#### SMALL INTERACTIVE SESSION

DIABETES AND CVD IN PRIMARY CARE:
LASSO-UP THE LATEST GUIDELINES FOR BEST
OUTCOMES!
A PATIENT CASE APPROACH TO INTEGRATING THE
ACC/AHA AND ADA/AACE RECOMMENDATIONS

AAPA Conference May 19, 2024 at 3:00PM to 5:00PM Houston, TX 2024

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DEPARTMENT OF MEDICINE | SCHOOL OF HEALTH PROFESSIONS | PHYSICIAN ASSISTANT PROGRAM





I have no relevant relationships with ineligible companies to disclose within the past 24 months.

### **OBJECTIVES**

## At the conclusion of this session, participants should be able to:

- 1. Identify patients with, or at high-risk of, diabetes and cardiovascular disease with screening methodologies from the USPSTF and other authorities
- Interpret current diagnostic guidelines for cardiovascular disease from ACC/AHA and diabetes from ADA and the AACE
- 3. Develop individualized treatment plans using shared-decision approaches rooted in evidence-based pharmacotherapeutics including the latest agents such as SGLT2-inhibitors, GLP1-receptor agonists, and PCSK9 inhibitors
- 4. Counsel and motivate diverse patient populations with culturally informed lifestyle modifications while considering social determinants of health

### Acronyms

- USPSTF United States Preventive Services Task Force
- ADA American Diabetes Association
- AACE American Association of Clinical Endocrinologists
- ACE American College of Endocrinologists
- ACC American College of Cardiologists
- ACP American College of Physicians
- AHA American Heart Association
- **GDM** Gestational Diabetes Mellitus
- **CKD** Chronic kidney disease
- ASCVD Atherosclerotic Cardiovascular Disease





#### HYPERTENSION SCREENING: USPSTF

Figure. Clinician Summary: Screening for Hypertension in Adults	Figure, Clinician Summar	v: Screening for Hypertension in Adults
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What does the USPSTF recommend?	Screen adults for hypertension. Grade: A
To whom does this recommendation apply?	Adults 18 years or older without known hypertension.
How to implement this	1. Screen: Measure blood pressure with an office blood pressure measurement.
recommendation?	Confirm: Take blood pressure measurements outside of the clinical setting to confirm a hypertension diagnosis before starting treatment.
	<ul> <li>Ways to measure blood pressure outside of the clinical setting include</li> <li>Ambulatory blood pressure monitoring: patients wear a programmed portable device that automatically takes blood pressure measurements, typically in 20- to 30-minute intervals over 12 to 24 hours while patients go about their normal activities or are sleeping.</li> </ul>
	<ul> <li>Home blood pressure monitoring: patients measure their own blood pressure at home with an automated device.</li> <li>Measurements are taken much less frequently than with ambulatory blood pressure monitoring (eg, 1 to 2 times a day or week, although they can be spread out over more time).</li> </ul>
	Blood pressure measurements should be taken at the brachial artery (upper arm) with a validated and accurate device in a seated position after 5 minutes of rest.
How often?	Although evidence on optimal screening intervals is limited, reasonable options include  • Screening for hypertension every year in adults 40 years or older and in adults at increased risk for hypertension (such as Black persons, persons with high-normal blood pressure, or persons who are overweight or obese).
	Screening less frequently (ie, every 3-5 years) as appropriate for adults aged 18 to 39 years not at increased risk for hypertension and with a prior normal blood pressure reading.

#### HYPERTENSION **SCREENING**: Categorize BP Readings

### **Blood Pressure Categories**



BLOOD PRESSURE CATEGORY	SYSTOLIC mm Hg (upper number)		DIASTOLIC mm Hg (lower number)
NORMAL	LESS THAN 120	and	LESS THAN 80
ELEVATED	120 – 129	and	LESS THAN 80
HIGH BLOOD PRESSURE (HYPERTENSION) STAGE 1	130 – 139	or	80 – 89
HIGH BLOOD PRESSURE (HYPERTENSION) STAGE 2	140 OR HIGHER	or	90 OR HIGHER
HYPERTENSIVE CRISIS (consult your doctor immediately)	HIGHER THAN 180	and/or	HIGHER THAN 120

### How to measure your blood pressure at home

Follow these steps for an accurate blood pressure reading



Avoid caffeine, cigarettes and other stimulants 30 minutes before you measure your blood pressure.

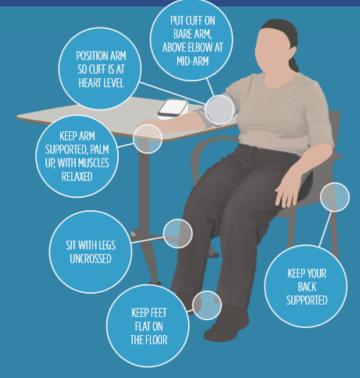
Wait at least 30 minutes after a meal.

If you're on blood pressure medication, measure your BP **before** you take your medication.

Empty your bladder beforehand.

Find a quiet space where you can sit comfortably without distraction.











Rest for five minutes while in position before starting.

Take two or three measurements, one minute apart.

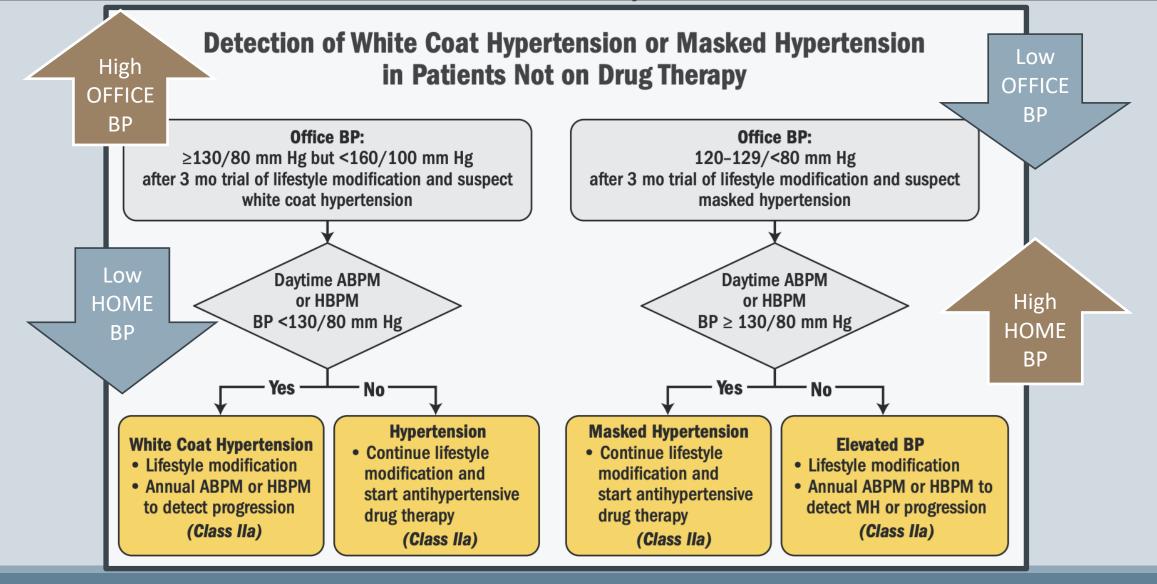
Keep your body relaxed and in position during measurements.

Sit quietly with no distractions during measurements—avoid conversations, TV, phones and other devices.

Record your measurements when finished.

This Prepare, position, measure handout was adapted with permission of the American Medical Association and The Johns Hopkins University. The original copyrighted content can be found at https://www.ama-assn.org/ama-johns-hopkins-blood-pressure-resources.

#### HYPERTENSION DIAGNOSIS: Verify White Coat vs Masked HTN







Mejorar y mantener la salud cardiovascular, o CVH (por sus siglas en inglés), puede ayudarlo a disfrutar de una vida más larga y saludable. Una mejor CVH también se asocia con la disminución del riesgo de enfermedades cardíacas, accidentes cerebrovasculares, cáncer, demencia y otros problemas de salud importantes.

> Life's Essential 8 describe algunos sencillos pasos que puede seguir para llevar un estilo de vida más saludable.



#### **✓** MEJORE SU ALIMENTACIÓN

Busque un patrón de alimentación saludable que incluya alimentos integrales, muchas frutas y verduras, proteínas magras, frutos secos, semillas y cocinar con aceites no tropicales, como el de oliva y el de canola.







#### SEA MÁS ACTIVO

Los adultos deben realizar 150 minutos de actividad física moderada o 75 minutos de actividad física intensa. Caminar es bueno para niveles de actividad moderada. Los niños deben realizar 60 minutos todos los días, incluidos juegos y actividades estructuradas.





#### **✓** DEJE EL TABACO

El uso de productos de administración de nicotina inhalada, que incluye cigarrillos tradicionales, cigarrillos electrónicos y vapeo, es la causa principal de muerte prevenible en los EE. UU., incluye aproximadamente un tercio de todas las muertes por enfermedades cardíacas. Además, aproximadamente un tercio de los niños de los EE. UU. entre los 3 y los 11 años se encuentran expuestos al humo y vapor indirectos.



#### **DUERMA LO SUFICIENTE**

Dormir bien todas las noches es vital para la salud cardiovascular. Los adultos deben tener como objetivo un promedio de 7 a 9 horas, u los bebés u niños necesitan más según su edad. Los estudios indican que el sueño excesivo o insuficiente está asociado con enfermedades cardíacas.



adultos varía entre 18.5 u menos de 25.

**✓** CONTROLE EL COLESTEROL

Los niveles altos de colesterol no transportado

por las lipoproteínas de alta densidad (HDL). o "malo", pueden provocar enfermedades

cardíacas. Su profesional de la salud puede

número preferido que monitorear, en lugar

del colesterol total, ya que se puede medir

considerar el colesterol no HDL como el

sin ayuno de antemano y se calcula de manera confiable en todas las personas.

Puede calcularlo en línea o consultar con un profesional de la salud.

La mayoría de los alimentos que comemos se convierten en glucosa (o glucemia) que nuestro cuerpo utiliza como energía. Con el tiempo, los niveles altos de glucemia pueden dañar el corazón, los riñones, los ojos y los nervios. Como parte de las pruebas, el monitoreo de la hemoglobina A1c puede reflejar mejor el control a largo plazo en personas con diabetes o prediabetes.



#### CONTROLE LA PRESIÓN ARTERIAL

Mantener la presión arterial dentro de márgenes aceptables puede mantenerlo saludable durante más tiempo. Los niveles inferiores a 120/80 mm Hg son los ideales. La presión arterial alta se define como una presión sistólica de 130-139 mm Hg (el número más alto en la lectura) o una presión diastólica de 80-89 mm Hg (el número inferior).



Obtenga más información en heart.org/lifes8





## AHA Life's Essential

## Lifestyle Modifications



#### What Can I Do To Improve My Blood Pressure?

Approximate SBP Modification Recommendation **Reduction Range** Maintain normal body weight Weight reduction 5 mm Hg  $(BMI=18.5-24.9 \text{ kg/m}^2)$ Diet rich in fruits, vegetables, low **DASH** eating plan 11 mm Hg fat dairy and reduced in fat Restrict sodium <1500 mg of sodium per day 5-6 mm Hg intake Be more physically active. Aim for at least 90 to 150 minutes **Physical activity** 5-8 mm Hg of aerobic exercise per week.



