

Applying the New ACC/AHA Cholesterol Guidelines in Practice

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Disclosure

Advisory Board: Novartis

Research Grant: DalCor

Stockholder: Centene; Walmart



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Learning Objectives

1. Understand guideline background and use in routine clinical practice.
2. Integrating subclinical atherosclerosis and risk-enhancing factors into practice.
3. Applying risk-based cholesterol targets to patients.



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Objective 1



Clinical Guidelines and How to Use Them in Clinical Practice

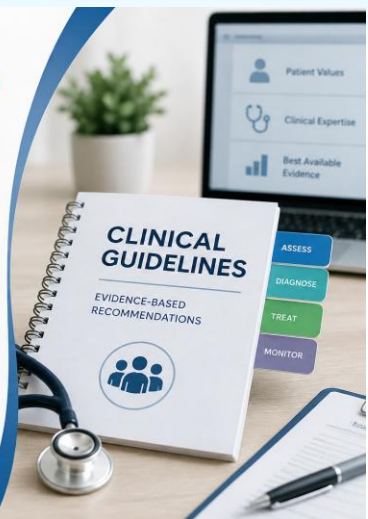


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2026 ACC/AHA/AACVPR/ABC/ACPM/ADA/AGS/A PhA/ASPC/NLA/PCNA Guideline on the Management of Dyslipidemia: A Report of the American College of Cardiology/American Heart Association Joint Committee on Clinical Practice Guidelines

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Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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What Are These Guidelines?



Guidelines are reference documents aimed at practicing clinicians to translate the best evidence into practice



Joint effort from multiple professional societies



123 pages of information—tables, figures, algorithms, etc...



Recommendations are graded, as is the evidence that supports them

Otto CM, et al. <https://doi.org/10.1161/CIR.0000000000001383>
Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Practical Applications

- Time is limited in clinic:
 - Calculating PREVENT takes time
 - Referencing multiple sources
 - Shared decision-making conversation
 - Risk discussion
- Can feel overwhelming
 - 68 tables, 18 figures

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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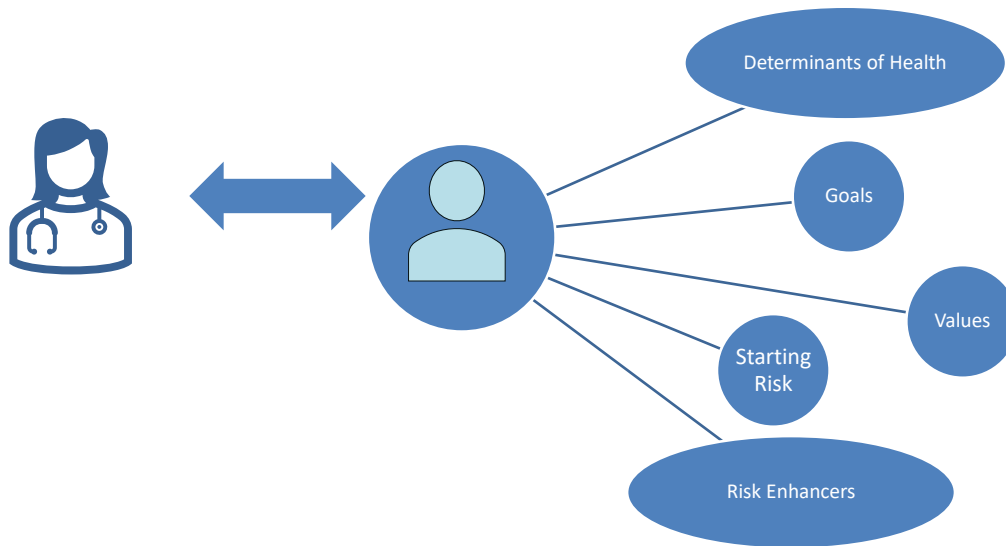
Make It Bite Sized

- Each visit as an opportunity
 - Lab testing and risk assessment
- Make risk personalization as useful as possible

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Shared Decision-Making



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Explaining Risk Is Confusing

Absolute Terms

- The risk of X is HIGH
- The risk of Y is LOW
- 1 in 100, every 6 people who has this done, etc...

Relative terms

- Compared to others your age, sex and race, your risk is Z.

Time

- In 5 years?, 10 years?, 30 years?

Although the benefit of risk reduction is related to absolute risk, people also think about relative risk.

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Cumulative Exposure

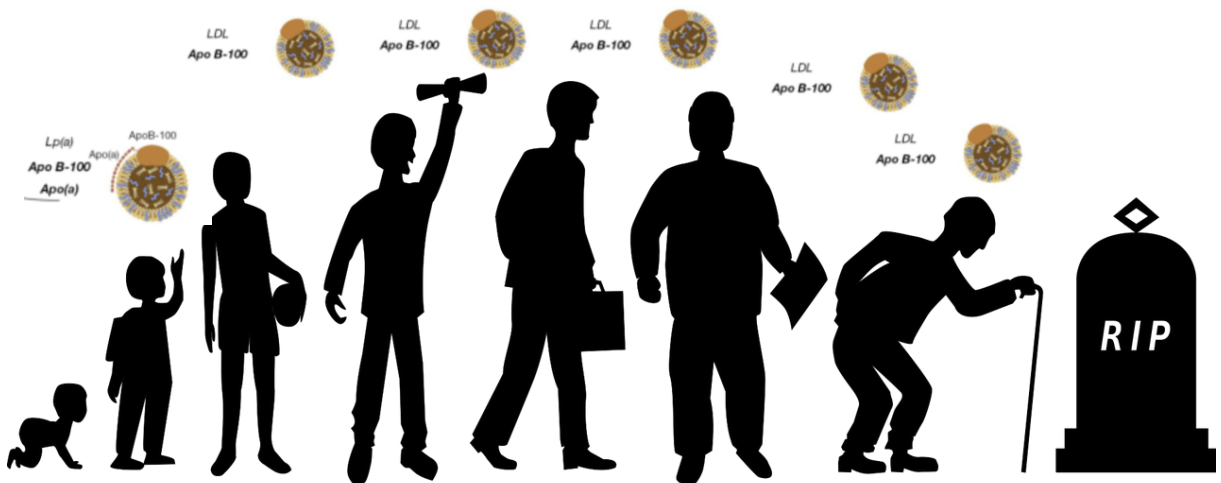


Pack Years
- How much for how long?

