;116:474–483.
- Gavin JRIII. New classification and diagnostic criteria for diabetes mellitus. *Clin Cornerstone*. 1998;1:1–12.
- Dustan HP, Rocella EJ, Garrison HH. Controlling hypertension: a research success story. Arch Intern Med. 1996;156:1926–1935.
- Chobanian AV, Bakris GL, Black HR, et al. and the National High Blood Pressure Education Program Coordinating Committee. The seventh report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure: the JNC 7 report. *JAMA*. 2003;289(19): 2560–2572.
- Burt VL, Whelton P, Rocella EJ, et al. Prevalence of hypertension in the US adult population: results from the Third National Health and Nutrition Examination Survey, 1988–1991. *Hypertension*. 1995;25:305–313.

- Williams RA. Cultural diversity in medicine. In: Williams RA ed. *Humane Medicine: A New Paradigm in Medical Education and Health Care Delivery.* Vol 2. Philadelphia, Pa: Lippincott Williams & Wilkins, 2001;13.
- Klag MJ, Whelton PK, Randall BL, Neaton JD, Brancati FL, Stamler J. End-stage renal disease in African American and White men: 16-year MRFIT findings. *JAMA*. 1997;277: 1293–1298.
- Ofili EO. Managing high blood pressure and cardiovascular disease in Blacks. Assoc Black Cardiol Digest Urban Cardiol. 1995;2: 220–225.
- Taylor HA Jr, Burchfield C, Fletcher B, et al. The Jackson Heart Study. Assoc Black Cardiol Digest Urban Cardiol. 1999;6:27–33.
- Kannel WB. Prevalence and natural history of electrocardiographic left ventricular hypertrophy. Am J Med. 1983;75(suppl 3A):4–11.
- Liao Y, Cooper RS, McGhee DL, Mensah GA, Ghali JK. The relative effects of left ventricular hypertrophy, coronary artery disease, and ventricular dysfunction on survival among Black adults. *JAMA*. 1995;273:1592–1597.
- American Diabetes Association. Diabetes among African Americans. Available at: http://www.diabetes.org/main. Accessed on December 3, 2002.
- Harris MI, Flegal KM, Cowie CC, et al. Prevalence of diabetes, impaired fasting glucose, and impaired glucose tolerance in US adults. The Third National Health and Nutrition Examination Survey, 1988–1994. *Diabetes Care*. 1998;21:518–524.
- Downs JR, Clearfield M, Weis S, et al. Primary prevention of acute coronary events with lovastatin in men and women with average cholesterol levels: results of AFCAPS/TexCAPS. Air Force Coronary Atherosclerosis Prevention Study. JAMA. 1998;279:1615–1622.
- Hebert PR, Gaziano JM, Chan KS, Hennekens CH. Cholesterol lowering with statin drugs, risk of stroke, and total mortality: an overview of randomized trials. *JAMA*. 1997; 278:313–321.
- The Long-Term Intervention with Pravastatin in Ischemic Heart Disease (LIPID) Study Group. Prevention of cardiovascular events and death with pravastatin in patients with coronary heart disease and a broad range of initial cholesterol levels. N Engl J Med. 1998;339:1349–1357.
- Sacks FM, Pfeffer MA, Moyé LA, et al. for the Cholesterol and Recurrent Events Trial Investigators. The effect of pravastatin on coronary events after myocardial infarction in patients with average cholesterol levels. *N Engl J Med.* 1996;335:1001–1009.
- 20. Scandinavian Simvastatin Survival Study Group. Randomized trial of cholesterol lowering in 4444 patients with coronary heart

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disease: the Scandinavian Simvastatin Survival Study (4S). *Lancet*. 1994;344:1383–1389.

- Shepherd J, Cobbe SM, Ford I, et al. for the West of Scotland Prevention Study Group. Prevention of coronary heart disease with pravastatin in men with hypercholesterolemia. *N Engl J Med.* 1995;333:1301–1307.
- 22. Grundy SM, Cleeman JI, Merz CN, et al. for the Coordinating Committee of the National Cholesterol Education Program. Implications of recent clinical trials for the National Cholesterol Education Program Adult Treatment Panel III guidelines. *Circulation*. 2004;110(2):227–239.
- 23. National Heart, Lung, and Blood Institute. National Heart, Lung, and Blood Institute Obesity Education Initiative. Clinical guidelines on the identification, evaluation, and treatment of overweight and obesity in adults. June 1998. Available at: www.nhlbi.nih. gov/guidelines/obesity/e_txtbk/txgd/algorthm/ algorthm.htm. Accessed on April 9, 2002.
- Flegal KM, Carroll MD, Kuczmarksi RJ, Johnson CL. Overweight and obesity in the United States: prevalence and trends, 1960– 1994. Int J Obes Relat Metab Disord. 1998;22: 39–47.
- Manson JE, Hu FB, Rich-Edwards JW, et al. A prospective study of walking as compared with vigorous exercise in the prevention of coronary heart disease in women. *N Engl J Med.* 1999; 341:650–658.
- Peto R, Lopez AD, Boreham J, Thun M, Heath CJr, Doll R. Mortality from smoking worldwide. *Br Med Bull.* 1996;52:12–21.
- LaCroix AZ, Land J, Scherr P, et al. Smoking and mortality among older men and women in three communities. *N Engl J Med.* 1991;324: 1619–1625.
- Fiore MC, Novotney TE, Pierce JP, Hatziandreu EJ, Patel KM, Davis RM. Trends in cigarette smoking in the United States: the changing influence of gender and race. *JAMA*. 1989;261:49–55.
- Taylor HA, Mickel MC, Chaitman BR, Sopko G, Cutter GR, Rogers WJ. Long-term survival of African Americans in the Coronary Artery Surgery Study (CASS). J Am Coll Cardiol. 1997;29:358–364.
- Eidelman RS, Lamas GA, Hennekens CH. The new National Cholesterol Education Program guidelines: clinical challenges for more widespread therapy of lipids to treat and prevent coronary heart disease. *Arch Intern Med.* 2002;162:2033–2036.
- 31. Douglas JG, Bakris GL, Epstein M, et al. Hypertension in African Americans Working Group of the International Society on Hypertension in Blacks. Management of high blood pressure in African Americans: consensus statement of the Hypertension in African Americans Working Group of the Interna-

tional Society on Hypertension in Blacks. Arch Intern Med. 2003;163(5):525-541.