Moderation of alcohol consumption

No more than 2 drinks/day for men and 1 drink/day for women

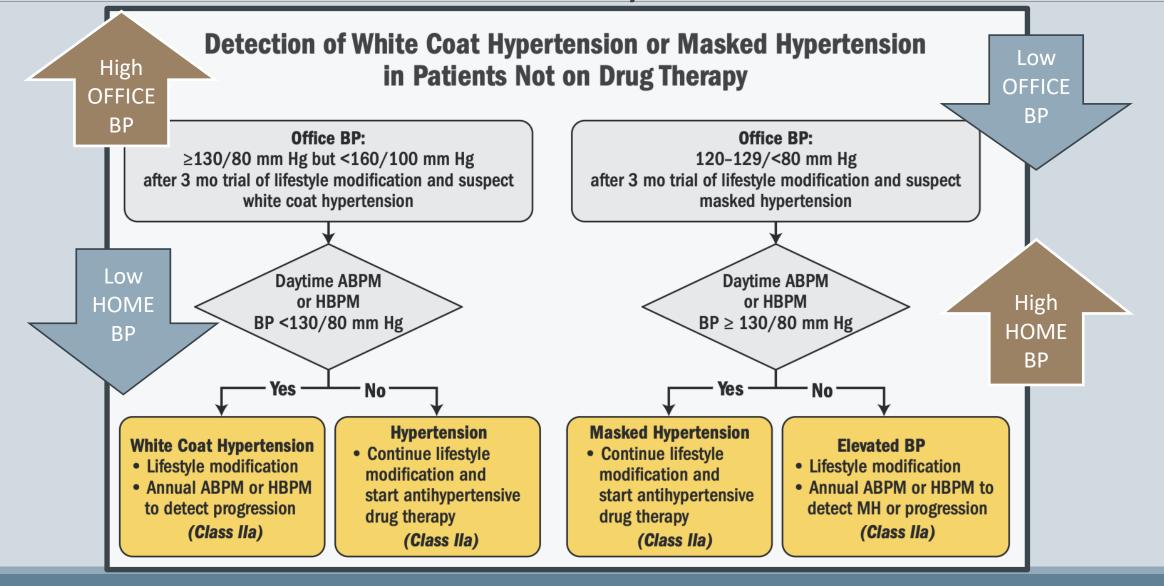
4 mm Hg

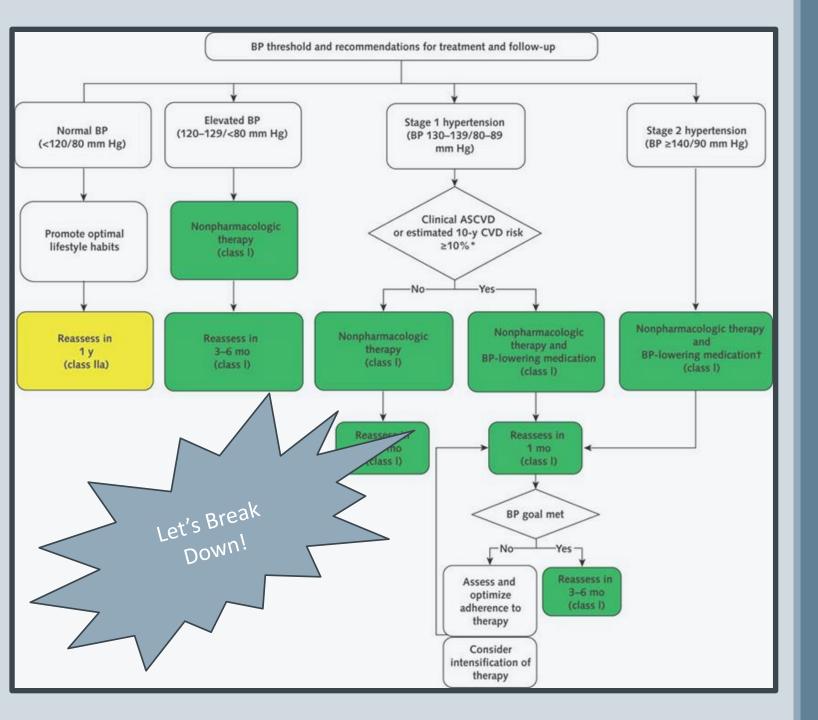


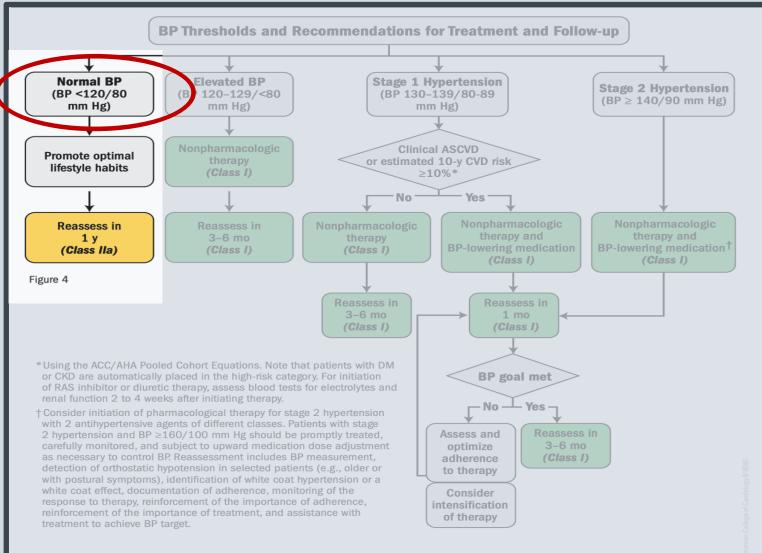
BP = Blood pressure, BMI = Body mass index, SBP = Systolic blood pressure, DASH = Dietary Approaches to Stop Hypertension



#### HYPERTENSION **DIAGNOSIS**: Verify White Coat vs Masked HTN



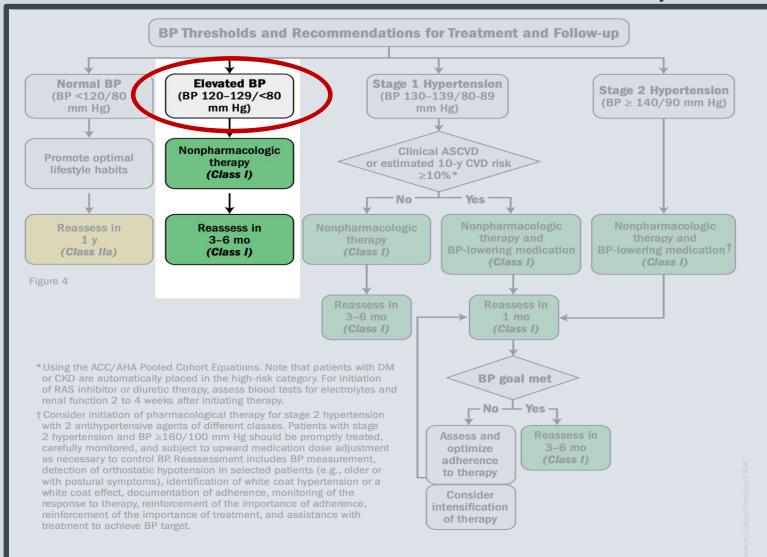




# Normal BP: <120/<80 mmHg

# Treatment: Promote Healthy Lifestyle Habits





# Elevated BP: 120-129/<80 mmHg

# Treatment: Lifestyle Modifications





#### What Can I Do To Improve My Blood Pressure?

Approximate SBP Modification Recommendation **Reduction Range** Maintain normal body weight **Weight reduction** 5 mm Hg  $(BMI=18.5-24.9 \text{ kg/m}^2)$ Diet rich in fruits, vegetables, low **DASH** eating plan 11 mm Hg fat dairy and reduced in fat Restrict sodium <1500 mg of sodium per day 5-6 mm Hg intake Be more physically active. Aim for at least 90 to 150 minutes **Physical activity** 5-8 mm Hg of aerobic exercise per week.



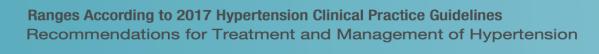
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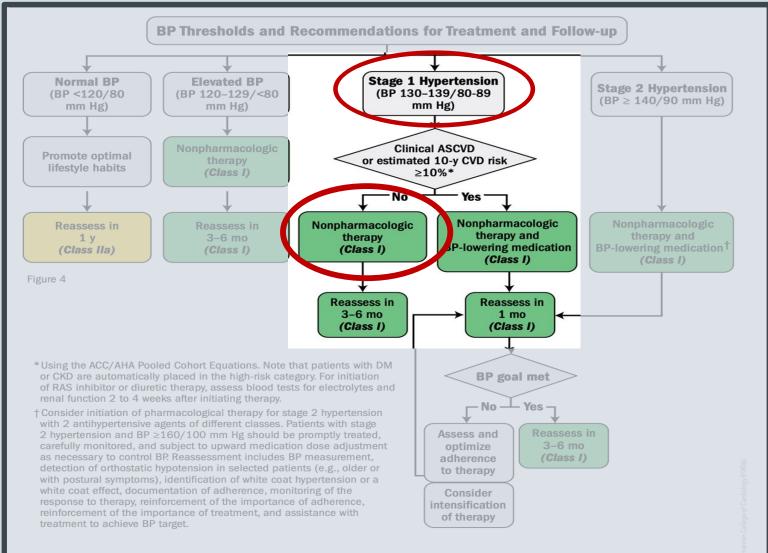
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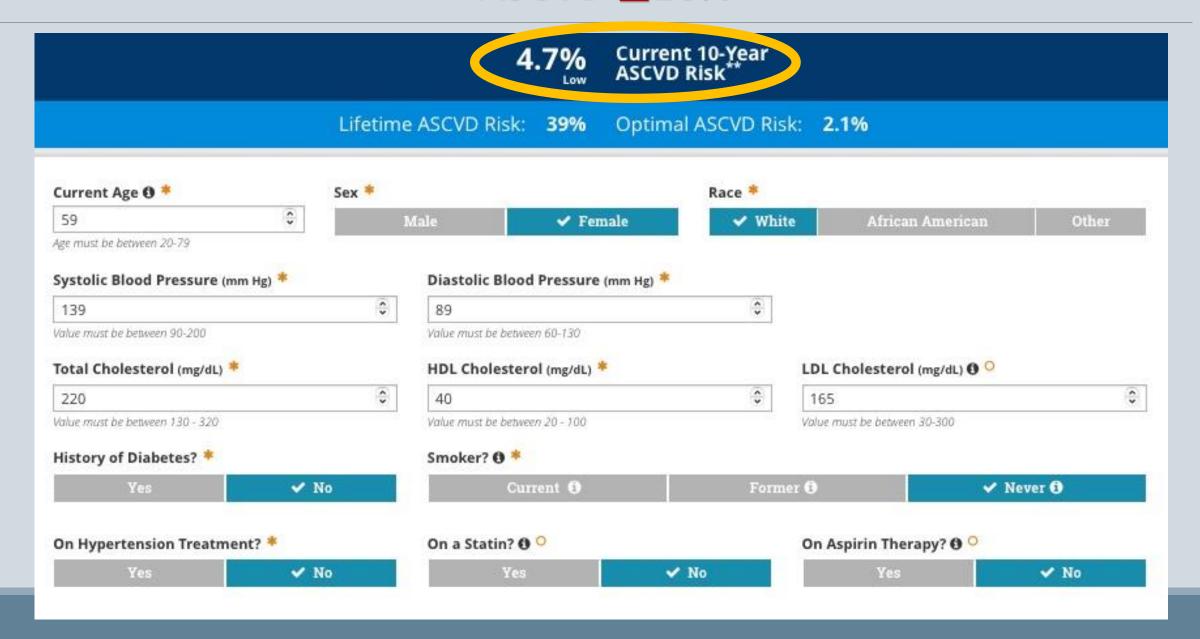