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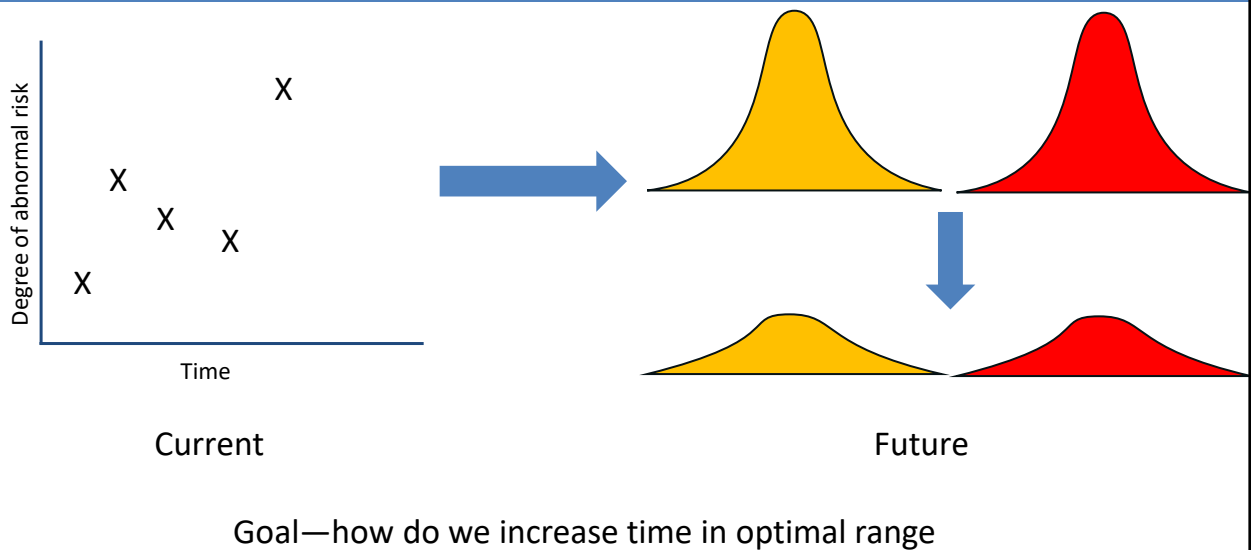
Why Not Cholesterol?



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<https://www.needpix.com/photo/855925/adult-age-baby-child-death-human-life-man-old>
https://commons.wikimedia.org/wiki/File:Remnant_Cholesterol.png

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Why Not Cholesterol?



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Takeaway #1

- Test Earlier
 - Lipid panel at least once for those 9-11
 - Risk personalization labs
 - Lp(a) in all
 - apoB in some
- Treat Earlier
 - Adherence to Life's Essential 8
 - Medications for those 30+ with increased risk
 - Intensify if not at goal

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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ARS Question 1

Which of the Following Is Not True?

- A. Consider obtaining first lipid panel as early as 9 years old
- B. Test Lp(a) in all individuals
- C. Test apoB in all individuals
- D. Consider medications as an adjunct to lifestyle as early as 30 years old



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Checking ApoB

2a: Reasonable to guide plan for those with ASCVD, DM, elevated TGLs on lipid-lowering therapy

2b: for risk personalization and guiding initiating lipid-lowering therapy

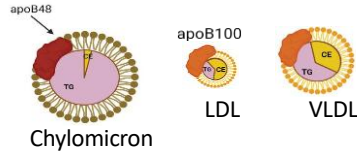
Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Use ApoB to Assess Residual Lipid-Related Risk

ApoB measures all atherogenic particles and is more accurate compared to LDL-cholesterol

Screening with Apolipoprotein B



All atherogenic lipoproteins have a single apoB100 protein. ApoB is not affected by fasting

In adults on lipid-lowering therapy, particularly those with ASCVD, type 2 DM, and/or elevated TG, measurement of apoB is reasonable to guide decisions regarding further therapeutic intensification once LDL-C and/or non-HDL-C goals are achieved (2a)

In adults not on lipid lowering therapy, measurement of apoB may be reasonable to enhance ASCVD risk assessment, guide initiation of lipid lowering therapy, and characterize inherited lipid disorders (2b)

Particularly for secondary prevention, metabolic syndrome, diabetes, elevated triglyceride > 200 mg/dL

Blumenthal RS, et al. J Am Coll Cardiol. 2026;Epub ahead of print.
Blumenthal, R.S., Morris, P.B., et al. 2026 ACC/AHA Guideline on the Management of Dyslipidemia. *Circulation*.

Objective 2

NEXT TOPIC

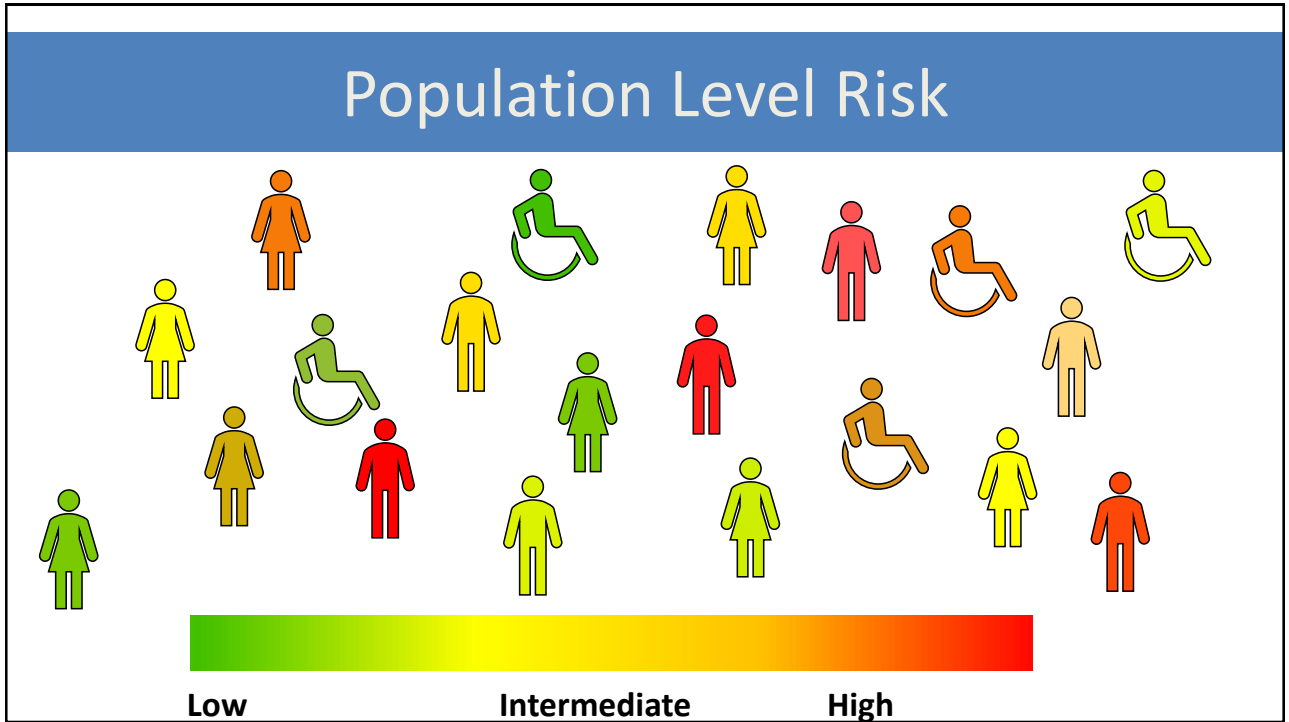
Integrating Subclinical Atherosclerosis into Decision Making

