



Outpatient Lipid Management

2026 ACC/AHA Multisociety Dyslipidemia Guideline · PREVENT replaces the pooled cohort equations

Earlier & lower
for longer

STEP 1 · THE CPR FRAMEWORK *the guideline's own model for primary prevention*

C — CALCULATE

PREVENT-ASCVD, ages 30-79, LDL 70-189. Bands: **low** < 3% · **borderline** 3-<5% · **intermediate** 5-<10% · **high** ≥ 10%. Race dropped as an input; CKD, HbA1c and BMI added; reports **10- and 30-year** risk.

P — PERSONALIZE

Apply **risk enhancers** PREVENT misses: family history of premature ASCVD, LDL ≥ 160, metabolic syndrome, CKD, elevated Lp(a), South Asian ancestry, **inflammatory disease** (RA, lupus, psoriasis, HIV) and **reproductive history** (preeclampsia, early menopause, gestational diabetes).

R — RECLASSIFY & REASSESS

When the decision is still uncertain, **CAC scoring** reclassifies. Then reassess the treatment plan. **Age 30-59, low risk, LDL < 160 and 30-year risk < 10%: lifestyle counseling alone** is the Class 1 answer.

STEP 2 · GOALS ARE BACK *LDL and non-HDL-C, plus a percent reduction*

Risk category	LDL-C goal	non-HDL-C goal	Also
Very-high-risk ASCVD Class 1	< 55	< 85	High-intensity statin, ≥ 50% LDL reduction. apoB < 55. Most patients with ASCVD land here.
ASCVD, not very high risk	< 70	< 100	A minority of ASCVD patients.
Primary prevention, high (≥ 10%)	< 70	< 100	High-intensity: atorvastatin 40-80, rosuvastatin 20-40 (≥ 50% reduction).
Borderline 3-<5% or intermediate 5-<10%	< 100	< 130	Moderate: atorvastatin 10-20, rosuvastatin 5-10 (30-49%). Intermediate: should be considered; borderline: may be, after discussion. Use CAC if undecided.

Very high risk = multiple major ASCVD events, or one major event plus multiple high-risk conditions (age > 65, prior revascularization, smoking, diabetes, heart failure, hypertension, LDL > 100 on maximal statin + ezetimibe).

STEP 3 · WHO GETS THERAPY *and at what intensity*

Indication	Intensity and notes
Clinical ASCVD	High-intensity. Add ezetimibe, then a PCSK9i, to reach goal. Combination is the expectation, not the exception.
LDL ≥ 190	High-intensity. Treat as familial hypercholesterolemia until proven otherwise; no risk calculation needed.
Diabetes, CKD stage 3-4, or HIV age 40-75	Recommended regardless of LDL level. After 75, therapy <i>can be considered</i> alongside lifestyle.
Early intervention NEW	HeFH at diagnosis (including children 8-10), young adults with LDL ≥ 160 , or strong family history of premature ASCVD. Exists because 10-year risk is always low in the young and systematically under-treats them.

CAC · NOW A CLASS 1 TOOL *upgraded from 2a in 2018*

WHO TO SCAN

Men ≥ 40, women ≥ 45, at **intermediate** risk or **selected borderline** risk, when the treatment decision remains uncertain. Both the **absolute score and the age/sex/race percentile** carry prognostic weight.

HOW IT CHANGES THE PLAN

CAC 0 supports withholding or postponing therapy in a low- or intermediate-risk patient with no other comorbidity. **Any calcium argues for treating**, and a higher score argues for treating harder. **Incidental calcium on a non-gated CT now counts**, so do not ignore “coronary calcifications” on a chest CT report.

ORDER ONCE, AND ONCE ONLY *Lp(a), plus apoB when LDL misleads*

Test	When, and what the number means
Lp(a) Class 1, all adults	Measure at least once in every adult. Genetically fixed, so one result settles it. ≥ 125 nmol/L (50 mg/dL) is a risk enhancer, roughly 1.4× risk; ≥ 250 nmol/L (100 mg/dL) is ≥ 2×. It has no drug of its own yet: it intensifies LDL lowering and risk-factor control. Cascade test first-degree relatives. In ASCVD with high Lp(a) not at goal on maximal statin, add a PCSK9 mAb (Class 1).
apoB	Useful once LDL and non-HDL goals are met , and especially when TG ≥ 150 , in diabetes , or when achieved LDL < 70 . In those settings LDL-C understates particle number, so a patient “at goal” can still carry residual risk.

SPECIAL SITUATIONS *the ones that change the drug*

Situation	What to do, and why
TG ≥ 500	Fenofibrate to prevent pancreatitis. Never gemfibrozil with a statin (rhabdomyolysis). At ≥ 1000 the urgency is pancreatitis, not cardiovascular risk. Statin remains the foundation for ASCVD risk in high TG.
ASCVD, TG 150-499	Icosapent ethyl (REDUCE-IT). Pure EPA, not interchangeable with OTC fish oil. PROMINENT closed the door on fibrates for event reduction.
“Statin intolerant”	Mostly placebo: SAMSON found identical symptoms on placebo. Rechallenge with a different agent or alternate-day dosing before accepting the label , which otherwise costs decades of protection. True intolerance: bempedoic acid, ezetimibe, PCSK9i.
Pregnancy	Stop statins, ezetimibe, fibrates, niacin and PCSK9 inhibitors. Bile acid sequestrants only, as they are not absorbed. Discuss contraception before starting a statin.
Dose caps	eGFR < 30: rosuvastatin max 10 mg. Asian ancestry: start rosuvastatin 5 mg. HIV: pitavastatin , minimal antiretroviral interaction.

MONITORING & PITFALLS

- Recheck lipids **4-12 weeks after starting or changing**, then every 3-12 months. Not at goal is the trigger to add a second agent, not to wait another year.
- Do not monitor LFTs or CK routinely.** Baseline ALT, then check CK **only if muscle symptoms** appear.
- Exclude secondary causes before labeling primary dyslipidemia:** hypothyroidism, nephrotic syndrome, cholestasis, uncontrolled diabetes, alcohol, and drugs (thiazides, steroids, retinoids, antipsychotics).
- Intensity is percent reduction, not milligrams.** Each dose doubling adds only about 6%.