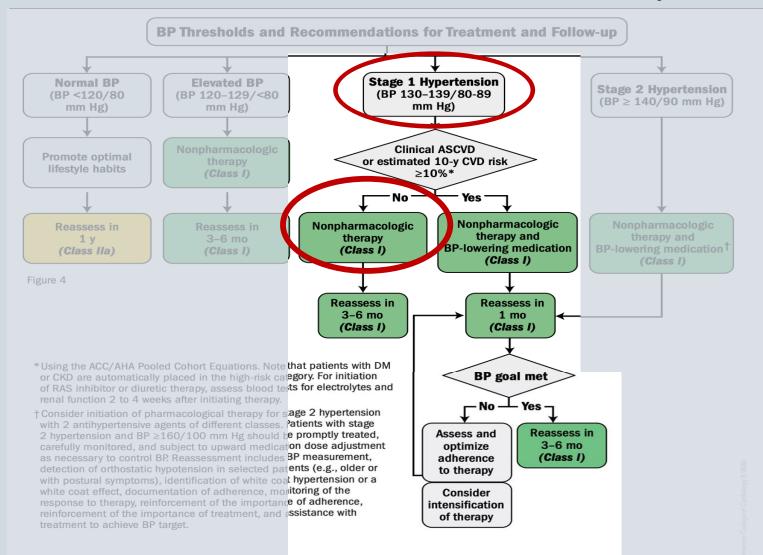


STAGE 1: 130-139/80-89 mm Hg and ASCVD ≤ 10%



#### ASCVD RISK ESTIMATOR ASCVD ≤10%





#### **STAGE 1:**

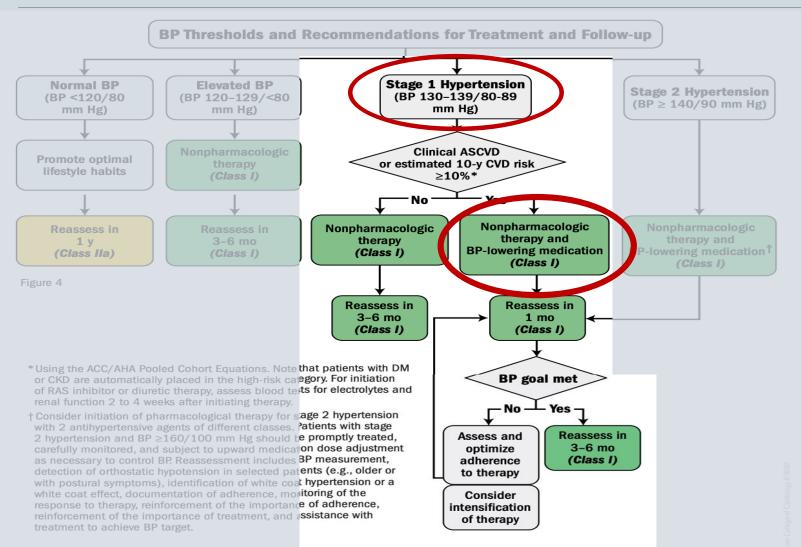
130-139/80-89 and

ASCVD ≤10%

#### **TREATMENT:**

Lifestyle
Modifications &
Recheck 3-6 mo



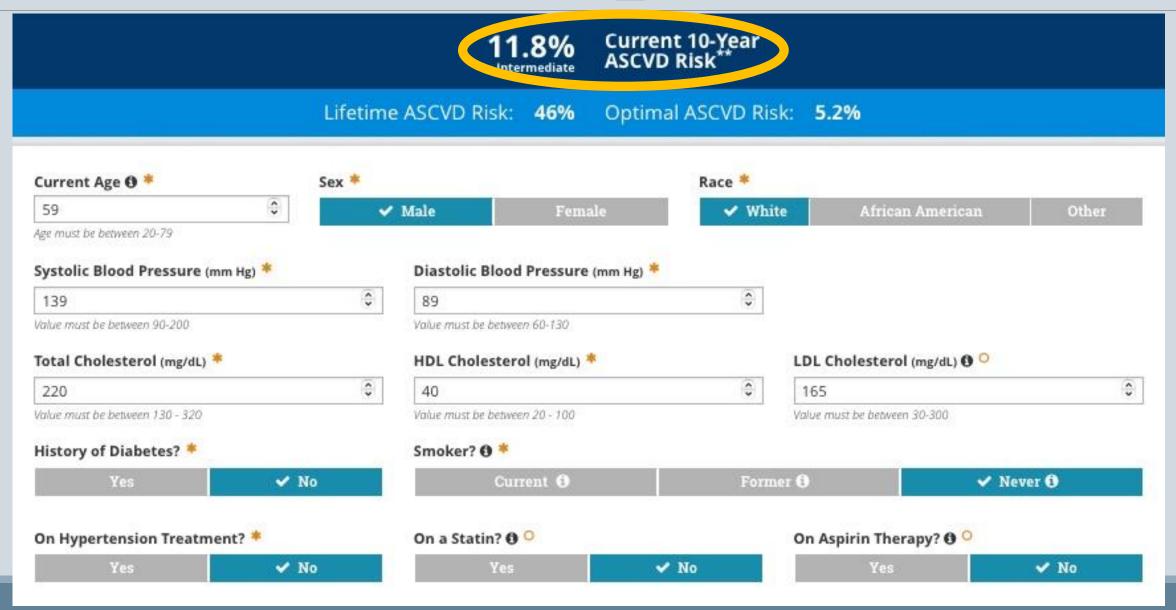


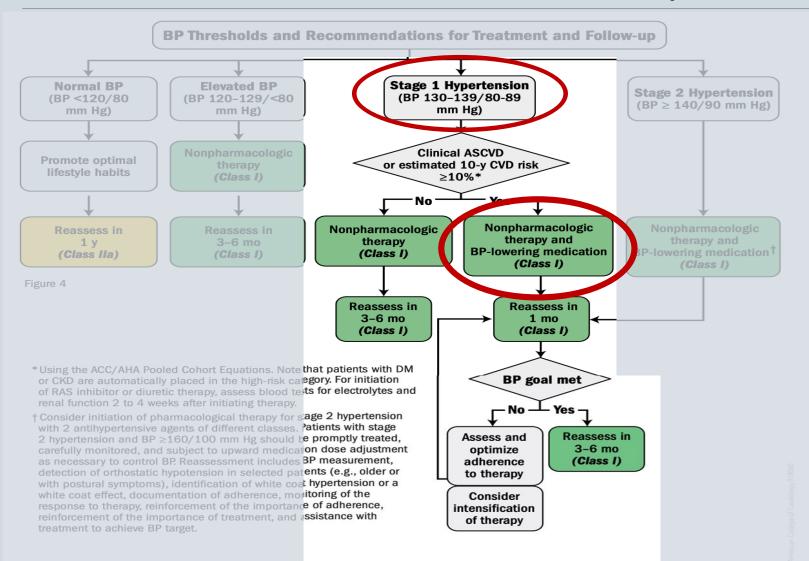
STAGE 1: 130-139/80-89 mm Hg and ASCVD ≥ 10%

ASCVD 2 10%



#### ASCVD RISK ESTIMATOR ASCVD ≥ 10%





#### STAGE 1:

130-139/80-89 and

ASCVD ≤10%

# TREATMENT: Lifestyle Modifications & Start Therapy



#### **ASCVD** ≥ 10% **STAGE** 1 HYPERTENSION TREATMENT

#### RENIN-ANGIOTENSIN-ALDOSTERONE (RAA) BLOCKING AGENTS

- ✓ Angiotensin-Converting Enzyme Inhibitor (-pril e.g., lisinopril)
- ✓ Angiotensin Receptor Blockers (-<u>sartan</u> e.g., olmesartan)

#### CALCIUM CHANNEL BLOCKERS

✓ Dihydropyridine (-pine e.g., amlodipine)

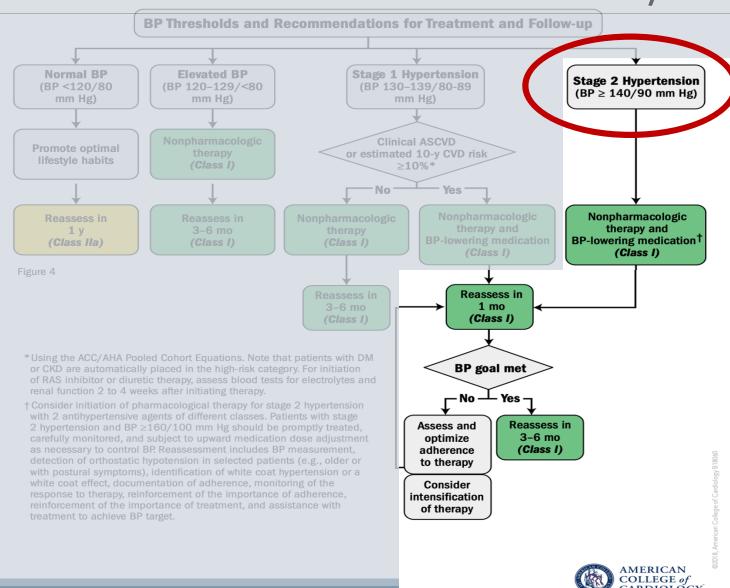
#### **DIURETICS**

✓ Thiazides (-ide e.g., hydrochlorothiazide)

#### What About Beta-Blockers?

#### **BETA-BLOCKERS**

- ✓ Not recommended as first-line
- ✓ MI or Ischemic Heart Disease (IHD): Considered first-line in this population
- ✓ HFrEF: Bisoprolol, metoprolol succinate & carvedilol preferred
- ✓ Do NOT combine with CCB-nondihydropyridines due to increased risk of bradycardia and heart block



STAGE 2: ≥140/90<sub>mmHg</sub>

TREATMENT:
Lifestyle
Modifications &
Optimize Meds



## STAGE 2 HYPERTENSION TREATMENT

ACC recommends starting **TWO** first-line agents of different classes:

- √ Stage 2 Hypertension
- ✓ BP more than 20/10 mm hg above their BP target



YOU STARTED TREATMENT, WHAT'S NEXT?

RECHECK
IN
1 MONTH





If NOT AT GOAL after ONE month:

✓ Confirm adherence to therapy

✓ Consider intensifying therapy

#### **STAGE 2 HYPERTENSION TREATMENT**

#### RENIN-ANGIOTENSIN-ALDOSTERONE (RAA) BLOCKING AGENTS

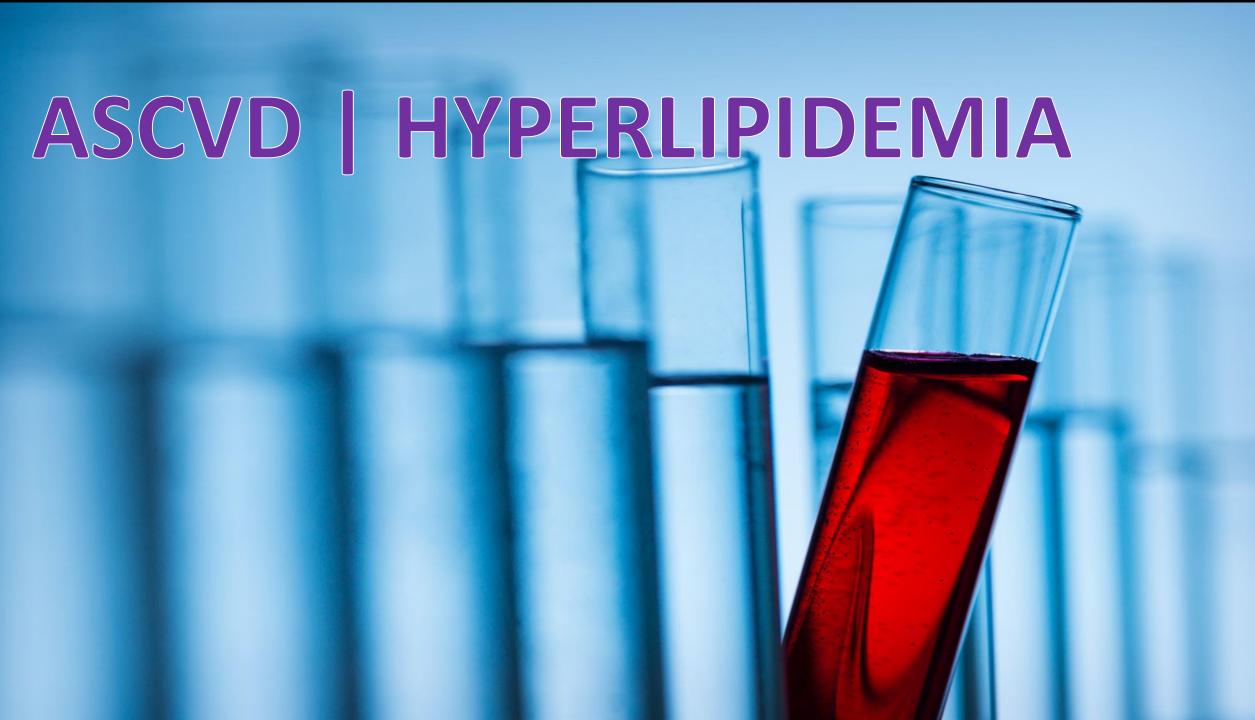
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### ASCVD/Lipids Screening USPSTF 2013

Final Recommendation Statement

#### Lipid Disorders in **Screening** noiesterol, Dysilo.

December 30, 2013

Recommendation ISPSTF are independent of the U.S. government. They show construed as an official Research and Quality or the U.S. Department of Health and position of the Services.