DETECT
Subclinical Atherosclerosis

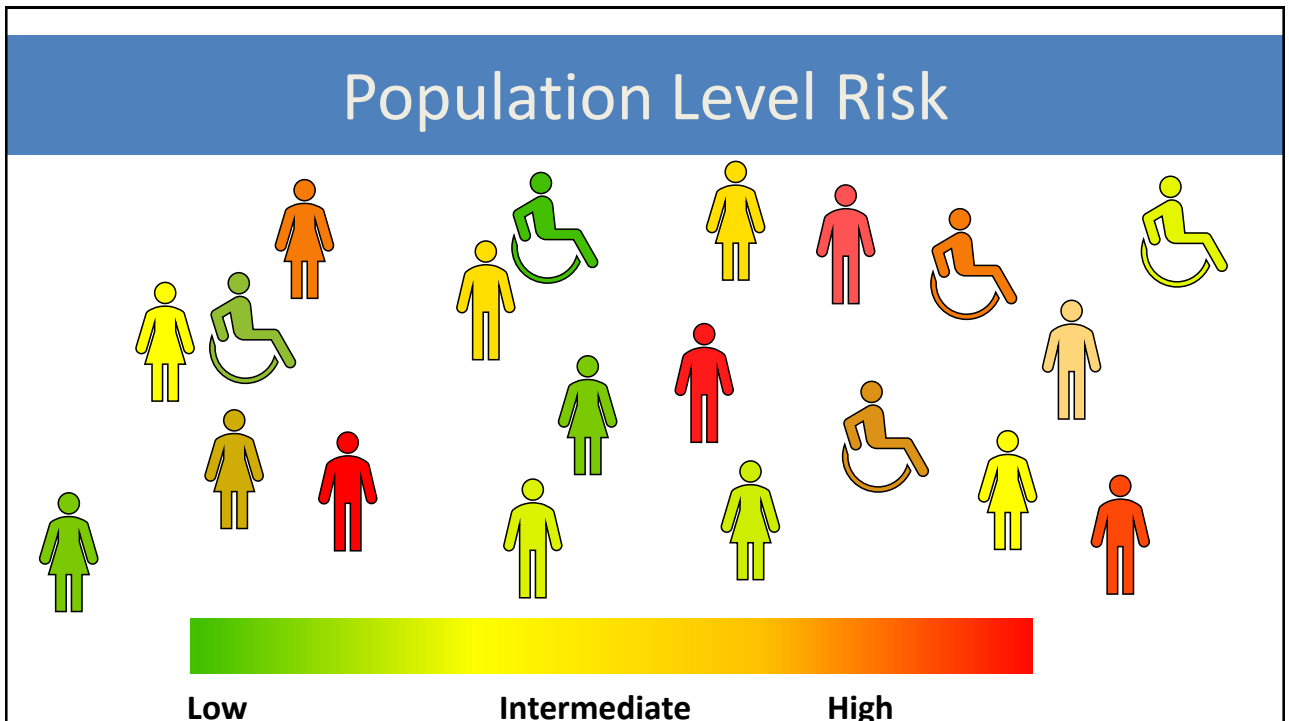
REFINE RISK
Beyond Traditional Risk Factors

INFORM
SHARED DECISION MAKING

PERSONALIZE
PREVENTION STRATEGIES



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2023 PREVENT

For adults 30-79, 10-year and 30-year risk

CVD
ASCVD
Heart Failure

Sex*
 Male Female

Current Smoking
 Any cigarette use within the last 30 days
 No Yes

Lipid-lowering medication
 Current use of statin medication to lower cholesterol
 No Yes

Age (years)*

HDL Cholesterol (mg/dL)*

BMI (kg/m²)*

Total Cholesterol (mg/dL)*

SBP (mmHg)*

eGFR (mL/min/1.73m²)*

Diabetes
 Any history of diabetes.
 No Yes

Anti-hypertensive medication
 Current use of any medication for hypertension
 No Yes

The following three predictors are optional for further personalization of risk assessment. When they are clinically indicated or available,
 If available or indicated, select "Yes" and enter the value.

UACR (mg/g)
 UACR is clinically indicated for individuals with chronic kidney disease, diabetes, or hypertension
 No Yes

HbA1C
 HbA1c is clinically indicated for individuals with diabetes, prediabetes, overweight, or obesity, or those with history of gestational diabetes
 No Yes

Zip Code
 valid 5-digit zip code is needed to estimate social deprivation index [SDI]
 No Yes

Calculate
Reset

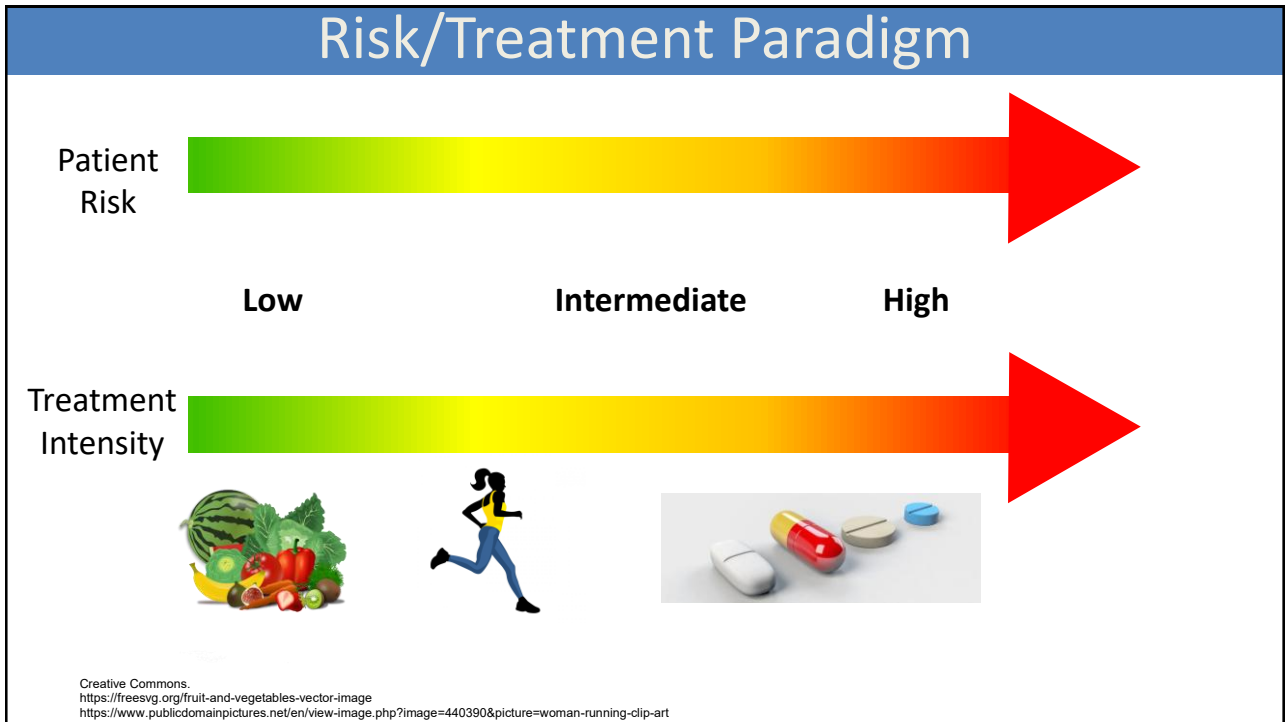
<https://professional.heart.org/en/guidelines-and-statements/prevent-calculator>

