

TABLE 1
Select major differences between 2013 and 2018 AHA/ACC cholesterol guidelines^{1,2} (cont'd)

Guideline parameter	2013 ²	2018 ¹
Severe hypercholesterolemia		
Nonstatin therapy	For adults 21-75 y who have an LDL-C level ≥ 190 mg/dL after maximizing statin therapy, adding a nonstatin drug can be considered to further lower the LDL-C level	Prescribe ezetimibe in patients 20-75 y who have an LDL-C level ≥ 190 mg/dL and who (1) achieve a $< 50\%$ reduction in LDL-C while receiving maximally tolerated statin therapy or (2) who have an LDL-C level ≥ 100 mg/dL (or both) Prescribe a PCSK9 inhibitor in patients who (1) have a baseline LDL-C level ≥ 220 mg/dL and (2) achieve an on-treatment LDL-C level ≥ 130 mg/dL while receiving maximally tolerated statin and ezetimibe therapy Prescribe a PCSK9 inhibitor in patients 30-75 y who have (1) heterozygous familial hypercholesterolemia and (2) an LDL-C level ≥ 100 mg/dL while taking maximally tolerated statin and ezetimibe therapy
Secondary prevention		
Very high-risk category of ASCVD	No equivalent recommendation	Adds specific recommendations for very high-risk patients
LDL-C targets	High- and moderate-intensity statin therapy recommendations did not specify LDL-C reduction targets but did recommend follow-up LDL-C testing for adherence to gauge adequacy of statin effect	Specifies importance of percentage reduction in LDL-C level when prescribing high ($\geq 50\%$) or moderate-intensity (30%-49%) statin therapy as well as follow-up LDL-C testing for adherence and to gauge effects of LDL-C lowering medication
Nonstatin therapy	Consider adding nonstatin therapy in adults at higher risk for ASCVD who are receiving maximally tolerated statin therapy with a less-than-anticipated therapeutic response Use nonstatin cholesterol-lowering drugs in adults who are candidates for statin therapy but are completely statin-intolerant	Very high-risk ASCVD patients should receive maximally tolerated statin therapy; if the LDL-C level remains ≥ 70 mg/dL, add ezetimibe before considering a PCSK9 inhibitor. If, despite ezetimibe, the LDL-C level is ≥ 70 mg/dL or non-HDL-C is ≥ 100 mg/dL, consider a PCSK9 inhibitor

ACC, American College of Cardiology; AHA, American Heart Association; ASCVD, atherosclerotic cardiovascular disease; CAC, coronary artery calcium; HDL, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; PCSK9, proprotein convertase subtilisin/kexin type 9.

¹Fifteen members of the 2013 guideline panel, including a vice chair, declared a conflict of interest. None of the authors of the 2018 guideline declared a conflict of interest.

²American College of Cardiology, American Heart Association, American Association of Cardiovascular and Pulmonary Rehabilitation, American Academy of Physician Assistants, Association of Black Cardiologists, American College of Preventive Medicine, American Diabetes Association, American Geriatrics Society, American Pharmacists Association, American Society for Preventive Cardiology, National Lipid Association, and Preventive Cardiovascular Nurses Association.

³Value statements address clinicians' and patients' need to consider the cost of drugs in determining the most appropriate treatment.

Aha Guidelines 2013 Criticism

**Maxine A. Papadakis, Stephen J.
McPhee, Michael W. Rabow**



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Clinical Lipid Management Jennifer G. Robinson, MD, 2023-10-11 Presents the newest thinking and data needed for clinicians to reduce atherosclerotic cardiovascular disease ASCVD risk This book provides detailed evidence based guidance for treating cholesterol to reduce ASCVD with a focus on recent ACC AHA guidelines an overview of evidence based treatment of other risk factors the management of genetic and acquired lipid disorders and complete efficacy and safety information on all available lipid lowering agents including clinical trial data and FDA approved labeling

Cardiovascular Diseases Y. Robert Li, 2015-02-25 Written in an accessible style and consistent format the book covers both the fundamentals and advances in the pharmacology of cardiovascular drugs as well as their integrated applications in the management of individual cardiovascular diseases Integrates fundamentals and recent advances regarding cardiovascular drugs blending basic and clinical sciences needed to effectively understand and treat cardiovascular diseases Facilitates understanding of drug action and mechanism by covering physiology pathophysiology and pharmacology Includes guidelines and algorithms for pharmacotherapeutic management of cardiovascular diseases Uses case presentations and study questions to enhance understanding of the material Serves as a resource for pharmaceutical and medical students and researchers interested in cardiovascular issues

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