

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

Axel Pressler, Josef Niebauer



Aha Guidelines 2013 Criticism:

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, 2015-04-06 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 Lipid Disorders: A Multidisciplinary Approach, Clinics Collections, 1e, (Clinics Collections) Joel J. Heidelbaugh, 2015-10-15 Clinics Collections Lipid Disorders draws from Elsevier's robust Clinics Review Articles database to provide multidisciplinary teams including general practitioners cardiologists endocrinologists pathologists cardiologists rheumatologists and hepatologists with practical clinical advice and insights on this highly prevalent condition and its comorbidities Clinics Collections Lipid Disorders guides readers on how to apply current primary research findings on lipid disorders to everyday practice to help overcome challenges and complications keep up with new and improved treatment methods and improve patient outcomes Areas of focus include lipids and biomarkers lipids and pharmacotherapy lipids and heart disease lipids and chronic disease lipids and pediatrics lipids and women's health and special considerations Each article begins with keywords and key points for immediate access to the most critical information Articles are presented in an easy to digest and concisely worded format Elsevier's Clinics Collections provide concise reviews of today's most prevalent conditions and significant medical developments Other Clinics Collections titles available include Type II Diabetes Mellitus Asthma Obesity and Pain Management Anesthesiology Critical Care Board Review George W. Williams, Navneet Kaur Grewal, Marc J. Popovich, 2019-10-21 Anesthesiology Critical Care Board Review fills a much needed niche offering a high yield review of content for the American Board of Anesthesiology's Critical Care board exam in a question and answer format Concise explanations and targeted references are provided for each question Arterial Disorders Adel Berbari, Giuseppe Mancia, 2015-03-19 As our knowledge about arterial disease is greatly expanding the aim of this book is to explore all aspects of arterial pathology including classification clinical manifestations pathogenesis and therapeutic options The discussion of

pathophysiologic mechanisms of arterial disease is wide ranging encompassing hemodynamic metabolic humoral inflammatory genetic and environmental factors Particular emphasis is placed on recent concepts such as the role of age associated arterial alterations in the initiation and progression of cardiovascular diseases in older persons the importance of mineral metabolism bone vascular interactions the clinical and prognostic significance of the renal resistive index retinal circulation toxemia of pregnancy as an arterial disease and the role of pulmonary vascular interaction in pulmonary hypertension and cross talk of macrocirculation and microcirculation in target organ involvement Evaluation procedures are carefully explained and the full range of currently available therapeutic options including lifestyle modifications and pharmacologic approaches are described and appraised **Yearbook of Pediatric Endocrinology 2014** K.

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