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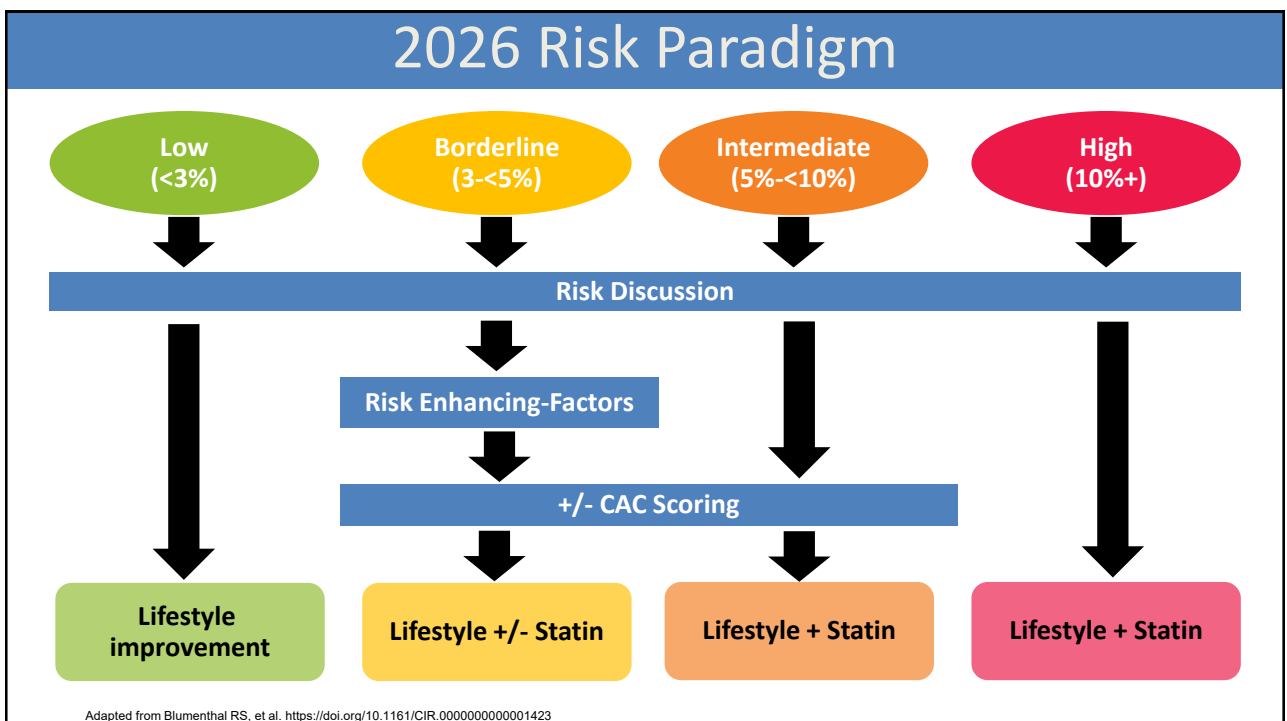
Side-by-Side Comparison

	2013 PCE	2023 PREVENT
Age	X	X
Sex	X	X
Race	X	—
Blood Pressure	X	X
Tobacco Use	X	X
Cholesterol	X	X
Diabetes	X	X
Hypertension Tx	X	X
Lipid Tx	—	X
BMI	—	X
eGFR	—	X
<i>Optional</i>		
UACR	—	X
Hgb A1C	—	X
Zip Code	—	X

