Update Round-up! Applying the Latest Guidelines for Diabetes and Cardiovascular Disease

AAPA WE ARE FAMILY (Medicine) Conference San Diego, California 2023

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BAYLOR COLLEGE OF MEDICINE

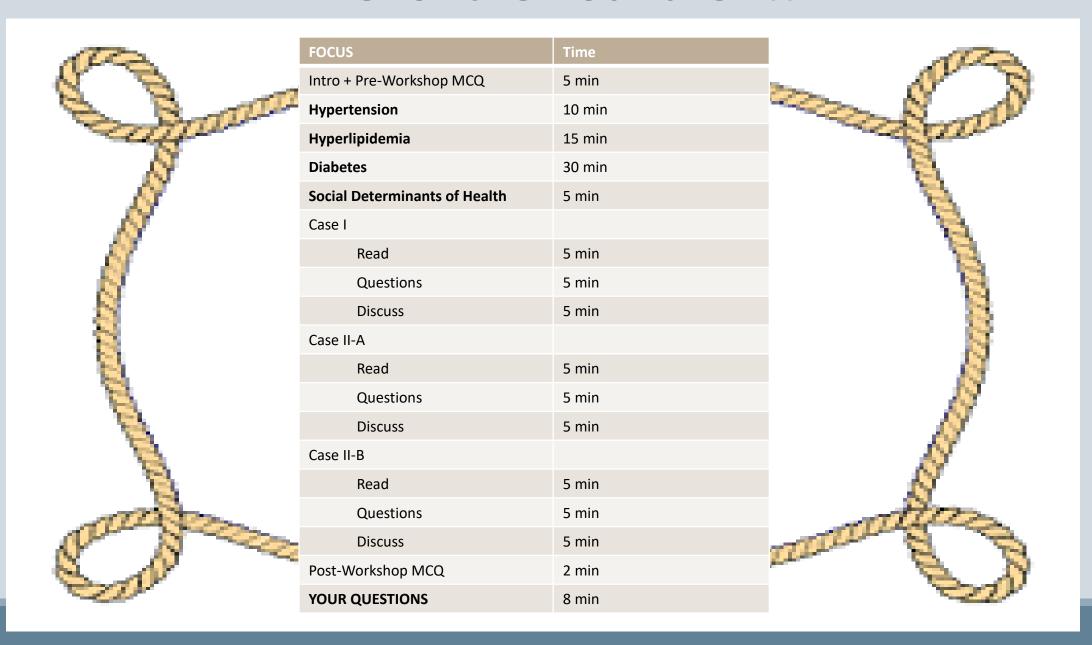
DEPARTMENT OF MEDICINE | SCHOOL OF HEALTH PROFESSIONS | PHYSICIAN ASSISTANT PROGRAM





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

Time for the Round-UP!!



OBJECTIVES

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

- Utilize recommended screening methodologies from the USPSTF and other authorities
- Apply current diagnostic guidelines
 for hypertension and atheroscierotic
 cardiovascular disease (including hyperlipidemia)
 from ACC/AHA and diabetes from ADA/AACE
- Implement individualized treatment plans using recommended algorithms for evidence-based pharmacotherapeutics and shared-decision approaches
- Counsel and motivate diverse patient populations on cardiovascular health 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

USPSTF RECOMMENDATIONS for HYPERTENSION (HTN) SCREENING		
Who?	All adults 18 +	
What?	Screen all adults for HTN (Grade A recommendation)	
Where?	 Screen in office, then confirm at home Educate BP should be taken at the brachial artery with an automated device in a seated position after 5 minutes of rest 	
When?	 Annual screening for adults 40+ Annual screening for adults at increased risk for HTN including: ✓ Black persons ✓ Persons with high-normal BP ✓ Persons who are overweight or obese Ok to screen less frequently (every 3-5 years) for adults 18 – 39 not at increased risk for HTN and with a prior normal BP reading 	

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.

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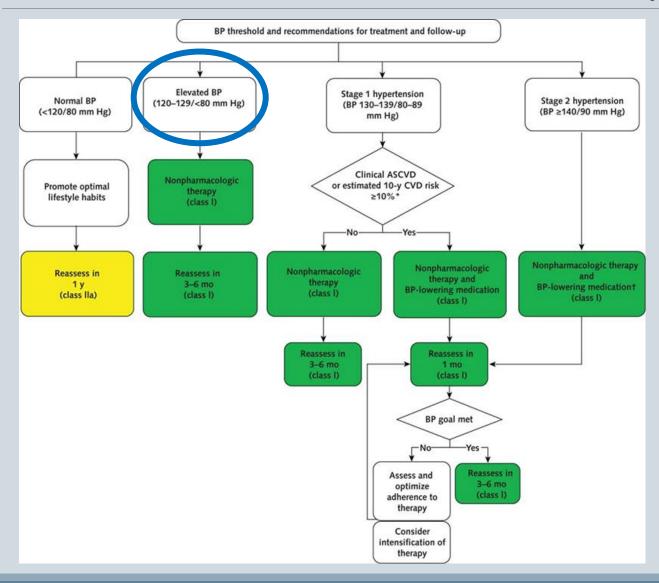
HYPERTENSION DIAGNOSTIC CRITERIA: ACC/AHA Guidelines