- 32. Collins R, Peto R, MacMahon S, et al. Blood pressure, stroke, and coronary heart disease. Part 2. Short-term reductions in blood pressure: overview of randomized drug trials in their epidemiological context. *Lancet.* 1990;335:827–838.
- 33. Bakris GL, Williams M, Dworkin L, et al. for the National Kidney Foundation Hypertension and Diabetes Executive Committees Working Group. Preserving renal function in adults with hypertension and diabetes: a consensus approach. *Am J Kidney Dis.* 2000;36: 646–661.
- Fogo AB. Progression and potential regression of glomerulosclerosis. *Kidney Int.* 2001;59: 804–819.
- 35. Heart Outcome Prevention Evaluation (HOPE) Study Investigators. Effects of ramipril on cardiovascular and microvascular outcomes in people with diabetes mellitus: results of the HOPE study and MICRO-HOPE study. *Lancet.* 2000;355:253–259.
- 36. Lewis EJ, Hunsicker IG, Bain RP, Rohde RD. The effect of angiotensin-converting enzyme inhibition on diabetic nephropathy. The Collaborative Study Group. *N Engl J Med.* 1993;329:1456–1462.
- UK Prospective Diabetes Study (UKPDS) Group. Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes (UKPDS 38). *BMJ*. 1998;317:703–713.
- Agodoa LY, Appel L, Bakris GL, et al. Effect of ramipril vs amlodipine on renal outcomes in hypertensive nephrosclerosis. *JAMA*. 2001; 285:2719–2728.
- American Diabetes Association. Standards of medical care in diabetes-2006. *Diabetes Care*. 2006;29(suppl 1):S4–S42.
- The DCCT Research Group. Effect of intensive therapy on the development and progression of diabetic nephropathy in the Diabetes Control and Complications Trial (DCCT). *Kidney Int.* 1995;47:1703–1720.
- 41. UK Prospective Diabetes Study (UKPDS) Group. Intensive blood glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33). *Lancet*. 1998;352:837–853.
- Royce JM, Hymowitz N, Corbett K, et al. Smoking cessation factors among African Americans and Whites. COMMIT Research Group. *Am J Public Health*. 1993;83(2):220–226.
- Mazas CA, Wetter DW. Smoking cessation interventions among African Americans: research needs. *Cancer Control.* 2003;10(5): 87–89.
- 44. Haynes RB, McKibbon KA, Kanani R. Systematic review of randomized trials of

interventions to assist patients to follow prescriptions for medications. *Lancet*. 1996;348:383.

- Fraser SN, Spink KS. Examining the role of social support and group cohesion in exercise compliance. J Behav Med. 2002;25(3): 233–249.
- Chen, MS, Jr. Informal care and the empowerment of minority communities: comparisons between the USA and the UK. *Ethm Health*. 1999;4(3):139–151.
- 47. National Center for Primary Care at the Morehouse School of Medicine. CRASH Course in Cultural Competency Skills. Available at: http://www.msm.edu/NCPC/crash/ crash_powerpointpage.htm. Accessed on August 28, 2006.

AUTHOR CONTRIBUTIONS

- Design concept of study: Williams, Flack, Gavin, Hennekens
- Acquisition of data: Williams, Schneider, Hennekens
- Data analysis and interpretation: Williams, Flack, Gavin, Hennekens
- Manuscript draft: Williams, Flack, Gavin, Schneider, Hennekens
- Statistical expertise: Hennekens
- Acquisition of funding: Williams
- Administrative, technical, or material assistance: Williams, Flack, Gavin, Schneider, Hennekens

Supervision: Williams, Hennekens

APPENDIX: ADVISORY BOARD MEMBERS

Norman Anderson, PhD; Andre Artis, MD; Mandeep Bajaj, MD; Narenda C. Bhalodkar, MD; James L. Brown, JD; Ludlow Creary, MD, MPH; Charles Curry, MD; Mayer B. Davidson, MD; Joyleen Earle, MD; James M. Falko, MD; Larry E. Fields, MD, MBA; Om Ganda, MD; Gairy Hall, MD; Joyce Harp, MD; Patricia Hebert, PhD; Terry A. Jacobson, MD; Frank James, MD; Michelle N. Johnson, MD; Edith Irby Jones, MD; Udaya Kabadi, MD; Norman N. Kaplan, MD; Mahmood Khan, MD; B. Waine Kong, PhD, JD; Robert Levine, MD; Philip Levy, MD; Anthony McCall, MD, PhD; Calvin McLarin, MD; Al Oberman, MD; Ola Odugbesan, MD; Kwame Osei, MD; Lawrence S. Phillips, MD; Marc Pohl, MD; James L. Pool, MD; Otelio T. Randall, MD; James W. Reed, MD; Richard S. Rigmaiden, III, MD; Victor Roberts, MD; John Siebel, MD; Craig W. Spellman, PhD, DO; Anne E. Sumner, MD; Frank Svec, MD; Anne Taylor, MD; Malcolm P. Taylor, MD; Aaron I. Vinik, MD, PhD; Laurence Watkins, MD, MPH; and Karol Watson, MD, PhD.