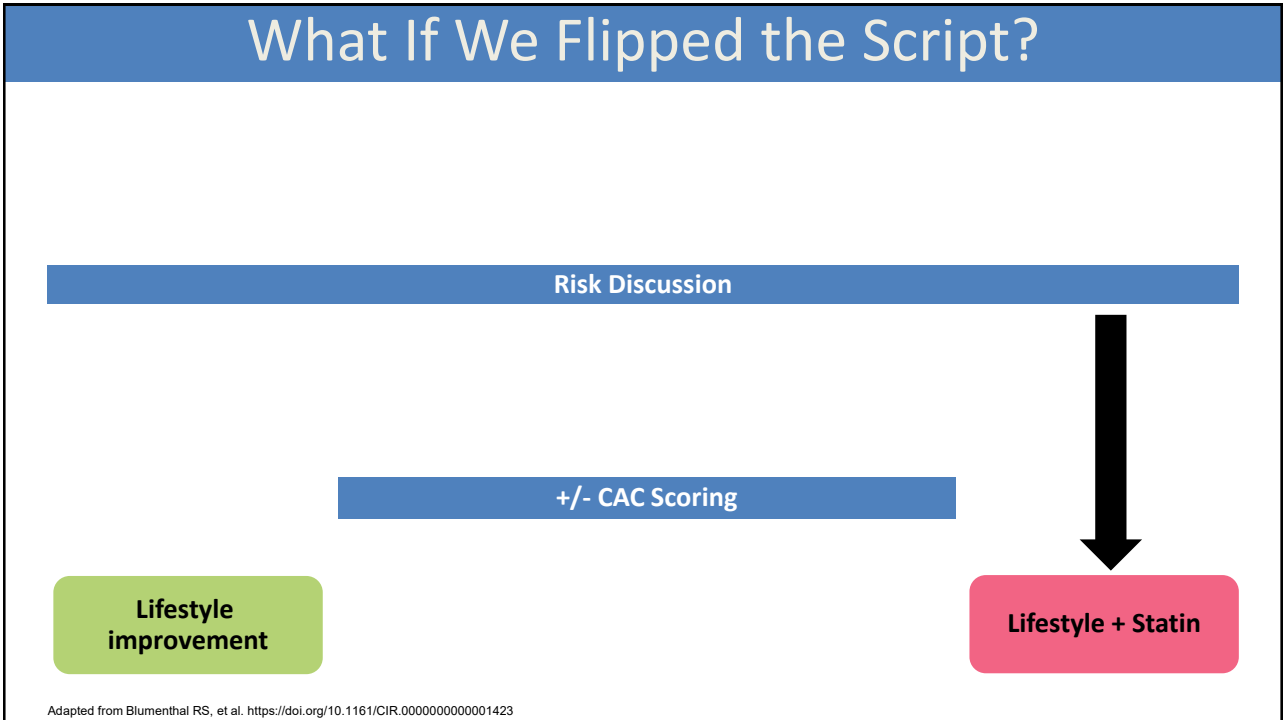
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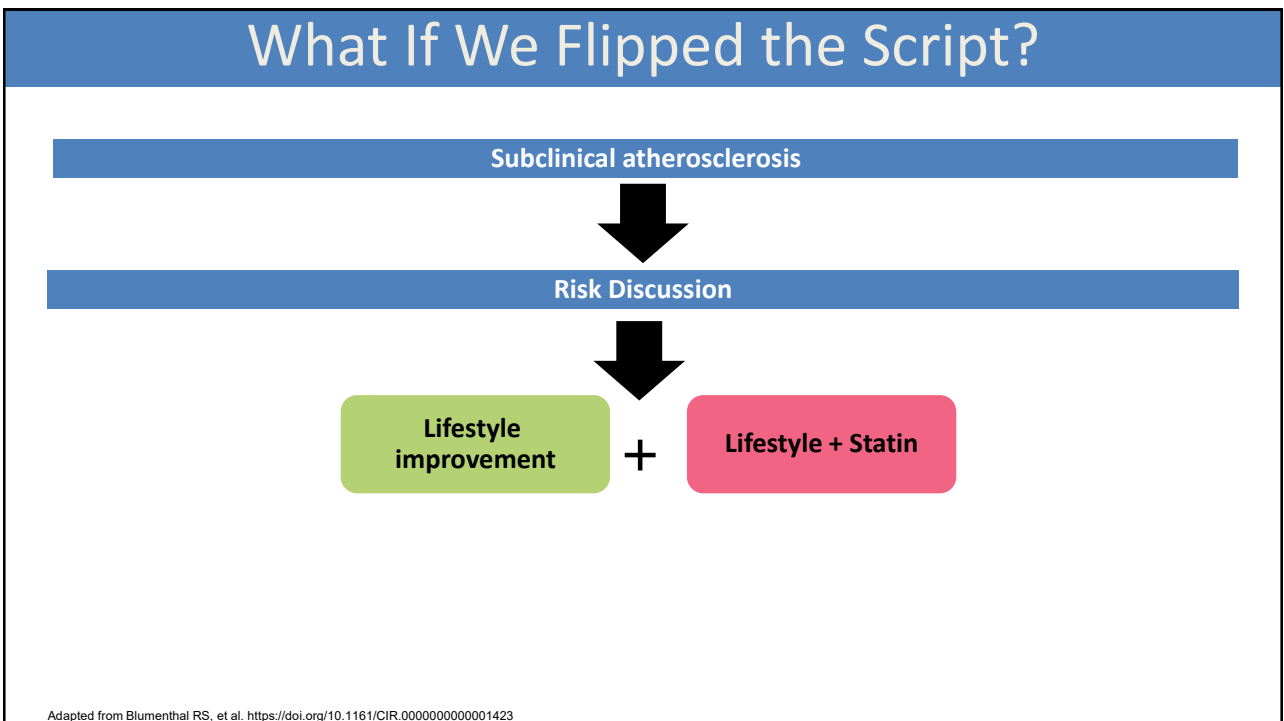
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In Clinic

- First, check for subclinical atherosclerosis
 - Have they had a CAC test done?
 - Have they had a non-gated CT with visual evidence or artificial intelligence evidence of CAC?
 - Other commented atherosclerosis?
- **NB—this does not mean start with CAC testing or order CAC scans on everybody**

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Incidental Atherosclerosis


Comparing two cohorts

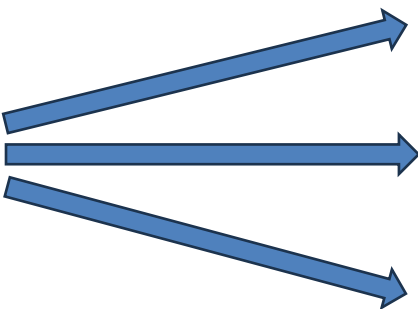
	Stanford iCAC	CAC Consortium
Age	60.5 ± 16.2	54 ± 11
Sex, female	51%	33%
Race/Ethnicity		
White	58%	89%
Black	3.5%	2%
Hispanic	13%	3%


Peng AW, Dudum R, et al. J AM Coll Cardiol. 2023;81(12):1192-1202.
Blaha MJ, et al. J Cardiovasc Comput Tomogr. 2016 Nov 11;11(1):54-61

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Burden of CAC








HOT



COLD



OK

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Adapted from Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Burden of CAC

	CAC Scan	Incidental CAC	LDL-C Goal
	1000+		<55mg/dL
	300+ to 999	Mod-Sev	<70mg/dL <70mg/dL
	100+ to 299	Mild	<70mg/dL <100mg/dL
	1 to 99	Mild	<100mg/dL



Adapted from Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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ARS Question 2

Which One of the Following Subclinical Atherosclerosis Severity/Targets Are Incorrect?

- A. CAC 1000+ and LDL-C goal <55mg/dL
- B. Mild CAC and LDL-C goal <70mg/dL
- C. CAC 100-299 and LDL-C goal <70mg/dL
- D. CAC 300-999 and LDL-C goal <70mg/dL

CAC Score Guided Rx of Subclinical Atherosclerosis

CAC Scoring Recommended if Borderline (3-<5%) or Intermediate (5-<10%) 10y ASCVD Risk In Men ≥40, Women ≥45 y.o

May defer LLT if not high risk: no DM, FH, tobacco use, family history of premature ASCVD, or LDL-C ≥190 mg/dL. Repeat CAC in 3-7 yr.

Incidental CAC on a nongated CT scan can also be used to guide therapy.

- If mild, LDL-C goal <100 mg/dL.
- If moderate or severe, LDL-C goal is <70 mg/dL.

Absence of incidental CAC should not be used to defer Rx given the low negative predictive value of nongated scans.

J Am Coll Cardiol. 2026;XX:XXX-XXX; *Circulation.* 2026;153:e00-e00
 Blumenthal, R.S., Morris, P.B., et al. 2026 ACC/AHA Guideline on the Management of Dyslipidemia. *Circulation.*

Takeaway #2

- Any CAC favors lipid lowering therapy in addition to lifestyle modifications
- If there is no evidence of subclinical atherosclerosis, then proceed with PREVENT-based risk assessment