Read t Recommendation Sta

#### ndation Summary Recon

Popula	Recommendation	Grade
Men 35 Older	The USPSTF strongly recomme ening men aged 35 and older for lipid ders.	A
Women d Older at Increased or CHD	The USPSTF strongly recommends screen men aged 45 and older for they are at increased risk for coronary hear	A
Women 20- Increased Risk	The USPSTF recommends screening women age for lipid disor they are at increased risk for coronary heart disease.	В
Men 20-35 at Inch Risk for CHD	The USPSTF recommends screening men aged 20-35 for have a screenin	В
Men 20-35, Women Not at Increased Risk	men. heart disea.  Screening for lipid disorders in are not at increased risk for coronary heart disea.	C





# ASCVD/Lipids Screening: USPSTF 2022

Population	Recommendation	Grade
Adults aged 40 to 75 years who have 1 or more cardiovascular risk factors and an estimated 10-year cardiovascular disease (CVD) risk of 10% or greater	The USPSTF recommends that clinicians prescribe a statin for the primary prevention of CVD for adults aged 40 to 75 years who have 1 or more CVD risk factors (ie, dyslipidemia, diabetes, hypertension, or smoking) and an estimated 10-year risk of a cardiovascular event of 10% or greater.	В
Adults aged 40 to 75 years who have 1 or more cardiovascular risk factors and an estimated 10-year CVD risk of 7.5% to less than 10%	The USPSTF recommends that clinicians selectively offer a statin for the primary prevention of CVD for adults aged 40 to 75 years who have 1 or more CVD risk factors (ie, dyslipidemia, diabetes, hypertension, or smoking) and an estimated 10-year risk of a cardiovascular event of 7.5% to less than 10%. The likelihood of benefit is smaller in this group than in persons with a 10-year risk of 10% or greater.	C
Adults 76 years or older	The USPSTF concludes that the current evidence is insufficient to assess the balance of benefits and harms of initiating a statin for the primary prevention of CVD events and mortality in adults 76 years or older.	I

## Steps for Implementing Screening:

- 1. Identify Risk Factors
- 2. Estimate ASCVD Risk in 40-75 y/o
- 3. Start *statin* for primary prevention for >10%
- 4. Use shared-decision making to start statin in 7.5% 10%









## What are USPSTF ASCVD Risk Factors?

- ✓ Diabetes
- ✓ Hypertension
- ✓ Dyslipidemia
- √ Smoking
- ✓ Obesity/overweight
- √ CKD and Albuminuria
- √ Family history of premature coronary disease





# What are ACC/AHA ASCVD Risk-Enhancing Factors?

- FHx premature ASCVD
  - Males <55 y; Females <65 y
- Primary hypercholesterolemia

LDL-C 160-189 mg/dL; non HDL-C 190-219 mg/dL

Metabolic syndrome

Three of any of the following: ↑ waist circumference, ↑ TG, ↑ BP, ↑ glucose, ↓ HDL-C < 40mg/dL in males or < 50mg/dL in females

- CKD
- High-risk race/ethnicities

South Asian ancestry

- Chronic inflammatory conditions
   RA, HIV/AIDS, Psoriasis
- Hx menopause before 40 yo and hx pregnancy-associated high-risk conditions (ie. preeclampsia)
- Lipid/biomarkers

Persistently elevated TG, 1 hs-CRP, 1 Lp(a), 1 apoB or ABI < 0.9

### Comparing ASCVD Risk Factors



- Diabetes
- Hypertension
- Dyslipidemia
- •Smoking
- Obesity/overweight
- CKD and Albuminuria
- •Family history of premature coronary disease



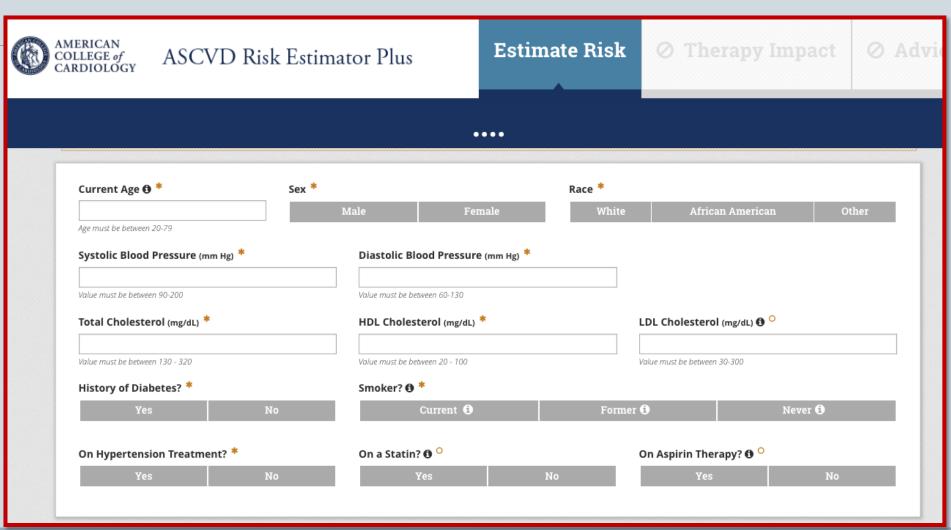
- FHx premature ASCVD
- Primary hypercholesterolemia
- Metabolic syndrome
- CKD
- Chronic inflammatory conditions
- Hx premature menopause and hx pregnancyassociated conditions that increase ASCVD risk
- High-risk race/ethnicities
- Lipid/biomarkers (Lp(a))



# **ACC ASCVD Risk Estimator**



ACC ASCVD Risk Estimator Calculator for internet browser



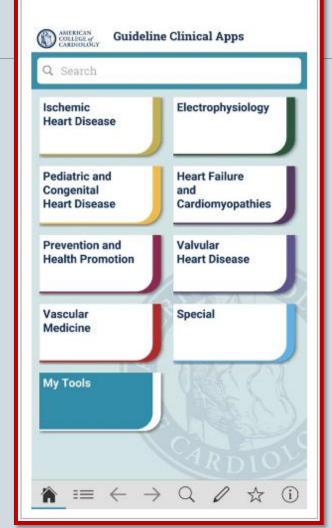


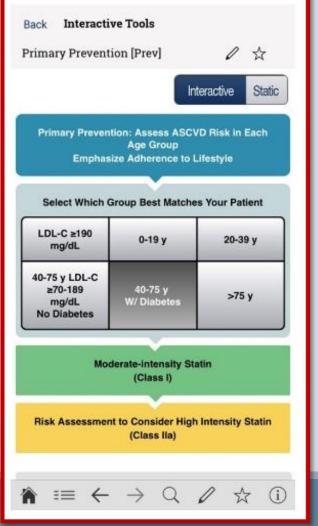
# **ACC Clinical Guideline**

# **Mobile App**

# COMPREHENSIVE MOBILE APP WITH ALL GUIDELINES AND CALCULATORS









# ACC/AHA Treatment: Severe Hypercholesterolemia LDL-C ≥ 190 mg/dL

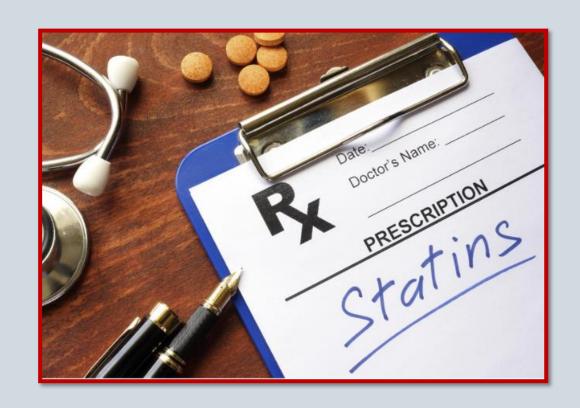
**Start** treatment with maximally tolerated **High Intensity** STATIN therapy

### **Add** Ezetimibe if:

- Patients achieves <50% LDL-C reduction</li>
- LDL-C is greater than 100 mg/dL

### **Add** PCSK9-inhibitor if:

- LDL-C is still greater than 100 mg/dL
- Multiple ASCVD enhancing risk factors

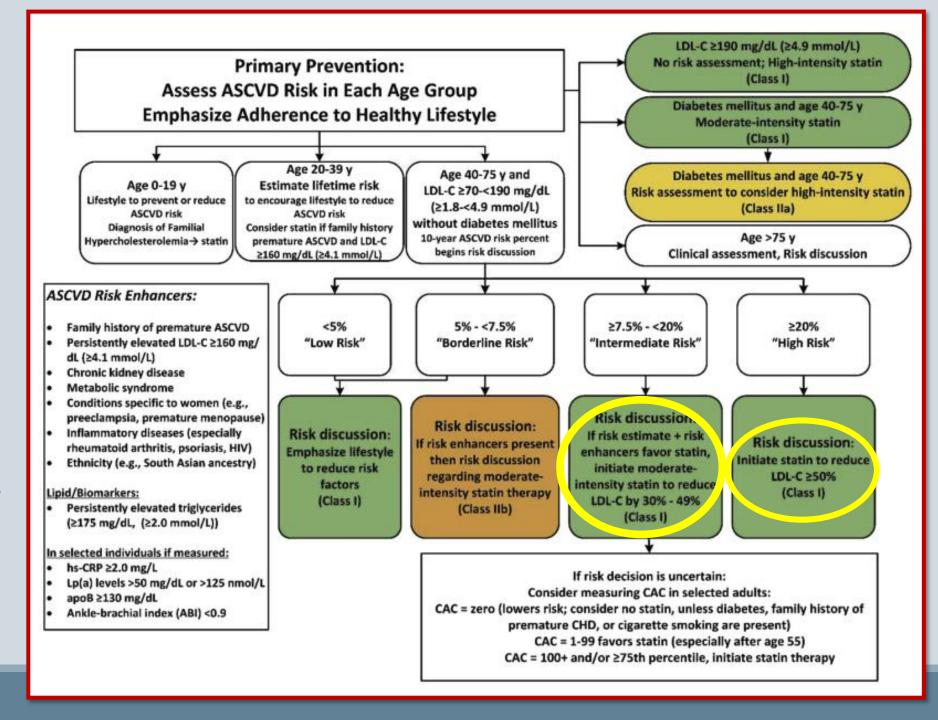




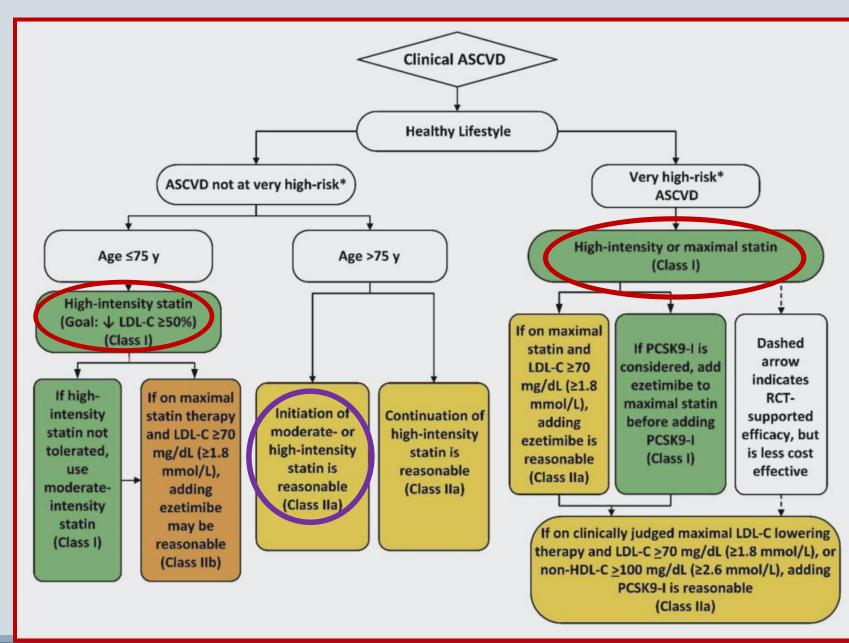
# ASCVD Primary Prevention

≥ 20%: Start High Intensity Statin

≥7.5% - <20%: SDM & Moderate Intensity Statin







# ASCVD Secondary Prevention

(ie-has had a cardiovascular event)