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

HYPERTENSION TREATMENT: ACC/AHA Guidelines



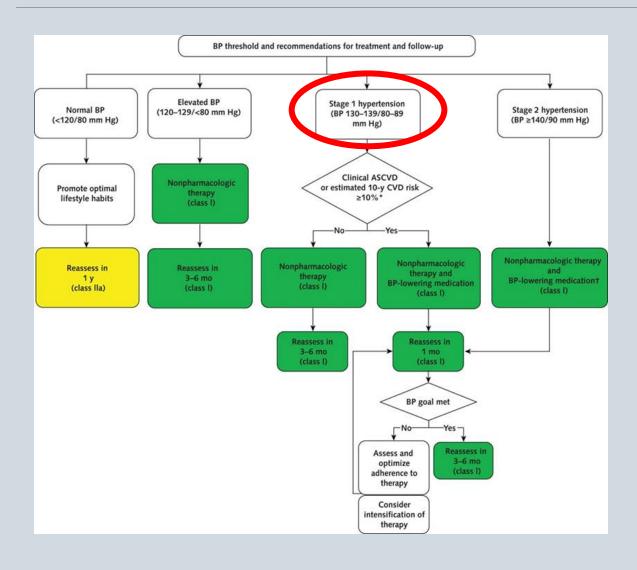
Elevated BP: 120-129/<80 mm Hg

LIFESTYLE MODIFICATIONS

HYPERTENSION TREATMENT: Lifestyle Modifications

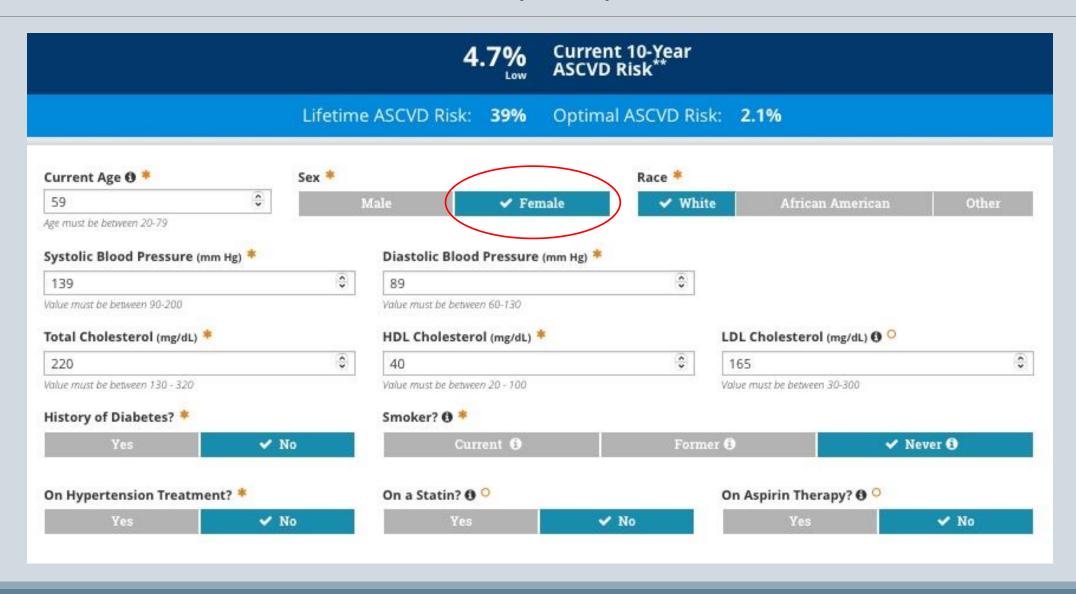
MODIFICATION	RECOMMENDATION	EXPECTED SBP REDUCTION
Weight Reduction	Maintain normal body weight (BMI 18.5-24.9)	5-20 mm Hg for each 10 kg weight loss
Adapt <u>DASH</u> eating plan	Consume diets rich in fruits, vegetables, low fat dairy and low saturated fat	8-14 mm Hg
Increase physical activity	Engage in regular aerobic activity such as walking (30 min/day on most days)	4-9 mm Hg
Dietary sodium reduction	Reduce sodium to no more than 1.5 g/day sodium	2-8 mm Hg
Moderate alcohol consumption	Limit alcohol to no more than 2 drinks/day for men and 1 drink/day for women.	2-4 mm Hg

HYPERTENSION TREATMENT: ACC/AHA Guidelines



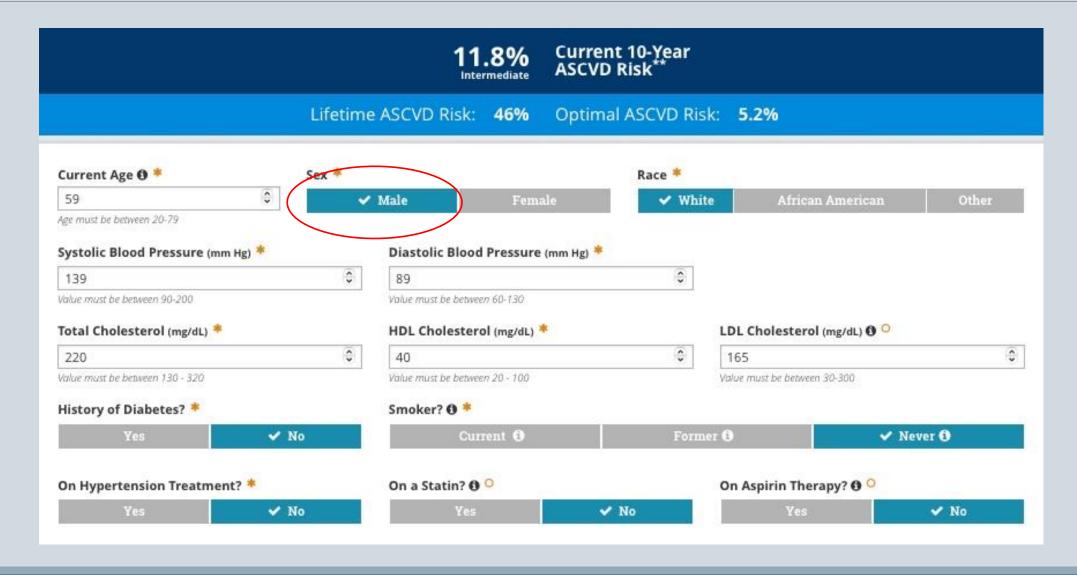
STAGE 1: 130-139/80-89 mm Hg

ASCVD RISK ESTIMATOR PLUS (ACC)



LIFESTYLE MODIFICATIONS!

ASCVD RISK ≥ 10%



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

LIFESTYLE MODIFICATIONS



ANTIHYPERTENSIVES

ASCVD RISK ≥ 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)

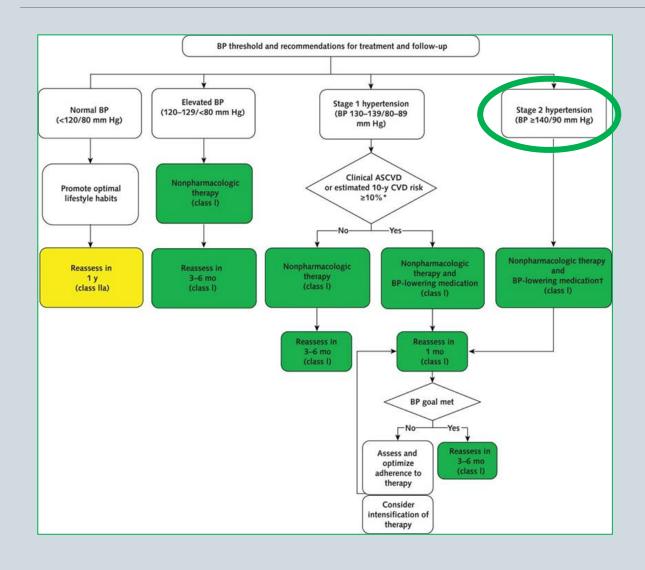
CALCIUM CHANNEL BLOCKERS

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

DIURETICS

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

HYPERTENSION TREATMENT: ACC/AHA Guidelines



STAGE 2: ≤140/90 mm Hg



FORGET THE CALCULATOR, MOVE STRAIGHT TO TREATMENT!