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Higher Risk Conditions

HIV

Adverse Pregnancy Outcomes

- Gestational Hypertension, Pre-Eclampsia, Eclampsia

Chronic inflammatory disorders

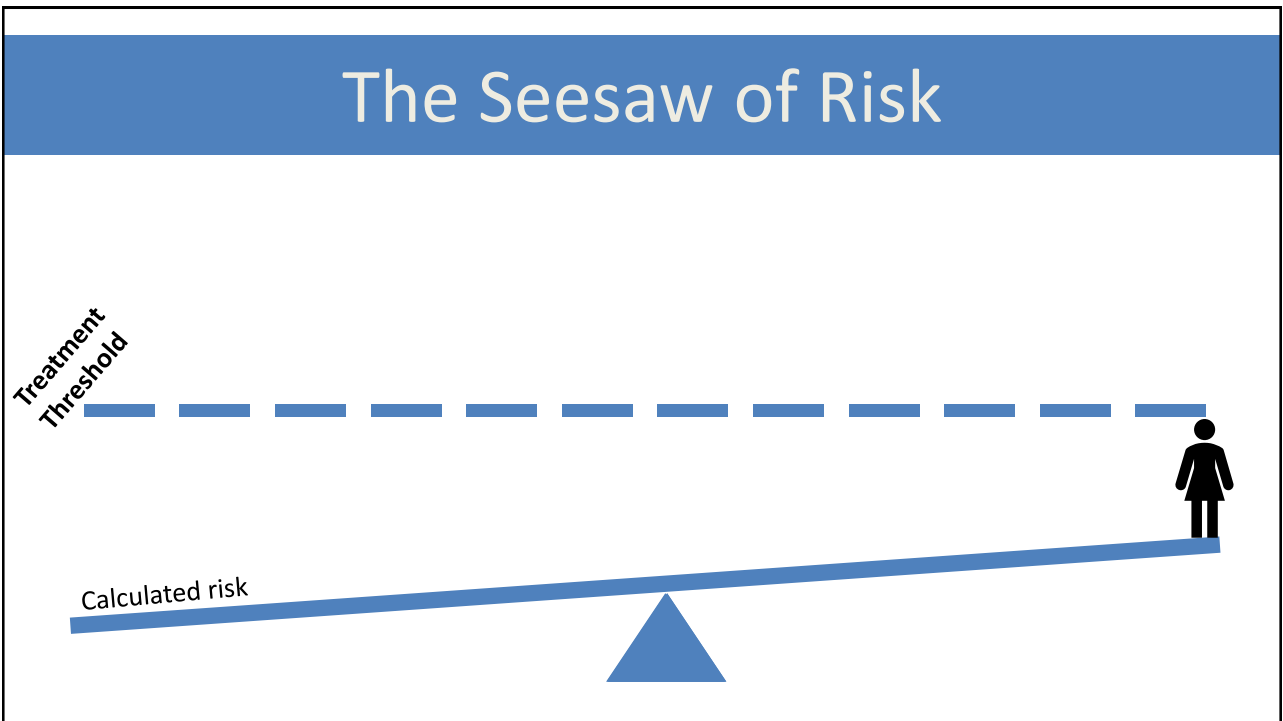
- RA, Lupus, Inflammatory Arthritis, etc.

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

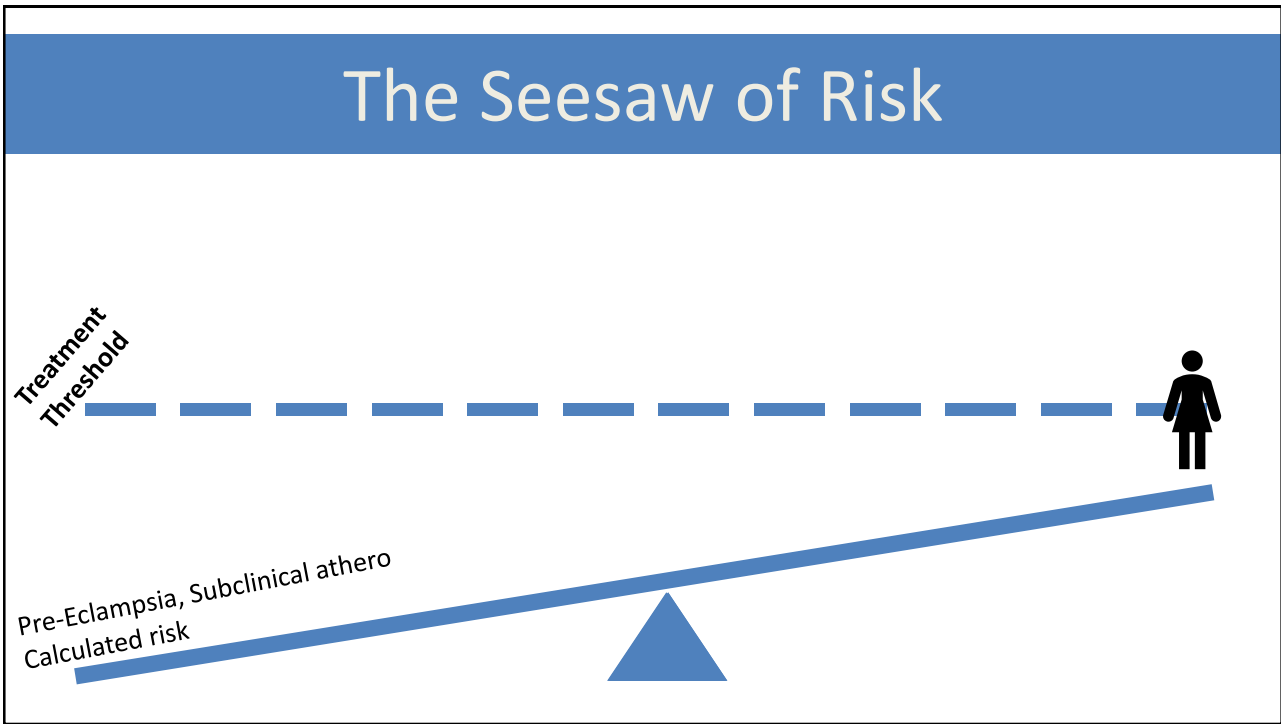
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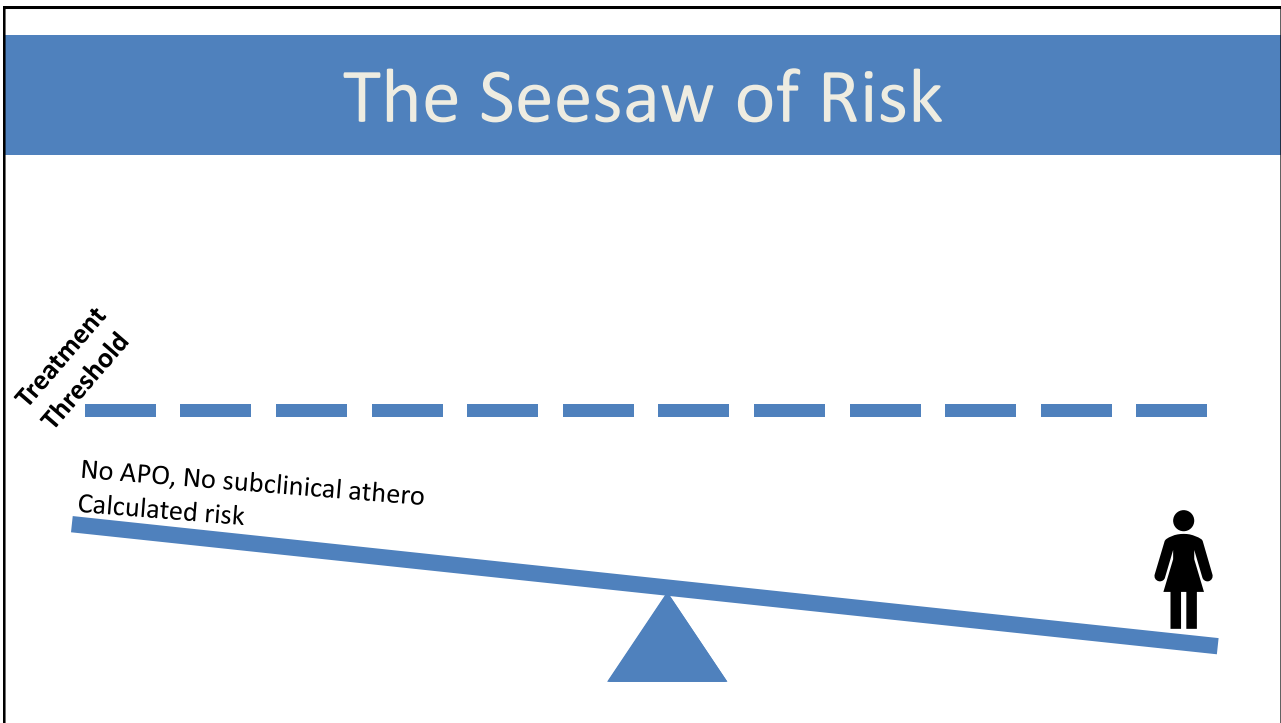
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Hypertriglyceridemia