# High Intensity Statin:

- \*Very HIGH Risk ASCVD
- \* ≤75 y/o & not very high risk
- \* >75 y/o may be reasonable

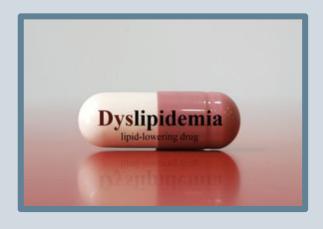


# **Statin Therapy by Intensity**

# **HIGH (≥50%)**

Atorvastatin 40mg, 80mg

Rosuvastatin 20mg, 40mg



# **MODERATE (30%-49%)**

Atorvastatin 10mg, 20mg

Rosuvastatin 5mg, 10mg

Simvastatin 20-40mg

Pravastatin 40mg, 80mg

Lovastatin 40mg

# LOW (<30%)

Simvastatin 10mg

Pravastatin 10mg-20mg

Lovastatin 20mg

# **Treatment Toolbag**:





# **Ezetemibe**

## When do we use this agent?

- •Severe primary hypercholesteremia already on statin to get to goal of less than 100 mg/dL
- •Very high risk ASCVD who have not achieved LDL goal of less than 70 mg/dL while on maximally tolerated statins

### How does this work?

Inhibits cholesterol absorption in the small intestine

### Side effects?

Headache, runny nose, sore throat. Less common: diarrhea, body aches, joint aches, fatigues, weakness

# **Treatment Toolbag**:



# **PCSK9 Inhibitors**



### When do we use this agent?

- •Very high risk ASCVD if already on max-tolerated statin and ezetimibe to get to LDL goal of less than 70 mg/dL
- Primary hypercholesteremia or familial hypercholesterolemia

### How does this work?

•PCSK9 proteins block LDL receptors in the liver that clear LDL

### Side effects?

Nasopharyngitis, injection-site reaction

# **Treatment Toolbag**:



# **Bempedoic Acid**



### When do we use this agent?

- Statin-intolerant patients who have not achieved LDL reduction
- Can be used in combination with statins and ezetimibe

### How does this work?

- •Inhibits ACL enzyme which decreases cholesterol production in the liver,
- Excreted by the kidneys

### **Side effects?**

Uric acid increase (gout?), tendon rupture (<0.5%)

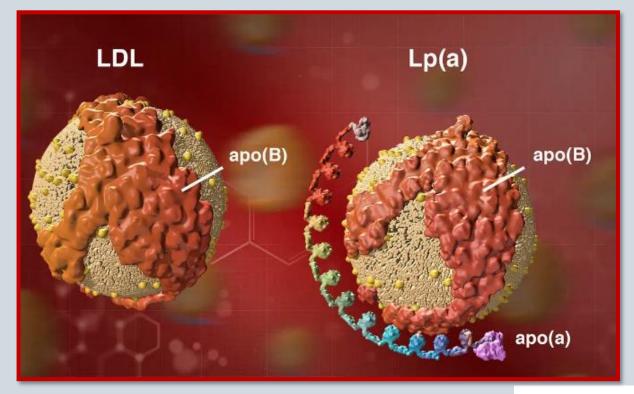
# Quick Word about ApoB and Lp(a)

### **ApoB**

Group of proteins strongly associated with ASCVD and atherosclerosis

# Lp(a)

- •Inherited trait that affects **20**% of population
- •Elevated Lp(a) ≥50 mg/dL or ≥125 nmol/L is considered **risk enhancing** property
- •Risk factor for early CAD and aortic stenosis





# Quick Word about Triglycerides

If Hypertriglyceridemia is,	Then,
Moderate (175-499 mg/dL)	Treat lifestyle factors such as obesity, secondary factors (such as diabetes, CKD, liver or thyroid disease) and address meds that increase TG in adults > 20 years old
Moderate to Severe (≥ 500 mg/dL) and ASCVD risk ≥ 7.5%	Start statin if TG are persistently elevated after lifestyle and secondary factors are addressed in adults 40-75 y/o
<b>Severe</b> (≥ 500 mg/dL, fasting) and ASCVD risk ≥ 7.5%	Start statin and address reversible causes of high TG

- •Notice: Fibrates and Niacin are no longer recommended
- •AACE 2022 Update recommends adding fibrates or icosapent ethyl if fasting TG are persistently over >200 mg/dL
- You could check triglycerides in both fasting and non-fasting states

- •If the triglycerides are VERY high, LDL-C will not be calculated by the lab
- •Still not at goal, refer your patients to endocrinologists or cardiologists

# Quick Word about CAC Score

### CORONARY CALCIUM SCORE CHART









NORMAL ARTERY

BEGINNING OF PLAQ FORMATION

FATTY I

NARROWED ARTERY BLOCKED BY A BLOOD CLOTH

The amount of calcium present in the coronary arteries is scored according to the Agatston scale, as follows:

0	No identifiable calcium deposits
1-10	Low Risk. Less than 10% chance of heart disease
11-99	MILD calcium deposits
100-399	MODERATE calcium deposits
400-999	SEVERE calcium deposits
1000+	25% chance of heart attack within a year

Helpful to further risk stratify patients into statin therapy (ie: "tie breaker")

CAC = 0 Low ASCVD risk

- Consider repeating in 5-10 years if statin therapy has not been started
- "Down Risk " Caution: Score could be 0 in patients with Risk Enhancing Factors

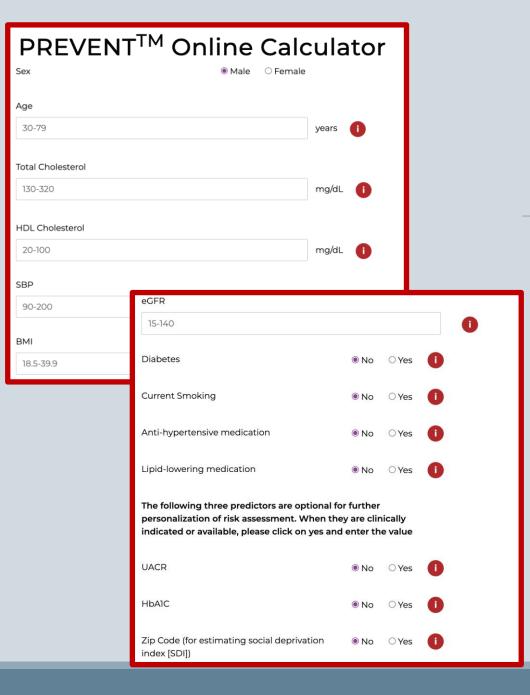
CAC = 1-99 Intermediate ASCVD risk

CAC ≥ 100 Considered High ASCVD risk

CAC scoring is **NOT useful** in patients taking statins because these will increase CAC score

# Quick word about Aspirin and ASCVD —

- Aspirin (75-162mg/day) can be used as <u>secondary</u> prevention in patients with DM and ASCVD
  - Shared decision making should be used to consider aspirin as primary prevention in patients at increased ASCVD risk
- DAPT: Dual antiplatelet therapy (low-dose Aspirin and Clopidogrel)
  - Reasonable for <u>1 year</u> after acute coronary syndrome (but could be extended)
  - Long term use should be considered in patients with high ischemic risk, prior coronary intervention and low bleeding risk to prevent further cardiovascular events



# PREVENT Online Calculator



- New Calculator: PREVENT AHA
   Predicting Risk of CVD Events
- Used in primary prevention
   patients ages 30-79 years and
   without coronary heart disease,
   stroke, or heart failure
- •Incorporates obesity, diabetes and CKD into the equation for calculating CVD risks in cardiovascular-kidney-metabolic syndrome.





# DIABETES



# **Diabetes Screening:** USPSTF

### **Recommendation (Grade B):**

- Screen for prediabetes/diabetes:
  - Age 35-70
  - WITH BMI >25-30 (or >23 in Asian American)
  - Without symptoms of diabetes

**<u>KEY CHANGE</u>**: Initial screening age 35 (previously age 40)

### **Screening tests:**

 Fasting blood glucose (FBG) or HbA1c or OGTT

### Frequency:

Interval is uncertain

### **Preventive Interventions:**

- Lifestyle modifications and metformin have both demonstrated efficacy in slowing progression to diabetes
- Metformin has not yet been approved for prediabetes treatment



USPSTF
Prediabetes &
Diabetes
Screening

What does the USPSTF recommend?	Adults aged 35 to 70 years who have overweight or obesity:  • Screen for prediabetes and type 2 diabetes, and offer or refer patients with prediabetes to effective preventive interventions. Grade: B
To whom does this recommendation apply?	Nonpregnant adults aged 35 to 70 years who have overweight or obesity are 1 no symptoms of diabetes.
What's new?	The USPSTF has lowered the starting age of screening from 40 to 35 years.
How to implement this recommendation?	<ol> <li>Assess risk:         <ul> <li>Obtain height and weight measurements to determine whether patient has overweight or obesity. Overweight and obesity are defined as a BMI ≥25 and ≥30, respectively.</li> </ul> </li> <li>Screen:         <ul> <li>If the patient is aged 35 to 70 years and has overweight or obesity.</li></ul></li></ol>
How often?	The optimal screening interval for adults with an initial norma glucose test result is uncertain. Screening every 3 years r be a reasonable approach for adults with normal blood glucose levels.
What are other relevant USPSTF recommendations?	The USPSTF has made a recommendation on behavioral weight loss interventions to prevent obesity-related morbidity and mortality in adults with a BMI ≥30. This recommendation is available at https://www.uspreventiveservicestaskforce.org
Where to read the full recommendation statement?	Visit the USPSTF website (https://www.uspreventiveservicestaskforce.org) to read the full recommendation statement. This includes more details on the rationale of the recommendation, including benefits and harms; supporting evidence; and recommendations of others.

The USPSTF recognizes that clinical decisions involve more considerations than evidence alone. Clinicians should understand the evidence but individualize decision-making to the specific patient or situation.



#### Figure Legend:

Clinician Summary: Screening for Prediabetes and Type 2 Diabetes BMI indicates body mass index (calculated as weight in kilograms divided by height in meters squared); HbA<sub>1c</sub>, hemoglobin A<sub>1c</sub>; USPSTF, US Preventive Services Task Force.

