

Lipid Screening and Management

The following guideline recommends risk assessment, stratification, education, counseling and pharmacological interventions for the management of low-density lipoprotein cholesterol (LDL-C).

Eligible Population	Key Components	Recommendation and Level of Evidence										
<p>Males ≥ 35 years of age</p> <p>Females ≥ 45 years of age</p> <p>Males and Females age ≥ 20 years of age if risk factors</p>	Risk Assessment	<p>Screening: Initial fasting lipid profile (i.e., total, LDL-C, HDL-C, triglycerides); If in normal range, repeat at least every 4-6 years. [D]</p> <p>Treatment is based on presence of clinical atherosclerotic cardiovascular disease (ASCVD), and ASCVD risk factors. [A]</p> <table border="1"> <tr> <td>Clinical ASCVD: TIA, Stroke Angina, MI Acute Coronary Syndrome Peripheral arterial disease, aortic aneurysm Revascularization procedure</td> <td>ASCVD Risk Factors: LDL-C ≥ 190 mg/dL and age ≥ 20, not caused by drugs or underlying medical condition Diabetes mellitus type 1 or 2, age 40-75 years of age with LDL-C 70-189 mg/dL 10-year ASCVD risk ≥ 7.5% for ages 40-75 years</td> </tr> </table>	Clinical ASCVD: TIA, Stroke Angina, MI Acute Coronary Syndrome Peripheral arterial disease, aortic aneurysm Revascularization procedure	ASCVD Risk Factors: LDL-C ≥ 190 mg/dL and age ≥ 20, not caused by drugs or underlying medical condition Diabetes mellitus type 1 or 2, age 40-75 years of age with LDL-C 70-189 mg/dL 10-year ASCVD risk ≥ 7.5% for ages 40-75 years								
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	Risk Stratification	<p>Calculate¹ 10-year ASCVD risk for patients 40-75 years of age without clinical ASCVD, diabetes mellitus (type 1 or 2) or LDL-C ≥ 190 mg/dL [D]</p> <table border="1"> <tr> <td>Statin treatment benefit group Clinical ASCVD: Age ≤ 75 years In very high risk ASCVD (multiple events or 1 main event and multiple risk factors), if LDL-C remains ≥ 70 mg/dL, consider addition of ezetimibe to statin</td> <td>Statin dosing intensity² High-intensity [A]</td> </tr> <tr> <td>Clinical ASCVD: Age > 75 years</td> <td>Moderate-intensity [D]</td> </tr> <tr> <td>LDL-C ≥ 190 mg/dL, age ≥ 21 years If LDL-C remains ≥ 100 mg/dL, consider addition of ezetimibe to statin</td> <td>High-intensity [A]</td> </tr> <tr> <td>Diabetes mellitus (type 1 or 2) and age 40-75 years with LDL-C 70-189 mg/dL</td> <td>Moderate-intensity [A], can consider high-intensity if 10-year ASCVD risk ≥ 7.5% [D]</td> </tr> <tr> <td>10-year ASCVD risk ≥ 7.5% and age 40-75 years</td> <td>Moderate-to-high intensity [A]</td> </tr> </table>	Statin treatment benefit group Clinical ASCVD: Age ≤ 75 years In very high risk ASCVD (multiple events or 1 main event and multiple risk factors), if LDL-C remains ≥ 70 mg/dL, consider addition of ezetimibe to statin	Statin dosing intensity² High-intensity [A]	Clinical ASCVD: Age > 75 years	Moderate-intensity [D]	LDL-C ≥ 190 mg/dL, age ≥ 21 years If LDL-C remains ≥ 100 mg/dL, consider addition of ezetimibe to statin	High-intensity [A]	Diabetes mellitus (type 1 or 2) and age 40-75 years with LDL-C 70-189 mg/dL	Moderate-intensity [A] , can consider high-intensity if 10-year ASCVD risk ≥ 7.5% [D]	10-year ASCVD risk ≥ 7.5% and age 40-75 years	Moderate-to-high intensity [A]
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Education and risk factor modification	<p>Promote a healthy lifestyle throughout life.</p> <p>If indicated: smoking cessation, reduce excessive alcohol [A]</p> <p>Recommend a dietary pattern that emphasizes intake of vegetables, fruits, and whole grains; includes low-fat dairy products, poultry, fish, legumes, non-tropical vegetable oils and nuts; and limits intake of sweets, sugar-sweetened beverages and red meats [A]</p> <p>Engage in at least 150 minutes per week of accumulated moderate-intensity physical activity or 75 minutes per week of vigorous-intensity [B]</p>											
Pharmacologic interventions	<p>Women of childbearing age who are treated with statins should be counseled to use a reliable form of contraception and to stop the statin 1-2 months before pregnancy is attempted.</p> <p>Assess adherence and LDL-C percentage response to therapy with repeat lipid measurement 4-12 weeks after statin initiation or dose adjustment.</p> <p>Obtain baseline ALT. If normal, no routine monitoring for patients on statin therapy is required. LFT at physician discretion for patients with abnormal baseline ALT, liver disease or risk factors.</p> <p>For prolonged myalgias, consider dosage reduction or statin change. Check creatine kinase (CK) only if symptomatic muscle aches/weakness.</p> <p>For patient > 75 years, statin use should be at patient/physician discretion.</p> <p>If statins not tolerated, consider alternate medical therapy including ezetimibe or PCSK9 inhibitor.</p>											

¹ACC/AHA [ASCVD Risk Estimator Tool](#)

²University of Michigan Ambulatory Adult Screening Management of Lipids Guidelines Table 6. [Statin Dose Intensity and Equivalency Chart Table](#)

Levels of Evidence for the most significant recommendations: A = randomized controlled trials; B = controlled trials, no randomization; C = observational studies; D = opinion of expert panel

This guideline represents core management steps. It is based on Grundy SM, et.al. 2018 AHA/ACC/AACVPR/AAPA/ABC/ACPM/ADA/AGS/APhA/ASPC/NLA/PCNA guideline on the management of blood cholesterol: executive summary: a report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation*. 2019; 139:e1046-e1081. and based on Arnett DK, et.al., 2019 ACC/AHA guideline on the primary prevention of cardiovascular disease: executive summary: a report of the American College of Cardiology/American Heart Association Task Force on Clinic Practice Guidelines. *Circulation*. 2019; 140:e563-e595. Individual patient considerations and advances in medical science may supersede or modify these recommendations.