- Risk-enhancing factor
 - >150mg/dL, if fasting
 - >175mg/dL, if non-fasting
- Consider lipid referral if >400mg/dL

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Hypertriglyceridemia Management

- Diet/Lifestyle in all
 - Increasing physical activity
 - Reduce the following:
 - Added Sugar
 - Total Fat
 - Alcohol
- Weight loss, if needed

Class 1: Potential addition of statin therapy (TGL \geq 150mg/dL)

Class 2a: Prescription strength omega-3s and fibrates (TGL \geq 500mg/dL)

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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When to Refer to a Specialist?

- Concern for heritable cholesterol disorders
- Extremes of risk, i.e. very early MI or very abnormal risk factors
- Significantly elevated triglycerides
- Concerns about drug interactions
 - Medical comorbidities
 - Considering pregnancy, pregnant, postpartum

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Takeaway #3

- Ask about associated conditions that increase ASCVD risk
- Incorporate into shared decision-making conversation
- Consider lipid specialist referral to assist in co-management

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Objective 3

A Step-Wise Approach

Using Medications to Hit Treatment Goals

GOAL ACHIEVED
Sustained risk reduction for better outcomes
refine as needed

5 REASSESS
Monitor response, tolerability, and adherence; refine as needed

4 TITRATE
Adjust dose and/or add therapy to achieve goals

3 START
Initiate appropriate medication therapy

2 SET GOALS
Define patient-centered lipid and risk reduction goals

1 ASSESS
Evaluate risk, comorbidities, and baseline lipids

ASSESS
Evaluate risk, comorbidities, and baseline lipids

SET GOALS
Define patient-centered lipid and risk reduction goals

START
Initiate appropriate medication therapy

TITRATE
Adjust dose and/or add therapy to achieve goals

REASSESS
Monitor response, tolerability, and adherence; refine as needed

Treatment Goals

- ✓ LDL-C reduction
- ✓ Apolipoprotein B reduction
- ✓ Non-HDL-C reduction
- ✓ Overall ASCVD risk reduction

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Risk/Treatment Paradigm

Patient Risk

Treatment Intensity

Low **Intermediate** **High**

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<https://www.publicdomainpictures.net/en/view-image.php?image=440390&picture=woman-running-clip-art>

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Types of Lipid Lowering Therapy

- Statins
 - Ezetimibe
 - Bempedoic acid
 - PCSK9-based therapies
- Non-statins
 - Ezetimibe
 - Bempedoic acid
 - PCSK9-based therapies

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Estimated LDL-C Reduction

LDL-C Reduction

Rosuvastatin 5-40mg

Atorvastatin 10-80mg

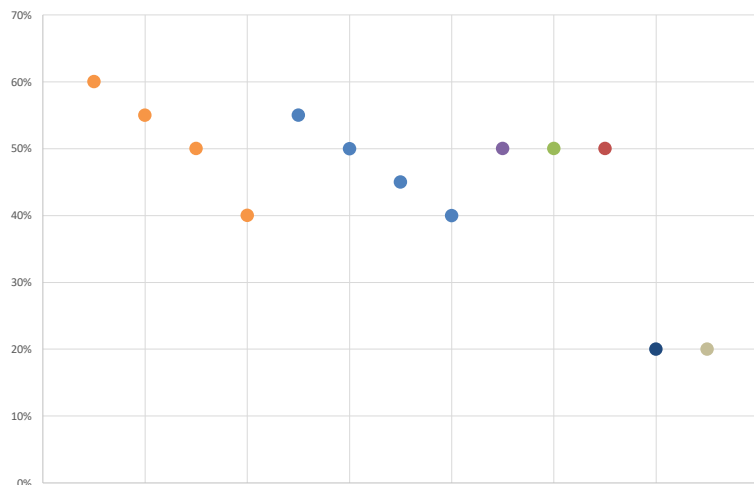
Evolocumab

Alirocumab

Inclisiran

Ezetimibe

Bempedoic Acid



Adapted from Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>
 Adapted from <https://www.uptodate.com/contents/search?search=statin%20comparison%20chart>

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Estimated LDL-C Reduction

- High-intensity statins (50+%)
 - Rosuvastatin 20mg or 40mg
 - Atorvastatin 40mg or 80mg
- Moderate-intensity statins (30%-50%)
 - Most other statin doses/combinations
- NB—not covering low-intensity statins

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Managing Statin Intolerance

- Good history
 - What happened? What dose? Time course?
 - Concurrent medications?
- Other statins tried?
- Was any work-up done at the time?
- Shared decision-making re: personal risks/benefits to determine if re-trialing is indicated

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Managing Statin Intolerance

- If only lipophilic statins have been tried, try hydrophilic statins
 - Rosuvastatin or pravastatin
- Start with lowest dose and ramp up
- Goal: Highest tolerated dose

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What About Supplements?