# Diabetes Screening: ADA Risk-Centered Approach

# Overweight/obese with ≥ 1 risk factors:

- First-degree relative with diabetes
- High-risk race/ethnicity
- History of CVD, HTN (treated or untreated)
- Abnormal Lipids
  - Low HDL (<35 mg/dL) and/or High TG (>250 mg/dL)
- Conditions associated with insulin resistance
  - PCOS
- Physical inactivity

### **Interval of Screening**

- Annually in patients with prediabetes
- Repeat at least every 3 years or more frequently based on results

### **Special Populations**

- History of gestational diabetes (GDM) get lifelong screening in 3-year intervals
- HIV patients

# Key Differences Between Authorities

### **USPSTF**

- Screening initiated at age 35 for any asymptomatic individual
- No optimal screening interval but 3-year interval may be reasonable

### **ADA**

- For initial screening, prioritizes risk factors over start age of 35
- Annual screening interval advised for people with prediabetes
- Three-year screening interval for patients who had GDM
- Three-year screening interval (OR more frequent) depending on risk factors or initial results
- Patients with HIV

# **DIABETES DIAGNOSTIC CRITERIA:**

# **ADA GUIDELINES**

# American Diabetes Association criteria for the diag nosis of diabetes

1 A1C ≥6.5%. The test should be performed in a laboratory using a method that is NGSP certified and standardized to the DCCT assay.\*

#### OR

2 FPG ≥126 mg/dL (7 mmol/L). Fasting is defined as no caloric intake for at least 8 hours.\*

#### OR

2-hour plasma glucose ≥200 mg/dL (11.1 mmol/L) during an OGTT. The test should be performed as described by the World Health Organization, using a glucose load containing the equivalent of 75 g anhydrous glucose dissolved in water.\*

#### OR

 In a patient with classic symptoms of hyperglycemia or hyperglycemic crisis, a random plasma glucose ≥200 mg/dL (11.1 mmol/L).

### **Diagnosis**

### requires **TWO** results

from the same or different/subsequent samples

(except in scenario #4)

#### Acronyms:

A1C: glycated hemoglobin; NGSP: National Glycohemoglobin Standardization Program; DCCT: Diabetes Control and Complications Trial; FPG: fasting plasma glucose; OGTT: oral glucose tolerance test.

\* In the absence of unequivocal hyperglycemia, diagnosis requires two abnormal test results from the same sample or in two separate test samples.

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# **DIABETES TARGET GOALS:** HbA1c

## **Treatment Goals**

ADA: < 7%</li>

• AACE: <6.5%

• ACP: 7% - 8%



## **Keep in mind:**

- A1c could be falsely elevated in conditions with decreased RBC turnover --Anemias (B12 or folate deficiency)
- A1c could be falsely decreased in conditions with increased RBC turnover -- End-stage renal disease (ESRD)



# **DIABETES TREATMENT:**

# Pharmacologic Options (Think TWO Buckets)

### **EVERYTHING ELSE:** (MOSTLY)

- Metformin
- GIP/GLP-1 receptor agonist (eg tirzepatide)
- GLP-1 receptor agonist (eg. semaglutide)
- SGLT-2 inhibitors (eg. dapagliflozin)
- Thiazolidinediones (TZD)
   (eg. pioglitazone)
- **DPP-4 inhibitors** (eg. sitagliptan)
- Sulfonylureas (SU)
   (eg. glipizide)

**INSULIN:** (OUTSIDE SCOPE OF TODAY'S WORKSHOP)

- Basal: Long acting
- **Prandial**: (aka: bolus)
  - Short acting, tons of options
- Start in cases of severe hyperglycemia
  - ∘ FBG >250 mg/dL
  - Random >300 mg/dL
  - HbA1c at or above 9%

# DIABETES TREATMENT: Two Authorities, One Mission

**ADA** 

AMERICAN DIABETES
ASSOCIATION

**AACE** 

AMERICAN ASSOCIATION OF CLINICAL ENDOCRINOLOGISTS





### **American Diabetes Association**



### USE OF GLUCOSE-LOWERING MEDICATIONS IN THE MANAGEMENT OF TYPE 2 DIABETES

HEALTHY LIFESTYLE BEHAVIORS; DIABETES SELF-MANAGEMENT EDUCATION AND SUPPORT (DSMES); SOCIAL DETERMINANTS OF HEALTH (SDOH)

REGULARLY (3-6 MONTHS)

Goal: Cardiorenal Risk Reduction in High-Risk Patients with Type 2 Diabetes (in addition to comprehensive CV risk management)\*

#### +ASCVD†

Defined differently across CVOTs but all included individuals with established CVD (e.g., MI, stroke, any revascularization procedure) Variably included: conditions such as transient ischemic attack, unstable angina. amputation, symptomatic or asymptomatic coronary artery disease.

GLP-1 RA# with proven

**CVD** benefit

proven CVD benefit or vice versa

TZD^

+ASCVD/Indicators of High Risk

EITHER/

If A1C above target

For patients on a GLP-1 RA, consider adding SGLT2i with

#### +Indicators of high risk

While definitions vary, most comprise ≥55 years of age with two or more additional risk factors (including obesity hypertension, smoking, dyslipidemia, or albuminuria)

SGLT2i§ with proven

CVD benefit

**Current or prior** symptoms of HF with documented HFrEF or HFpEF

+HF

SGLT2i§

with proven

HF benefit

in this

population

#### +CKD

eGFR <60 mL/min per 1.73 m<sup>2</sup> OR albuminuria (ACR ≥3.0 mg/mmol [30 mg/g]). These measurements may vary over time; thus, a repeat measure is required to document CKD.

+CKD (on maximally tolerated dose of ACEi/ARB)

#### **PREFERABLY**

SGLT2i§ with primary evidence of reducing CKD progression

Use SGLT2i in people with an eGFR ≥20 mL/min per 1.73 m<sup>2</sup>; once initiated should be continued until initiation of dialysis or transplantation

GLP-1 RA with proven CVD benefit if SGLT2i not tolerated or contraindicated

- - - - - NR -

If A1C above target, for patients on SGLT2i, consider incorporating a GLP-1 RA or vice versa

#### Glycemic Management: Choose approaches that provide the

Metformin OR Agent(s) including COMBINATION therapy that provide adequate EFFICACY to achieve and maintain treatment goals

efficacy to achieve goals:

Consider avoidance of hypoglycemia a priority in high-risk individuals

In general, higher efficacy approaches have greater likelihood of achieving glycemic goals

Efficacy for glucose lowering

#### Very High:

Dulaglutide (high dose), Semaglutide, Tirzepatide

Insulin

Combination Oral, Combination Injectable (GLP-1 RA/Insulin)

GLP-1 RA (not listed above), Metformin, SGLT2i, Sulfonylurea, TZD

> Intermediate: DPP-4i

#### **Achievement and Maintenance of Weight Management Goals:**

Set individualized weight management goals

General lifestyle advice: medical nutrition therapy/eating patterns/ physical activity

Goal: Achievement and Maintenance of Glycemic and Weight Management Goals

Intensive evidencebased structured weight management program

**Consider medication** for weight loss

Consider metabolic surgery

#### When choosing glucose-lowering therapies:

Consider regimen with high-to-very-high dual glucose and weight efficacy

Efficacy for weight loss

#### Very High:

Semaglutide, Tirzepatide

#### High:

**Dulaglutide**, Liraglutide

#### Intermediate:

GLP-1 RA (not listed above). SGLT2i

#### Neutral:

DPP-4i, Metformin

If A1C above target

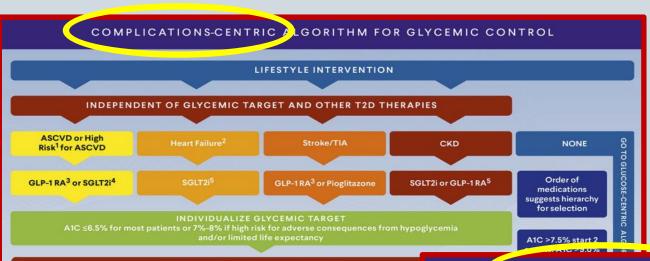
γευριε with mr. thu, established CVD or multiple risk factors for CVD, the decision to use a GLP-1 RA or SGLT2i with proven benefit should be independent of background use of metformin;† A strong

If additional cardiorenal risk reduction or glycemic lowering needed

#### Identify barriers to goals:

- Consider DSMES referral to support self-efficacy in achievement of goals
- Consider technology (e.g., diagnostic CGM) to identify therapeutic gaps and tailor therapy
- · Identify and address SDOH that impact achievement of goals

mendation is warranted for people with CVD and a weaker recommendation for those with indicators of high CV risk. Moreover, a higher absolute risk reduction and thus lower numbers needed to treat are seen at higher levels of baseline risk and should be factored into the shared decision-making process. See text for details; ^ Low-dose TZD may be better tolerated and similarly effective; & For SGLT2i, CV. ral outcomes trials demonstrate their efficacy in reducing the risk of composite MACE, CV death, all-cause mortality, MI, HHF, and renal outcomes in individuals with T2D with established/high risk of CVD; GLP-1 RA, CVOTs demonstrate their efficacy in reducing composite MACE, CV death, all-cause mortality, MI, stroke, and renal endpoints in individuals with T2D with established/high risk of CVD.







Continue or start metformin if appropriate

If not at glycemic target at <3 months, titrate to maximum tolerated dose or add agent

SGLT2i4 or GLP-1 RA

Pioglitazone<sup>2</sup> or GLP-1 RA

GL

IF NOT AT GOAL: CONTINUE TO GLUCOSE-CENTRIC ALGORITHM FOR OR ALGORITHM FOR ADDING/INTENSIFYING INSU

High risk for ASCVD: albuminuria or proteinuria, hypertension and left ventricular (LV) hypertrophy, LV systolic or diastolic dysfunction, <sup>2</sup>TZDs are contraindicated in NYHA Class III/IV HF. <sup>3</sup>ASCVD: liraglutide/semaglutide/dulaglutide or Stroke: semaglutide/dulagl 4canagliflozin/empagliflozin. 5Use SGLT2i or GLP-1 RA with proven benefit.

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# GLUCOSE-CENTRIC ALGO ITHM FOR GLYCEMIC CONTROL

#### LIFESTYLE INTERVENTION

Start or continue metformin if appropriate1

#### INDIVIDUALIZE GLYCEMIC TARGET

Patients may Overweight or Obesity<sup>2</sup> Hypoglycemia Risk<sup>3</sup> Access / Cost Severe Hyperglycemia4 present with >1 scenario Order of medications suggests hierarchy for selection A1C >7.5% start 2 DPP-4i<sup>8</sup> or TZD DPP-4i8 or TZD9 agents, A1C >9.0% or >1.5% above goal start 2-3 agents GLP-1 RA | GIP/GLP-1 RA | Other agents likely Avoid SU/GLN Avoid SU/GLN or Not Preferred SGLT2i COLSVL ineffective in the setting of glucotoxicity BRC-QR Titrate to maximum tolerated dose. If not at glycemic target at ≤3 months, add best available agent not in use<sup>7</sup>

GLP-1 RA | GIP/GLP-1 RA | SGLT2i | TZD | DPP-4i | SU/GLN | COLSVL | BRC-QR | PRAML11

#### IF NOT AT GOAL: CONTINUE TO ALGORITHM FOR ADDING/INTENSIFYING INSULIN

<sup>1</sup>Take with food with dose titration for enhanced tolerance. <sup>2</sup>See also COMPLICATIONS-CENTRIC MODEL FOR THE CARE OF PERSONS WITH OVERWEIGHT/OBESITY and PROFILES OF WEIGHT-LOSS MEDICATIONS table, 3 Evaluate for issues leading to hypoglycemia or hypoglycemia unawareness and manage with patient-centered strategies, 4 If A1C > 10% and/or BG ≥300 with symptomatic hyperglycemia, reduce glucose/AIC as promotivand safely as possible. See also ALGORITHM FOR ADDING/INTENSIFYING INSULIN 6 GLP-1 RA requires titration phase which can delay glycemic control. After glucose toxicity is resolved consider adding other agents, 7 See also PROFILES OF ANTIHYPERGLYCEMIC MEDICATIONS table, 8 GLP-1 RA and DPP-4i should not be combined. 9TZD can cause fluid retention but have benefit for NAFLD, CVD prevention, dyslipidemia. 10 Access/Cost are dependent on location of the market. Insulin costs vary widely with devices (e.g., pens versus vials) and formulations (e.g., analogues versus combinations such as 70/30). 11 PRAML is used as an adjunct with prandial insulin.