LIFESTYLE MODIFICATIONS



ANTIHYPERTENSIVES

STAGE 2 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)

SO, YOU STARTED TREATMENT, NOW WHAT?

REASSESS IN 1 MONTH

REASSESS AGAIN IN 3-6 MONTHS

IF **NOT AT** GOAL AFTER 1 MONTH, THEN

ASSESS & OPTIMIZE ADHERANCE TO THERAPY

+

CONSIDER INTENSIFYING THERAPY



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 en 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 Rish	The USPSTF recommends screening women age for lipid disor they are at increased risk for coronary heart disease.	В
Men 20-35 at Incre Risk for CHD	The USPSTF recommends screening men aged 20-35 for have a fitney are at sreased risk for coronary heart disease.	В
Men 20-35, Women Not at Increased Risk	men. A screening for lipid disorders in are not at increased risk for coronary heart dises.	C

ASCVD/Lipids Screening: USPSTF 2022 CU.S. Preventive Services



Steps for implementing screening:

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

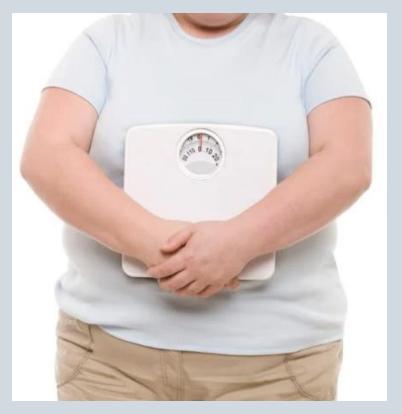
Population	Recommendation	Grade
Adults aged 40 to 75 years who have 1 or more cardiovascular risk factors and an estimated 10-year cardiovascular disease (CVD) risk of 10% or greater	The USPSTF recommends that clinicians prescribe a statin for the primary prevention of CVD for adults aged 40 to 75 years 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.	С
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.	l



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? American Resociation. AMERICAN COLLEGE of CARDIOLOGY.

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 pregnancy-associated conditions that increase ASCVD risk
- High-risk race/ethnicities
- Lipid/biomarkers





ACC ASCVD Risk Estimator Calculator for internet browser

AMERICAN COLLEGE of CARDIOLOGY ASCVD Risk Estin	nator Plus Estimate Ris	Sk Ø Therapy Impact Ø A
vananaanaanaanaanaanaanaanaanaanaanaanaa	••••	
Current Age • * Sex *	Race * Male Female Wi	nite African American Other
Age must be between 20-79 Systolic Blood Pressure (mm Hg) *	Diastolic Blood Pressure (mm Hg) *	7
Value must be between 90-200 Total Cholesterol (mg/dL) *	Value must be between 60-130 HDL Cholesterol (mg/dL) *	LDL Cholesterol (mg/dL) ᠪ ^O
Value must be between 130 - 320 History of Diabetes? *	Value must be between 20 - 100 Smoker? **	Value must be between 30-300
Yes No		ormer 1 Never 1
On Hypertension Treatment? * Yes No	On a Statin? 🚱 ^O Yes No	On Aspirin Therapy? 🐧 ^O Yes No

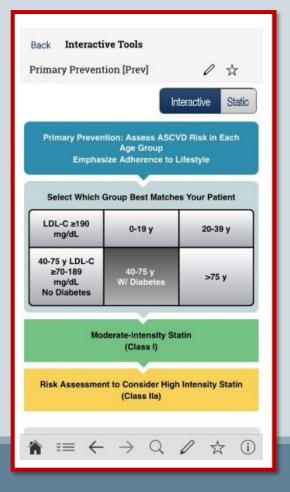


ACC Clinical Guideline Mobile App

COMPREHENSIVE MOBILE APP WITH ALL GUIDELINES AND CALCULATORS





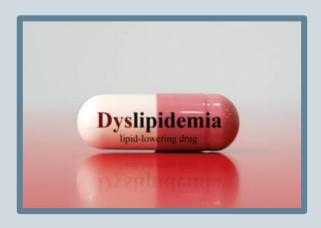


Statin Therapy by Intensity

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

ACC/AHA Treatment:



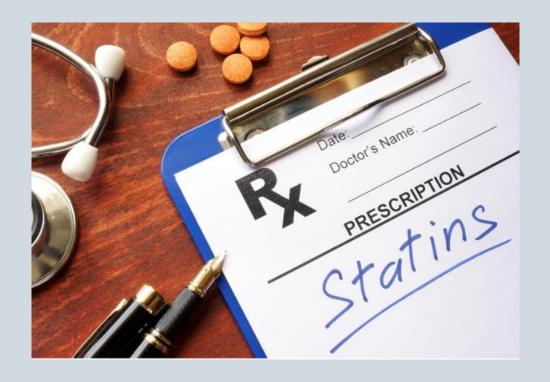
Severe Hypercholesterolemia LDL-C ≥ 190 mg/dL

Start treatment in 20 to 75 y/o with maximally tolerated **High Intensity*** STATIN therapy

Add EZETIMIBE if:

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

Add BILE ACID SEQUESTRANT if all the above and have fasting TG >300 mg/dL

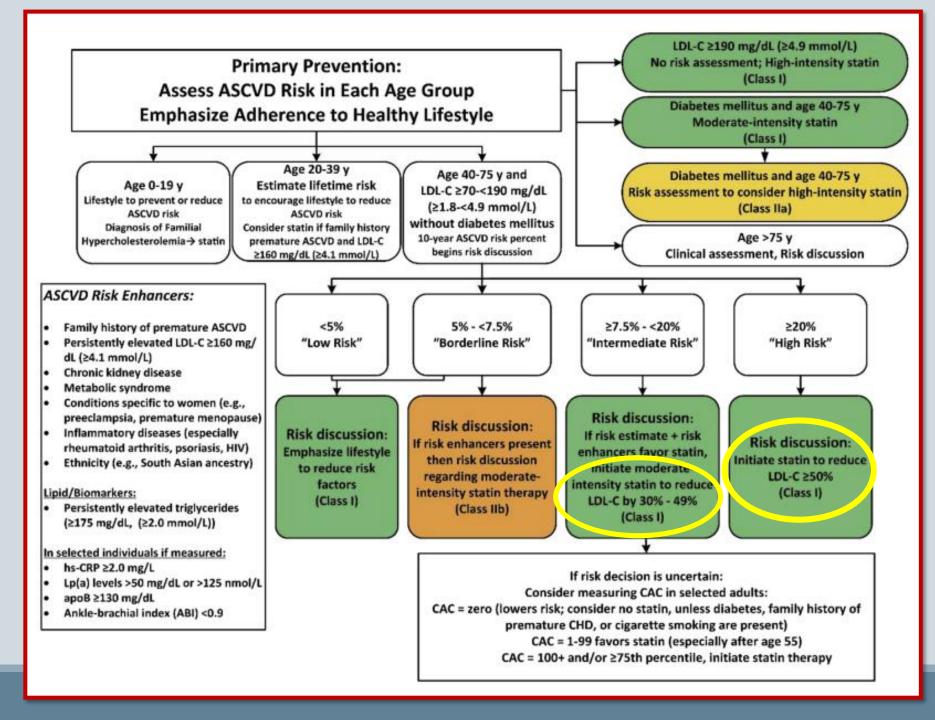


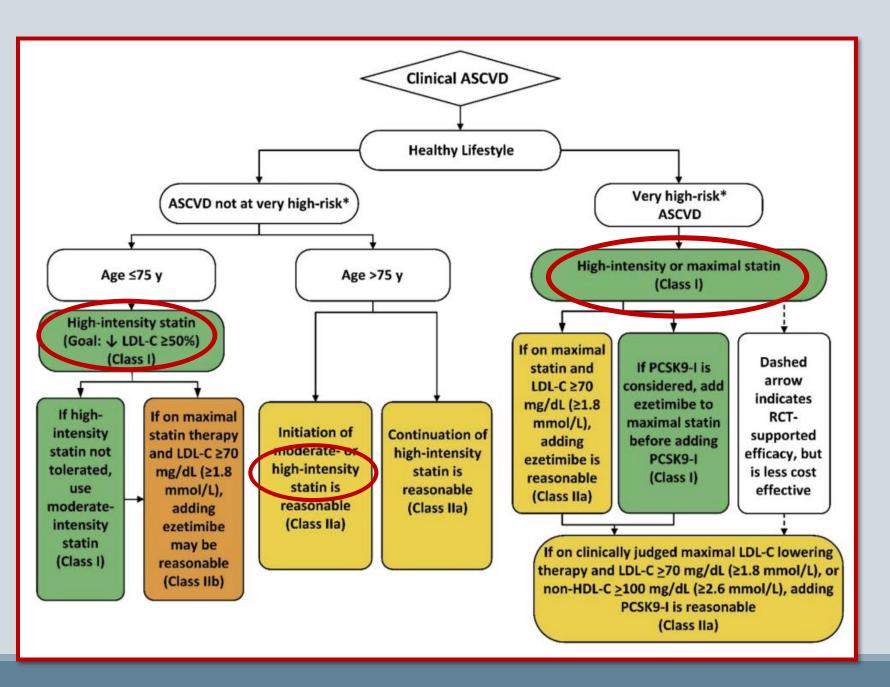
ACC/AHA Treatment:





ASCVD Primary Prevention





ACC/AHA



ASCVD Secondary Prevention

(ie-has had a cardiovascular event)



Quick word about Triglycerides from ACC

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
Severe (≥ 500 mg/dL, fasting) and ASCVD risk ≥ 7.5%	Start statin and address reversible causes of high TG

- AACE 2022 Update recommends adding fibrates or icosapent ethyl if fasting TG are persistently over > 200 mg/dL
- You can 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 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
- 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



Diabetes Screening: USPSTF CU.S. Preventive Services



Recommendation (Grade B):

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

KEY CHANGE: 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?	 Assess risk: 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. Screen: If the patient is aged 35 to 70 years and has overweight or obesity.
How often?	The optimal screening interval for adults with an initial normal glucose test result i uncertain. Screening every 3 years may 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.
statement?	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_{1c}, hemoglobin A_{1c}; 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

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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UpToDate°

DIABETES TARGET GOALS: HbA1c

Treatment Goals

ADA: < 7%

• 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%

GLYCEMIC CONTROL ALGORITHM

INDIVIDUALIZE GOALS

A1C ≤6.5%

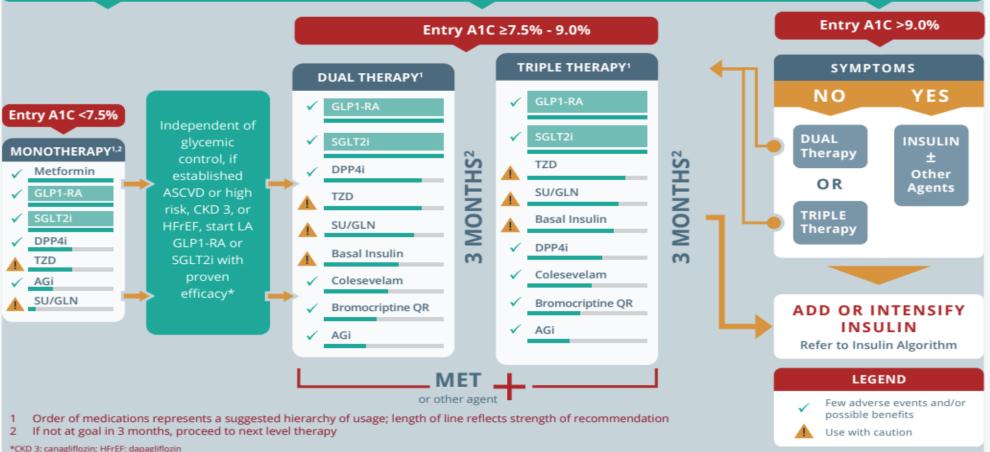
For patients without concurrent serious illness and at low hypoglycemic risk

A1C >6.5%

For patients with concurrent serious illness and at risk for hypoglycemia

LIFESTYLE THERAPY AND ONGOING GLUCOSE MONITORING (CGM preferred)

INDEPENDENT OF GLYCEMIC CONTROL, IF ESTABLISHED OR HIGH ASCVD RISK AND/OR CKD, RECOMMEND SGLT2i AND/OR LA GLP1-RA



(AA) CE







"CKD 3: canagimozin; HFTEF: dapagimozin

CKD 3 = stage 3 chronic kidney disease; HFTEF = heart failure with reduced ejection fraction; LA = long-acting (≥24 hour duration)



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†

+ASCVD/Indicators of High Risk

EITHER/

If A1C above target

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

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^

+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 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

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

---- OR --

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

efficacy to achieve goals:

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)

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

In people with HF, CKD, 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 recommendation 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 renal 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; # For 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.

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

If additional cardiorenal risk reduction or glycemic lowering needed

DIABETES TREATMENT:

(All Authorities Agree - ADA/ACE/AACE)

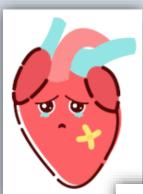
FIRST LINE TREATMENTS (EVERYONE!)