- Commonly used supplements, include red yeast rice, berberine, plant sterols, fish oil, etc.
- Head-to-head trial of moderate-intensity rosuvastatin and common supplements found that none were as efficacious as rosuvastatin 5mg

Class 3: No benefit for using dietary supplements for treating lipids

Laffin LJ et al. <https://doi.org/10.1016/j.jacc.2022.10.013>
Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Takeaway #4

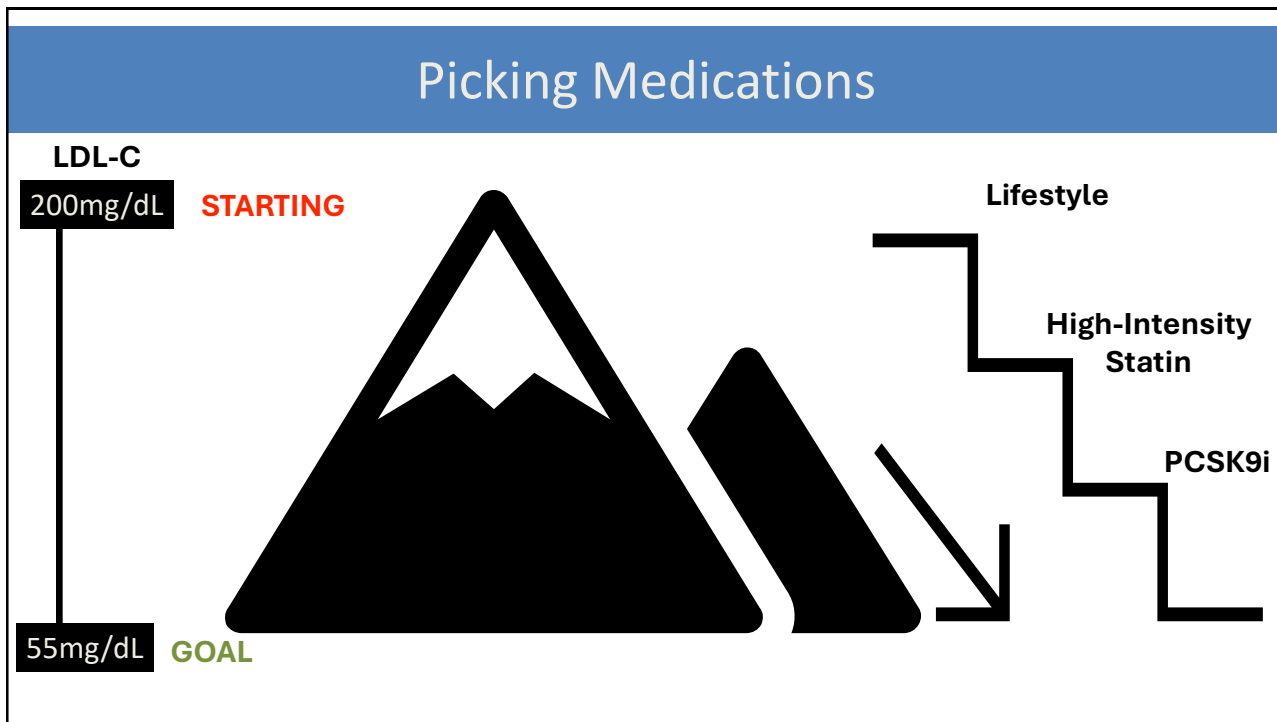
- Concern for statin related adverse events are common
- Obtain a good history and discuss with patients to determine if and when to re-challenge
- If re-challenging, consider low-dose and a ramp up

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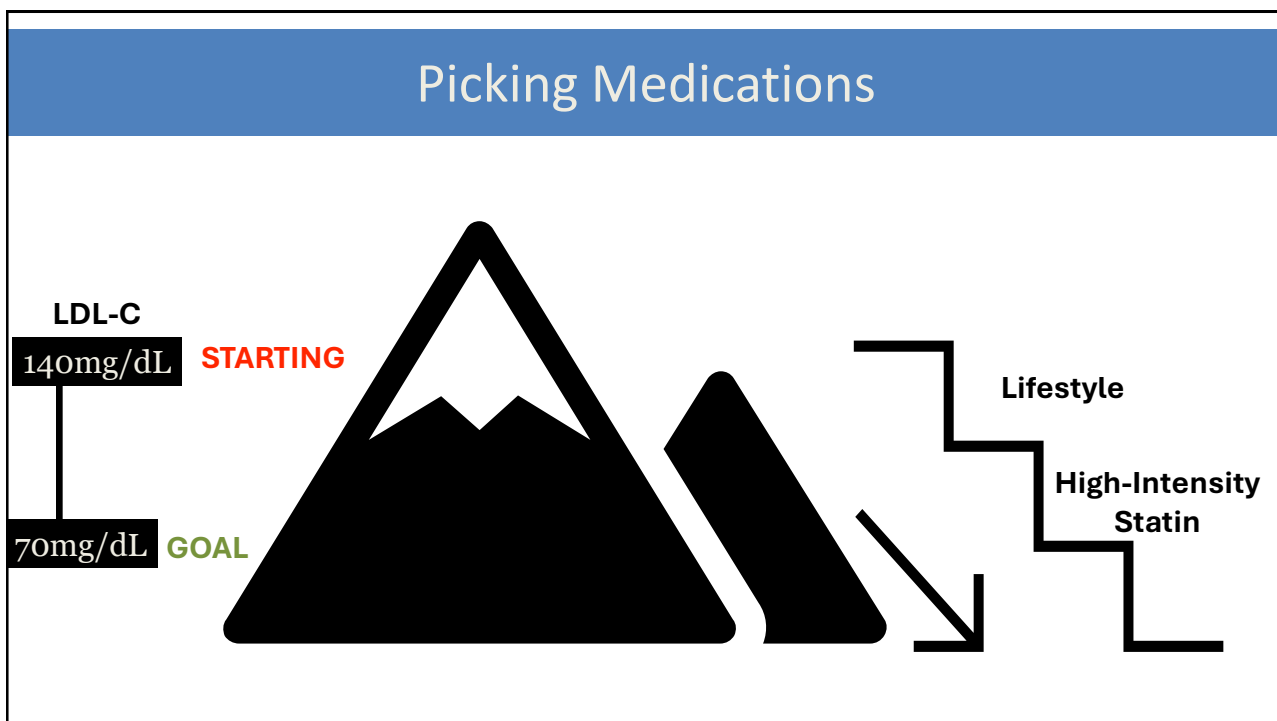
Picking Medications



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When to Check Labs?

- At baseline
- If lifestyle changes only, after 3-6 months
- >4 weeks after lipid lowering therapy change
- Include apoB when metabolic milieu present to assess for apoB/LDL-C discordance

Blumenthal RS, et al. <https://doi.org/10.1161/CIR.0000000000001423>

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Takeaway #5

- Think of where you're starting and where you need to go for lipid lowering
- Use a combination of lifestyle and medication(s) to get there
- If discordance, use apoB goal

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ARS Question 3**Which of These Combinations Is Unlikely to Achieve More Than a 50% LDL-C Reduction?**

- A. Alirocumab 150mg every other week
- B. Atorvastatin 20mg + Ezetimibe 10mg
- C. Bempedoic Acid 180mg daily
- D. Rosuvastatin 40mg daily

Summary

- Personalize risk and think of the cumulative exposure of risk factors as well as incremental risk increase
 - Higher Risk Conditions
- Subclinical atherosclerosis, if present, can help simplify risk assessment and recommendations
- Lifestyle therapy and medications, as needed
 - Remember → Where are you starting, Where are you going?