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Algorithm Figure 7-Glucose-Centric Glycemic Control





#### COMPLICATIONS-CENTRIC ALGORITHM FOR GLYCEMIC CONTROL

#### LIFESTYLE INTERVENTION INDEPENDENT OF GLYCEMIC TARGET AND OTHER T2D THERAPIES ASCVD or High Heart Failure<sup>2</sup> Stroke/TIA NONE CKD Risk<sup>1</sup> for ASCVD 7 GLUCOSE-CENTRIC Order of GLP-1 RA3 or SGLT2i4 SGLT2i5 GLP-1 RA<sup>3</sup> or Pioglitazone SGLT2i or GLP-1 RA<sup>5</sup> medications suggests hierarchy for selection INDIVIDUALIZE GLYCEMIC TARGET A1C ≤6.5% for most patients or 7%-8% if high risk for adverse consequences from hypoglycemia ALGORITHM FOR GLYCEMIC and/or limited life expectancy A1C >7.5% start 2 agents, A1C > 9.0% or >1.5% above goal Continue or start metformin if appropriate start 2-3 agents If A1C >10% and/or If not at glycemic target at <3 months, titrate to maximum tolerated dose or add agent not in use glucose >300 mg/dL with symptomatic hyperglycemia, use basal insulin SGLT2i4 or GLP-1 RA Pioglitazone<sup>2</sup> or GLP-1 RA GLP-1 RA or SGLT2i5 GLP-1RA +/- GLP-1 RA IF NOT AT GOAL: CONTINUE TO GLUCOSE-CENTRIC ALGORITHM FOR GLYCEMIC CONTROL OR ALGORITHM FOR ADDING/INTENSIFYING INSULIN High risk for ASCVD; albuminuria or proteinuria, hypertension and left ventricular (LV) hypertrophy, LV systolic or diastolic dysfunction, ankle-brachial index <0.9.</p> <sup>2</sup>TZDs are contraindicated in NYHA Class III/IV HF. <sup>3</sup>ASCVD: liraglutide/semaglutide/dulaglutide or Stroke: semaglutide/dulaglutide.









4canagliflozin/empagliflozin. 5Use SGLT2i or GLP-1 RA with proven benefit.

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#### GLUCOSE-CENTRIC ALGORITHM FOR GLYCEMIC CONTROL

#### LIFESTYLE INTERVENTION

Start or continue metformin if appropriate<sup>1</sup>

#### INDIVIDUALIZE GLYCEMIC TARGET

A1C ≤6.5% for most persons or 7%-8% if high risk for adverse consequences from hypoglycemia and/or limited life expectancy

Overweight or Obesity<sup>2</sup>

Hypoglycemia Risk<sup>3</sup>

Access / Cost

Severe Hyperglycemia4

Patients may present with >1 scenario

Preferred

GLP-1 RA or GIP/GLP-1 RA or SGLT2i GLP-1 RA or GIP/GLP-1 RA or SGLT2i

TZD or SU/GLN

Basal Insulin<sup>5</sup> + Prandial Insulin or + GLP-1 RA I GIP/GLP-1 RA<sup>6</sup> Order of medications suggests hierarchy for selection<sup>7</sup>

Alternatives

DPP-4i<sup>8</sup> or TZD<sup>9</sup>

DPP-4i8 or TZD

Insulin or DPP-4i10

Basal Insulin + other agent(s) A1C >7.5% start 2 agents, A1C >9.0% or >1.5% above goal start 2-3 agents

Concerns or Not Preferred

Avoid SU/GLN

Avoid SU/GLN

GLP-1 RA | GIP/GLP-1 RA | SGLT2i | COLSVL BRC-QR

Other agents likely ineffective in the setting of glucotoxicity<sup>5</sup>

Titrate to maximum tolerated dose. If not at glycemic target at ≤3 months, add best available agent not in use <sup>7</sup> GLP-1 RA | GIP/GLP-1 RA | SGLT2i | TZD | DPP-4i | SU/GLN | COLSVL | BRC-QR | PRAML<sup>11</sup>

#### IF NOT AT GOAL: CONTINUE TO ALGORITHM FOR ADDING/INTENSIFYING INSULIN

<sup>1</sup>Take with food with dose titration for enhanced tolerance. <sup>2</sup>See also COMPLICATIONS-CENTRIC MODEL FOR THE CARE OF PERSONS WITH OVERWEIGHT/OBESITY and PROFILES OF WEIGHT-LOSS MEDICATIONS table. <sup>3</sup>Evaluate for issues leading to hypoglycemia or hypoglycemia unawareness and manage with patient-centered strategies. <sup>4</sup>If AIC >10% and/or BG ≥300 with symptomatic hyperglycemia, reduce glucose/AIC as promptly and safely as possible. <sup>5</sup>See also ALGORITHM FOR ADDING/INTENSIFYING INSULIN. <sup>6</sup>GLP-1 RA requires titration phase which can delay glycemic control. After glucose toxicity is resolved, consider adding other agents. <sup>7</sup>See also PROFILES OF ANTIHYPERGLYCEMIC MEDICATIONS table. <sup>8</sup>GLP-1 RA and DPP-4i should not be combined. <sup>9</sup>TZD can cause fluid retention but have benefit for NAFLD, CVD prevention, dyslipidemia. <sup>10</sup>Access/Cost are dependent on location of the market. Insulin costs vary widely with devices (e.g., pens versus vials) and formulations (e.g., analogues versus combinations such as 70/30). <sup>11</sup>PRAML is used as an adjunct with prandial insulin.

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Algorithm Figure 7-Glucose-Centric Glycemic Control











# **DIABETES TREATMENT:**

(All Authorities Agree - ADA/AACE)

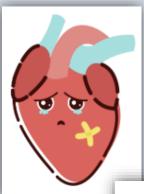
## FIRST LINE TREATMENTS (EVERYONE!)

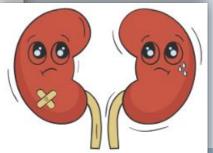
- Lifestyle modifications
  - Diet
  - Exercise
    - AHA recommends weekly goals of:
      - 150 mins of moderate intensity exercise
      - OR 75 mins of vigorous intensity exercise



### American Diabetes Association







Goal: Cardiorenal Risk Reduction in High-Risk Patients with Type 2 Diabetes (in addition to comprehensive CV risk management)\*

#### +ASCVD† +Indicators of high risk +CKD +HF **Defined differently across** While definitions vary, most eGFR <60 mL/min per 1.73 m<sup>2</sup> OR **Current or prior** CVOTs but all included albuminuria (ACR ≥3.0 mg/mmol comprise ≥55 years of age symptoms [30 mg/g]). These measurements individuals with established with two or more additional of HF with CVD (e.g., MI, stroke, any risk factors (including obesity, may vary over time; thus, a repeat documented measure is required to document CKD. revascularization procedure). hypertension, smoking, **HFrEF or HFpEF** dyslipidemia, or albuminuria) Variably included: conditions such as transient ischemic attack, unstable angina. +CKD (on maximally tolerated dose amputation, symptomatic of ACEi/ARB) or asymptomatic coronary +HF artery disease. **PREFERABLY** SGLT2i§ SGLT2i§ with primary evidence of with proven reducing CKD progression **HF** benefit +ASCVD/Indicators of High Risk in this Use SGLT2i in people with an eGFR ≥20 mL/min per 1.73 m<sup>2</sup>; once initiated population EITHER/ should be continued until initiation GLP-1 RA# with proven SGLT2i§ with proven of dialysis or transplantation **CVD** benefit CVD benefit GLP-1 RA with proven CVD benefit if SGLT2i not tolerated or contraindicated If A1C above target If A1C above target, for patients on SGLT2i, consider incorporating a • For patients on a GLP-1 RA, consider adding SGLT2i with GLP-1 RA or vice versa proven CVD benefit or vice versa TZD^

If additional cardiorenal risk reduction or glycemic lowering needed

# FIRST LINE THERAPEUTICS ASCVD or HIGH-RISK INDICATORS

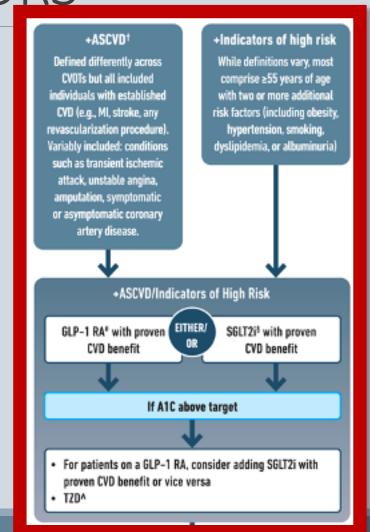


### **GLP-1 RA**

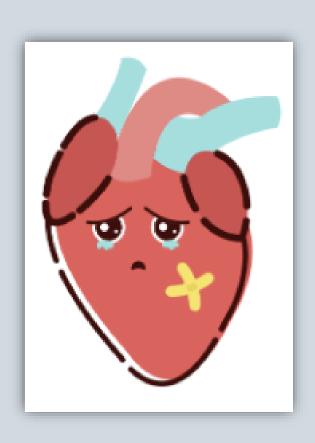
- Do NOT use in thyroid cancer, pancreatic cancer or MEN syndrome
- Side effects: GI issues such as nausea, reflux, diarrhea/constipation

### SGLT2i

- Side effects: GU yeast infx
- Pos: slight weight loss, diuresis may lower BP
- Do not use in GFR <30</li>

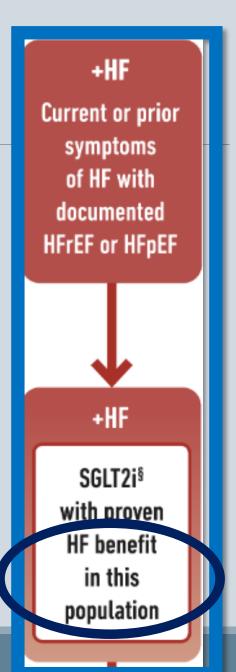


# FIRST LINE THERAPEUTICS Heart Failure



### **Use SGLT2i**

- Neg: GU yeast infx
- Pos: slight weight loss, diuresis may lower
   BP
- Do not use in GFR <30</li>
- As of 2022, consider medications with proven HF benefit such as canagliflozin, dapagliflozin, empagliflozin or ertugliflozin



# FIRST LINE THERAPEUTICS Chronic Kidney Disease (CKD)



- Use SGLT2i with evidence of reducing CKD progression
- •As of 2022, consider medications with **CKD reduction benefit** (canagliflozin, dapagliflozin or empagliflozin)
  - Hold 3-4 days before surgery or during critical illness or prolonged fasting
  - *RARE*: Euglycemic DKA or Fournier gangrene
  - *Side Effect*: GU mycotic infections and glycosuria

#### +CKD

eGFR <60 mL/min per 1.73 m² OR albuminuria (ACR ≥3.0 mg/mmol [30 mg/g]). These measurements may vary over time; thus, a repeat measure is required to document CKD.