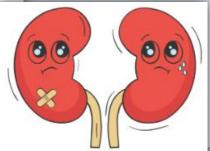
- Lifestyle modifications
 - Diet
 - Exercise
 - AHA recommends weekly goals of:
 - 150 mins of moderate intensity exercise
 - OR 75 mins of vigorous intensity exercise



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 m² 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 m²; 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

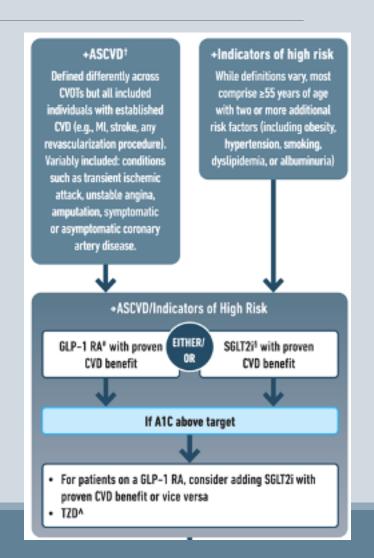


GLP-1 RA

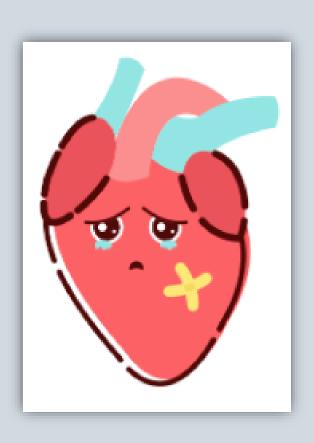
- Do NOT use in thyroid cancer, pancreatic cancer or MEN syndrome
- Side effects: GI issues such as nausea, reflux, diarrhea/constipation

SGLT2i

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

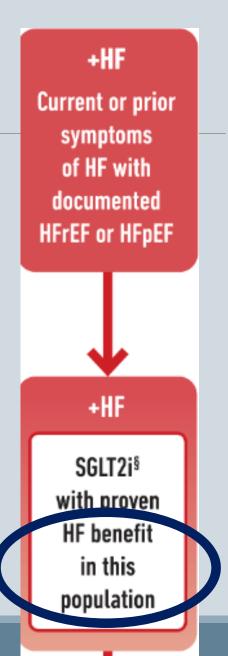


FIRST LINE THERAPEUTICS Heart Failure



Use SGLT2i

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



First Line Therapeutics: Chronic Kidney Disease (CKD)



CKD with aminuria

- SGL
- G RA if Se is not tolerated

CK vithout album. sia

- P-1 RA with CVD ben
- T2i with CVD benefit

+CKD**

CKD and albuminuria (e.g., ≥200 mg/g creatinine) CKD without albuminuria (e.g., eGFR <60 mL/min/1.73 m²)

PREFERABLY

T2i with primary evidence cing CKD progression

- OR

SGL revidence of reducing rogression in

GLP-1 RA with ven CVD benefit¹ if SGLT: tolerated or contrain ed

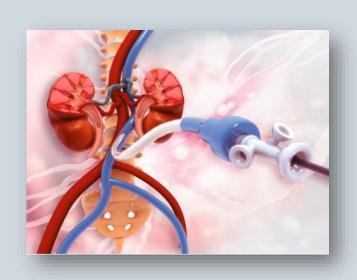
For patients with (e.g., eGFR <60 mL/min/1.7 without albuminuria, referenced ardiovascular

yith yith SGLT2i with proven CVD

benefit¹

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

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





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







Glycemic Management: Choose approaches that provide the 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 glucose 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-4

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

DIABETES TREATMENT: 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
 - Reduce dose between eGFR 30-45
- Recommended for prediabetes by ADA and AACE

DIABETES TREATMENT:What about Tirzepatide?

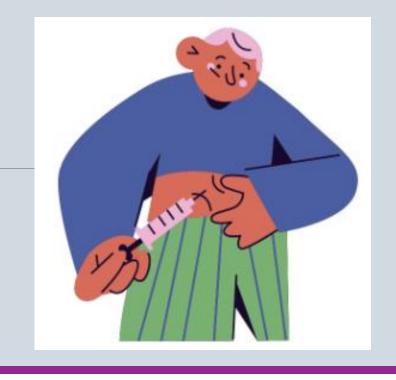
Classification

 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 Hypogly-		Weight change ²	CV effects		Renal effects	
	Efficacy cemia	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

DIABETES TREATMENT:Avoiding Hypoglycemia



Definition

- Decrease in glucose concentration <70 mg/dL
- 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

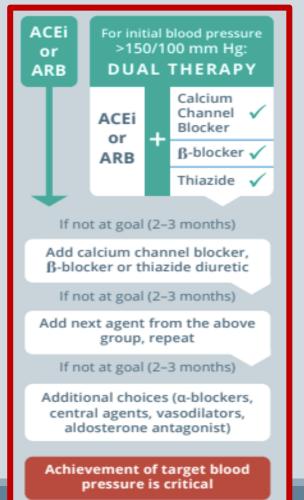
Sugar/honey

Hard candies

DIABETES CAVEATS: HYPERTENSION

LESS THAN 130/80

- ADA
 - 10-year ASCVD risk score at or greater than 15% or,
 - If established ASCVD
- AACE
 - Appropriate target for most individuals



LESS THAN 140/90

- ADA
 - 10-year ASCVD risk score *less* than 15%
 - **=AACE**
 - Aim for <130/80



DIABETES CAVEATS: ASCVD

AACE

Stratification to determine goals

HIGH: Diabetes alone and > 40 y/o

VERY HIGH: Diabetes + one more

risk factor

EXTREME: Diabetes + prior ASCVD event such as CVA or MI =or= CKD3/4

Treatment

- Maximally tolerated statin >
 Intensify statin to reach goal if tolerated
- Add ezetimibe or PCSK9inhibitors or colesevelam as needed

ADA

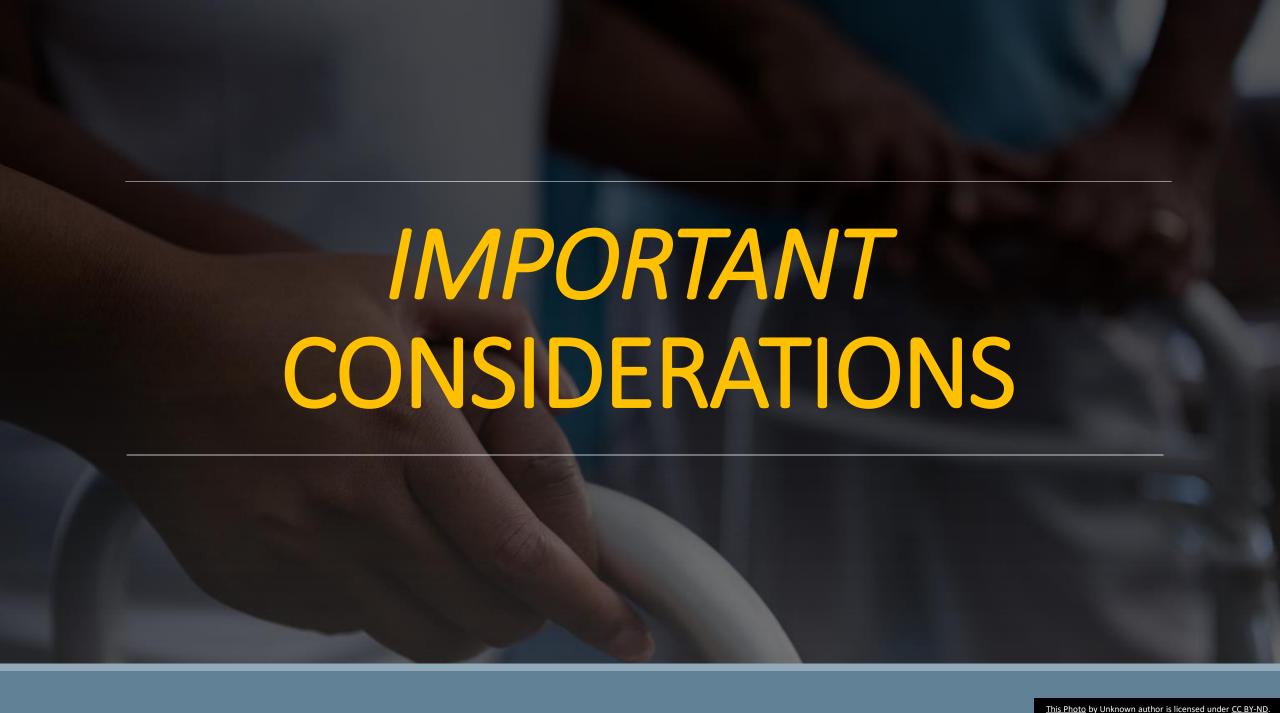
Primary Prevention

- Moderate-intensity statin in ages
 40-75 without ASCVD
- High intensity statin in patients with multiple ASCVD risk factors or ages 50-70
- Add ezetimibe in adults with ASCVD risk of ≥20%
- Reasonable to continue statin or initiate statin after cost/benefit discussion in pts
 >75 y/o

ACC/AHA

Primary Prevention

- Moderate-intensity statin in ages
 40-75 regardless of ASCVD risk
- High intensity statin in patients
 with multiple ASCVD risk factors
- Add ezetimibe in adults with ASCVD risk of ≥20%
- Reasonable to continue statin or initiate statin after cost/benefit discussion in pts >75 y/o



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







Case Content QR Code



CC: Headaches every morning with Claudia

Claudia is a 56-year-old female with prior GDM, controlled migraines and mild psoriatic arthritis.

She is presently concerned with a headache over the last 4 days that "comes and goes, usually only in the mornings." She reports the headaches are different from her migraines in that they are located across the forehead or on the back of her neck. She added that her headache yesterday caused some blurry vision, mild nausea and dizziness that were self-limiting. These headaches are not associated with vision loss, vomiting, photophobia or phonophobia. States no worsening factors but improvement either on their own or with acetaminophen. Denies fever, chills, unintentional weight loss, congestion, weakness, speech changes, numbness/tingling, chest pain, dyspnea, orthopnea, abdominal pain, bowel changes, dysuria or imbalance.

She had a similar symptoms a few months ago and was seen in the ER but was told "I was fine. They said I didn't have a stroke."

CC: Headaches every morning

PMH: As per above

PSH: Total hysterectomy without oophorectomy

Fam Hx: Father-unknown; Mother-HTN, thyroid disease. Brother-HTN, CAD and T1DM

Social Hx: Denies smoking vaping or chewing tobacco. Denies illicit drug use or

substance abuse. Drinks 1-2 glasses of red wine twice a week

Allergies: ciprofloxacin

Medications:

Daily: escitalopram 10mg, estradiol cream 0.1mg vaginal cream every 3 days, meloxicam 7.5mg, women's multivitamin.

PRN: Rizatriptan 5mg + Toradol 10mg for migraine

CC: Headaches every morning

Vitals

BP 138/89 | HR 86 and regular | Wt 203 lb | Ht 5'6" | BMI 32.8 | Waist Circumference 39" | Pulse ox 100%

Physical Exam

GEN: no acute distress; alert and oriented x 3

Skin: warm, dry skin without tenting; no lesions or rashes

HN: Normocephalic, atraumatic; neck supple without rigidity and without carotid bruits; no thyromegaly

EENT: Conjunctiva clear; PERRLA; funduscopic exam with discs flat and sharp; OP with pink moist buccal mucosa; Nasal turbinates pink and moist

CV: Distinct S1, S2; no murmurs, gallops or rubs

Lungs: CTA-B without wheezing, rales or rhonchi

Neuro: Normal speech; cranial nerves II through XII intact; DTR's symmetric; Babinski sign negative; motor and sensory grossly normal bilaterally; normal muscle tone, no tremors, BUE and BLE strength 5/5, Romberg sign negative, normal gait and station.

CC: Headaches every morning

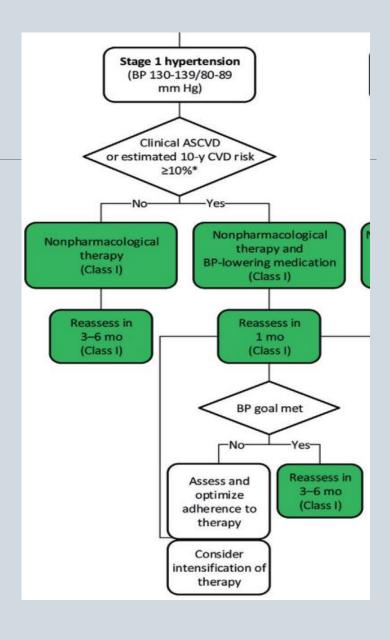
Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value
Glucose	122	Protein	6.9	TSH	3.5	Chol	260
BUN	19	Albumin	4.1	HgA1c	5.5	HDL	29
Creatinine	0.99	Globulin	2.3	Vitamin D	18	LDL-C	170
eGFR	76	A/G Ratio	1.8			TG	306
Sodium	145	Bilirubin, total	0.3	WBC	8.1		
Potassium	3.8	AST	28	RBC	4.1		
Chloride	105	ALT	35	Hg	12.6		
Carbon dioxide	25			Hct	39.7		
Calcium	9.9						

Lifetime ASCVD Risk: 39% Optimal ASCVD Risk: 2.0%

Current Age 😝 *	Sex *		Race *		
56	Male	✓ Female	White	✓ African	American Other
Age must be between 20-79					
Systolic Blood Pressure (mm Hg) *	Diastolic Blo	od Pressure (mm Hg) *			
138	89				
Value must be between 90-200	Value must be betw	veen 60-130			
Total Cholesterol (mg/dL) *	HDL Cholesto	erol (mg/dL) *		LDL Cholesterol (mg	g/dL) 🚯 🖰
260	29			170	
Value must be between 130 - 320	Value must be betw	veen 20 - 100		Value must be between 30-3	00
History of Diabetes? *	Smoker? 🛭 *	k			
Yes 🗸 N	To (Current 19	Forme	er 🔁	✓ Never ①
On Hypertension Treatment? *	On a Statin?	0 °		On Aspirin Therapy	r? 0 °
Yes 🗸 N	To Yo	es 🗸	No	Yes	✓ No

Discussion questions

- 1. What are the diagnoses?
- 2. How would you approach the treatment for HTN?
- 3. Does this patient have ASCVD?