+CKD (on maximally tolerated dose of ACEi/ARB)

#### **PREFERABLY**

SGLT2i§ with primary evidence of reducing CKD progression

Use SGLT2i in people with an eGFR ≥20 mL/min per 1.73 m²; once initiated should be continued until initiation of dialysis or transplantation

OR

GLP-1 RA with proven CVD benefit if SGLT2i not tolerated or contraindicated

If A1C above target, for patients on SGLT2i, consider incorporating a GLP-1 RA or vice versa





Goal: Achievement and Maintenance of Glycemic and Weight Management Goals

Glycemic Management: Choose approaches that provide the efficacy to achieve goals:

COMBINATION therapy that provide adequate EFFICACY to achieve and maintain treatment goals

Consider avoidance of hypoglycemia a priority in high-risk individuals

In general, higher efficacy approaches have greater likelihood of achieving glycemic goals

Efficacy for glucose lowering

#### **Very High:**

Dulaglutide (high dose), Semaglutide, Tirzepatide

Insulin

Combination Oral, Combination Injectable (GLP-1 RA/Insulin)

#### High:

GLP-1 RA (not listed above), Metformin, SGLT2i, Sulfonylurea, TZD

> Intermediate: DPP-4i

#### Achievement and Maintenance of Weight Management Goals:

Set individualized weight management goals

General lifestyle advice: medical nutrition therapy/eating patterns/ physical activity Intensive evidencebased structured weight management program

Consider medication for weight loss Consider metabolic surgery

#### When choosing glucose-lowering therapies:

Consider regimen with high-to-very-high dual glucose and weight efficacy

Efficacy for weight loss

Very High:

Semaglutide, Tirzepatide

High:

**Dulaglutide**, Liraglutide

Intermediate:

GLP-1 RA (not listed above), SGLT2i

Neutral:

DPP-4i, Metformin





# **NEXT LINE THERAPEUTICS**

Glycemic Management

•EFFICACY without/less hypoglycemia

 Metformin OR Agent(s) such as combination medications that effectively achieve and maintain glycemic goals







efficacy to achieve goals:

Metformin OR Agent(s) including COMBINATION therapy that provide adequate EFFICACY to achieve and maintain treatment goals

Consider avoidance of hypoglycemia a priority in high-risk individuals

In general, higher efficacy approaches have greater likelihood of achieving glycemic goals

Efficacy for alucose lowering

Very High:

Dulaglutide (nigh dose), Semaglutide, Tirzepatide

Insulin

Combination Oral, Combination Injectable (GLP-1 RA/Insulin)

High:

GLP-1 RA (not listed above), Metformin, SGLT2i, Sulfonylurea, TZD

> Intermediate: DPP-4i

# **NEXT LINE THERAPEUTICS**

# Weight Management

•Select diabetes therapies that will control BOTH glucose and weight

General lifestyle modification counseling

•Consider referral to surgery or weight loss medications

•Implement evidence-based weight management

### Achievement and Maintenance of Weight Management Goals:

Set individualized weight management goals

General lifestyle advice: medical nutrition therapy/eating patterns/ physical activity Intensive evidencebased structured weight management program

Consider medication for weight loss

Consider metabolic surgery

When choosing glucose-lowering therapies:

Consider regimen with high-to-very-high dual glucose and weight efficacy

Efficacy for weight loss

Very High:

Semaglutide, Tirzepatide

High:

**Dulaglutide**, Liraglutide

Intermediate:

GLP-1 RA (not listed above), SGLT2i

Neutral:

DPP-4i, Metformin

# **NEXT LINE THERAPEUTICS**

# Weight Management

Select therapies based on WEIGHT LOSS EFFICACY

## **Very High**

Semaglutide, Tirzepatide

## High

Dulaglutide, Liraglutide

### **Intermediate**

GLP-1 not listed above (such as Exentide), SGLT2i

### **Neutral**

DPP-4i, Metformin

#### Achievement and Maintenance of Weight Management Goals:

Set individualized weight management goals

General lifestyle advice: medical nutrition therapy/eating patterns/ physical activity Intensive evidencebased structured weight management program

Consider medication for weight loss Consider metabolic surgery

#### When choosing glucose-lowering therapies:

Consider regimen with high-to-very-high dual glucose and weight efficacy

Efficacy for weight loss

#### Very High:

Semaglutide, Tirzepatide

#### High:

Dulaglutide, Liraglutide

#### Intermediate:

GLP-1 RA (not listed above), SGLT2i

#### Neutral:

DPP-4i, Metformin

## But what about Metformin as first line?



- Neutral to potential effect on cardiovascular risk
- Low risk of hypoglycemia, low to modest weight loss
- Take with food to lessen GI side effects and monitor for B12 Deficiency
- •CKD implications
  - Can be used in eGFR >30
  - Do NOT start if eGFR <45</li>
  - Reduce dose between eGFR 30-45
- Recommended for prediabetes by ADA and AACE

# What about Tirzepatide?

### Classification

 Dual glucagon-like peptide 1/glucose dependent insulinotropic polypeptide receptor agonist (gip/glp-1)

Very high weight loss potential

Cost: Very High

Cardiovascular and Renal benefit studies are underway



	Efficacy <sup>1</sup>	Hypogly- cemia	Weight change <sup>2</sup>	CV effects		Renal effects	
				Effect on MACE	HF	Progression of DKD	Dosing/use considerations*
GIP and GLP-1 RA	Very high	No	Loss (very high)	Under investigation	Under investigation	Under investigation	See label for renal dose considerations     No dose adjustment     Monitor renal function when initiating or escalating doses in patients with renal impairment reporting severe adverse GI reactions

# Avoiding Hypoglycemia



### **Definition**

- Decrease in glucose concentration <70 mg/dL</li>
- However, this is a moving target based on the individual

### **Risk Factors**

- Medications (Insulin/sulfonylurea)
- Exercise
- Frequency of Meals (food insecurities)

### Signs/Symptoms

Tremor

**Palpitations** 

Sweating

Confusion

Dizziness

Anxious/angry/hangry

### **Treatment Sugar!**

Glucose tablets

Soda/juices

Raisins

Sugar/honey

# Avoiding Hypoglycemia

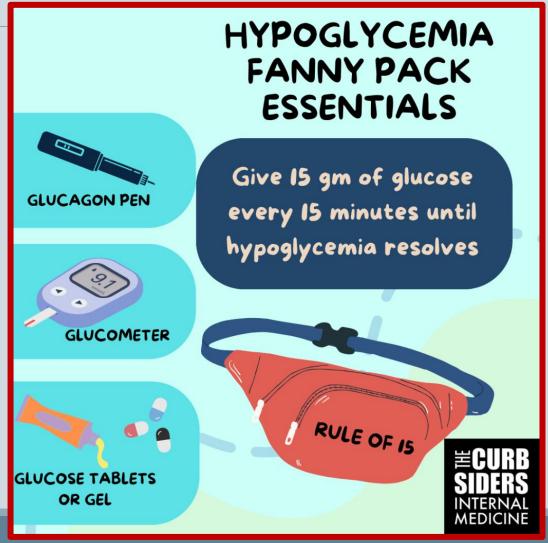
### **Definition**

- Decrease in glucose concentration <70 mg/dL</li>
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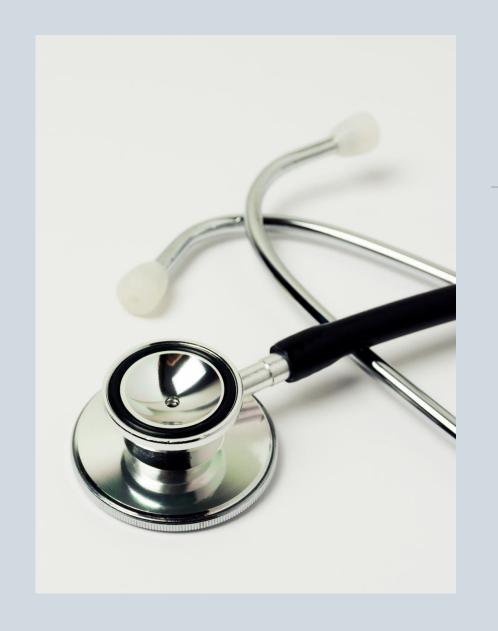
### **Risk Factors**

- Medications (Insulin/sulfonylurea)
- Exercise
- Frequency of Meals (food insecurities)









# Objective

## Counsel and motivate

diverse patient populations on cardiovascular health

with culturally informed lifestyle modifications

while considering social determinants of health

# Language, Disease and Inclusivity

Mindful goal: Avoid labeling our patients

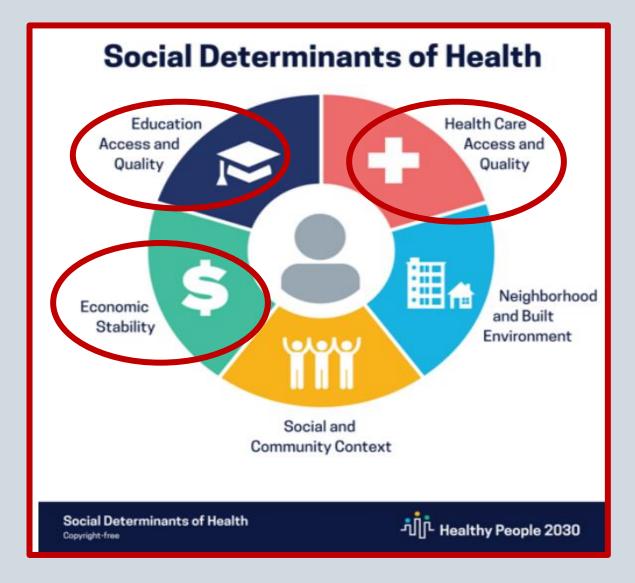
Patients are **NOT** 'diabetics' - they have a disease called diabetes

Patients are **NOT** 'obese' - they have a disease called obesity

In summary, use PEOPLE FIRST language

What do we mean by social determinants of health?





# **Patient Considerations**

### **Health Care Access and Quality**

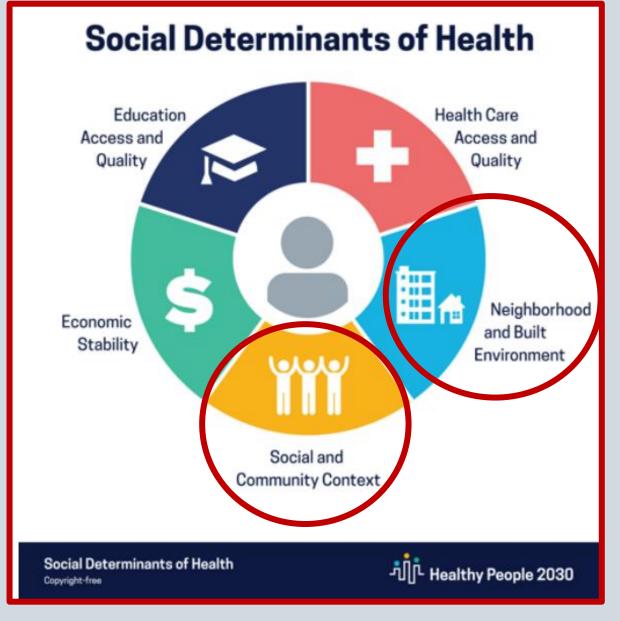
- Patients who live alone/limited assistance
- Distance to nearest clinics or hospitals
- Cost of medications and testing
  - Consider telemedicine visit, generic medications

### **Education Access and Quality**

- Consider health literacy of patient AND caregivers
- Language barriers
  - Provide materials in native language, consider limited access to internet

### **Economic Stability**

- Fixed income
- Unemployed
  - Engage team in identifying patient assistance options (GOODRX or COVERAGE app)



# **Patient Considerations**

### **Social and Community Context**

- Patients who live alone/limited assistance
- Multi-generation households
- Recent immigrants/Refugees
- Homelessness
  - ASK ASK ASK!

### **Neighborhood and Built Environment**

- Food deserts and access to healthy foods
- Transportation
- Faith-based community outreach
- Neighborhood safety and its implication on exercise
- Homelessness
  - Reconsider language of 'noncompliance'
  - Change locus of control what can we do to help them?

# Isabel's 5Cs of Treatment Considerations

#### Collect ALL Data

- History
- Exam
- Screening Results
- Lab Results

#### **Comorbidities**

- Cardiovascular
  Disease
- CKD
- Obesity

### **Complications**

- Complications from the disease
- Complications from the Rx = Side effects

#### Compliance with Treatment

- Select treatments that offer easier compliance
- Avoid labeling as "non-compliant"

### **Cost and Coverage**

 Consider possibility of limited resources/financial constraints

- Meet each person where they are
- **GOAL:** Best Outcomes for every patient regardless of situation

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