CC: Follow up on back pain and labs with Mario

Mario is 46-years-old with OSA and HTN, who presents to clinic after going to the ED for back pain. His random blood glucose was 133 and he was advised to follow up with his PCP.

He reports trying diet and exercise to control his glucose since his last physical but finds it hard to make time to exercise.

Currently, he denies polydipsia, polyphagia and polyuria as well chest pain, shortness of breath, palpitations, headache, fevers, unintentional weight loss, abdominal pain, diarrhea, or rashes.

CC: Follow up on back pain and labs

PMH: As per above

PSH: Tonsillectomy as a child

Fam Hx: Father-ASCVD; died at 62 of an AMI; Mother-diabetes and

breast cancer; sister-RA and HTN

Social Hx: Previous 22-pack year history. Quit smoking tobacco two years ago; does not vape or chew tobacco. Denies illicit drug use or alcohol use

Allergies: NKDA

Medications: Daily: loratadine 10mg, HCTZ 25mg, vitamin D 2000

IU, aspirin 81mg. PRN ibuprofen 600mg.

CC: Follow up on back pain and labs

Vitals

BP 143/79 | HR 86 | Wt 223 lb | Ht 5'9" | BMI 32.9 | Waist Circumference 45" | Pulse ox 100% RA

Physical Exam

GEN: no acute distress; alert and oriented x 3

Skin: dry, skin tenting noted

HEENT: conjunctiva clear, PERRLA; OP with pink, moist mucosa; Nasal turbinates pink, moist

CV: RRR with normal S1/S2; no murmur or gallops noted

Lungs: CTA-B without wheezing, rales or rhonchi

Abd: soft, non-distended, non-tender with normal bowel sounds, no palpable masses, no hepatosplenomegaly, no rebound or guarding, no CVA-T

CC: Follow up on back pain and labs

Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value
Glucose	118	Protein	7.2	TSH	2.6	Chol	256
BUN	22	Albumin	4.6	HgA1c	6.3	HDL	31
Creatinine	1.00	Globulin	2.6	Vitamin D	27	LDL-C	169
eGFR	87	A/G Ratio	1.77			TG	282
Sodium	138	Bilirubin, total	0.9	WBC	5.3		
Potassium	4.2	AST	26	RBC	4.1		
Chloride	99	ALT	32	Hg	13.1		
Carbon dioxide	26			Hct	39.2		
Calcium	8.9						

Lifetime ASCVD Risk: 69% Optimal ASCVD Risk: 1.3%

App should be used for primary prevention patients (those without ASCVD) only.

Current Age 🛭 *	ex *		Race *	_		
46	✓ Male	Female	✓ White	African American	Other	
Age must be between 20-79						
Systolic Blood Pressure (mm Hg) *	Diastolic Blood	Pressure (mm Hg) *				
143	79					
Value must be between 90-200	Value must be between	n 60-130				
Total Cholesterol (mg/dL) *	HDL Cholester	ol (mg/dL) *	L	DL Cholesterol (mg/dL) ᠪ ^O		
256	31			169		
Value must be between 130 - 320	Value must be between	n 20 - 100	V	Value must be between 30-300		
History of Diabetes? *	Smoker? 🛭 *		F	low long ago did patient quit s	moking? *	
Yes 🗸 No	Current 1	✓ Former 🛈 N	lever 🛈	2.5-3.5 years ago		
On Hypertension Treatment? *	On a Statin? 🚯	0	c	n Aspirin Therapy? 🛭 O		
✓ Yes No	Yes	✓	No	✓ Yes	No	

Discussion questions



- 1. What are the diagnoses?
- 2. Does this patient have ASCVD? If so, how would you address this diagnosis?
- 3. What are the implications of this pt's BMI? How can this be addressed?
- 4. What are the implications of the pt's glucose? How could this be addressed?
- 5. Does this pt need to be on aspirin?

Six-month follow-up with Mario

On routine follow-up, Mario reports he has changed his lifestyle by drinking fewer soft drinks and but still eats out for lunch 3-4 times a week. He has not had time to exercise. He is compliant with his **new medications** *



Medications

Daily: loratadine 10mg, HCTZ 25mg, Vitamin D 2000 IU, valsartan 160mg*, atorvastatin 20mg*

PRN: ibuprofen 600mg, famotidine 10mg

Six-month follow-up with Mario

Vitals

BP 122/78 | HR 74 | Wt 218 lb | Ht 5'9" | BMI 32.2 | Waist Circumference 45" | Pulse ox 100%

Physical Exam

GEN: no acute distress, alert and oriented x 3; obese

Skin: dry, skin tenting noted

HEENT: conjunctiva with PERRLA; pink oral mucosa, dry; Nasal turbinates pink, moist

CV: RRR with normal S1/S2; no murmur or gallops noted

Lungs: CTA-B without wheezing, rales or rhonchi

Abd: soft, non-distended, non-tender with normal bowel sounds, no palpable masses, no hepatosplenomegaly, no rebound or guarding, no CVAT

Six-month follow-up with Mario

Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value	Fasting Lab	Value
Glucose	145	Protein	7.5	TSH	3.2	Chol	209
BUN	23	Albumin	4.3	HgA1c	7.2	HDL	41
Creatinine	1.04	Globulin	2.7			LDL-C	123
eGFR	87	A/G Ratio	1.59	WBC	4.8	TG	223
Sodium	136	Bilirubin, total	0.9	RBC	3.9		
Potassium	3.9	AST	36	Hg	13.7		
Chloride	6	ALT	41	Hct	38.5		
Carbon dioxide	24						
Calcium	8.3						

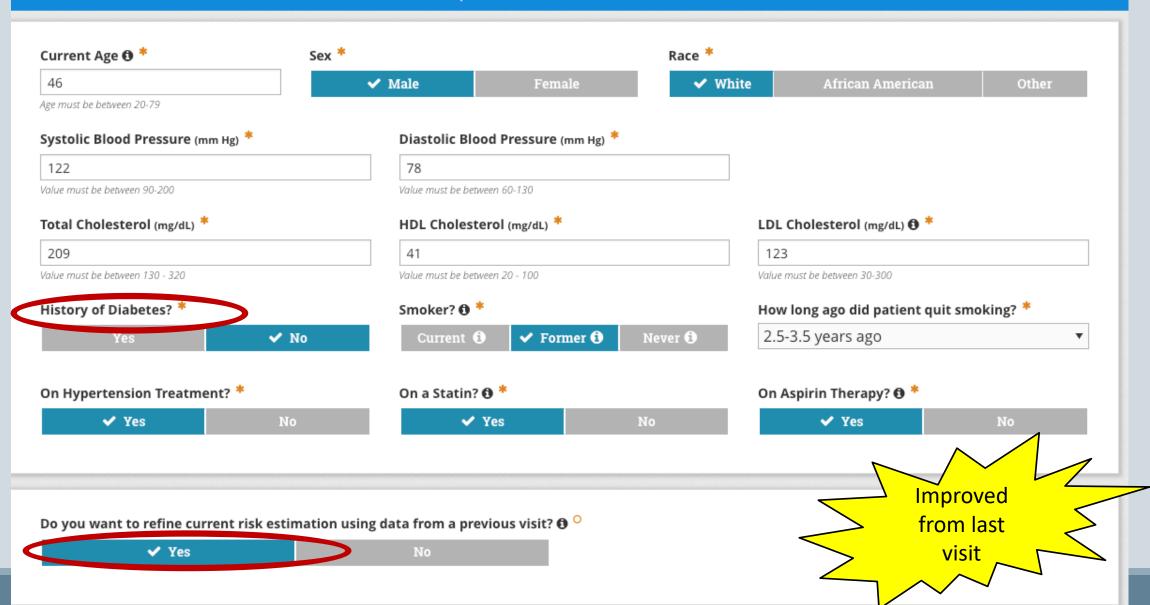
Lifetime ASCVD Risk: 50%

Optimal ASCVD Risk: 1.3%

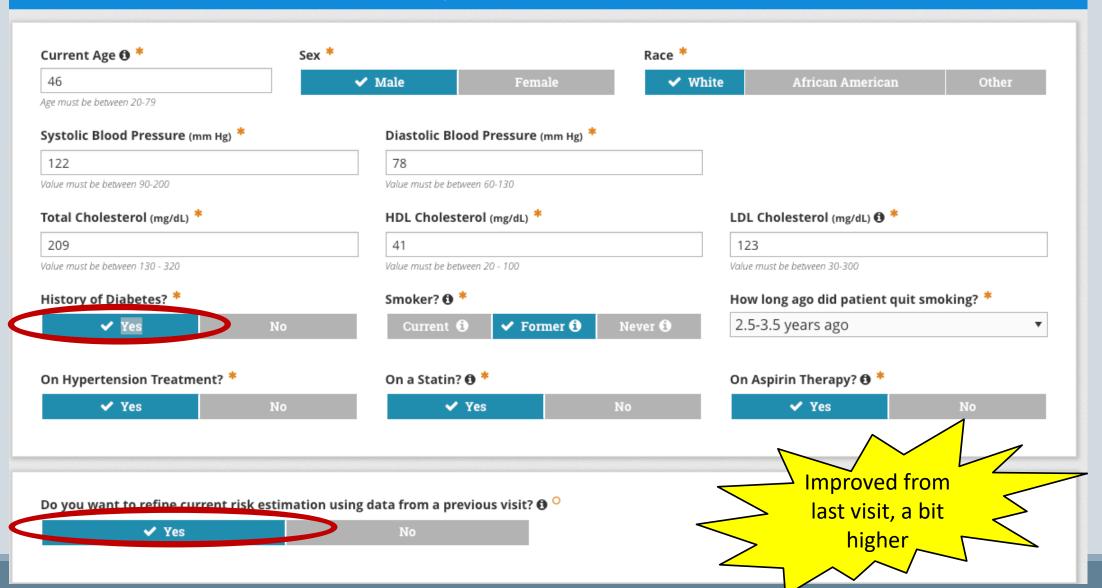
Current Age ᠪ *	Sex *			Race *		
46		✓ Male	Female	✓ White	African American	Other
Age must be between 20-79						
Systolic Blood Pressure (mm Hg) *		Diastolic Blo	od Pressure (mm Hg) *			
122		78				
Value must be between 90-200		Value must be betv	veen 60-130			
Total Cholesterol (mg/dL) *		HDL Cholest	erol (mg/dL) *		LDL Cholesterol (mg/dL) ᠪ ^O	
209	\$	41			123	
Value must be between 130 - 320		Value must be betv	veen 20 - 100		Value must be between 30-300	
History of Diabetes? *		Smoker? 🗗 🦥	•		How long ago did patient quit s	moking? *
Yes	✓ No	Current C	✓ Former ①	Never 🚯	2.5-3.5 years ago	•
On Hypertension Treatment? *		On a Statin?	6 °		On Aspirin Therapy? 🛭 ^O	
✓ Yes	No		Yes	No	✓ Yes	No

Do you want to refine current risk estimation using data from a previous visit? 19 O

Optimal ASCVD Risk: 1.3%



Optimal ASCVD Risk: 1.3%



Discussion questions



- 1. What are the diagnoses?
- 2. What are your impressions of his ASCVD risk?
- 3. What are the implications of the new glucose readings?
- 4. Discuss the implications of the lipid panel after starting statin therapy? How do we approach the elevated TG?
- 5. Does he need to be on aspirin?

TAKE HOME POINTS

- ASCVD Risk Estimation drives diagnoses and management of hypertension, hyperlipidemia and diabetes.
- Lifestyle modifications are still the mainstay in treatment.
- Statin therapy is the preferred treatment for hypercholesteremia and hypertriglyceridemia.
- Novel treatments for diabetes that reduce the risks of ASCVD, HF and CKD and that promote weight loss are considered first-line therapy.
- Meet your patients where they are based on SDOH and find creative solutions to help them meet goals.

7

Thank you!

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References and Acknowledgments

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ALGORITHM FOR ADDING/INTENSIFYING INSULIN

Glycemic

Control Not

at Goal*

START BASAL (Long-Acting Insulin)

A1C <8%

A1C >8%

TDD 0.1-0.2 U/kg

TDD 0.2-0.3 U/kg

Insulin titration every 2-3 days to reach glycemic goal:

- · Fixed regimen: Increase TDD by 2 U
- Adjustable regimen:
- FBG >180 mg/dL: add 20% of TDD
- FBG 140-180 mg/dL: add 10% of TDD
- FBG 110-139 mg/dL; add 1 unit
- · If hypoglycemia, reduce TDD by:
- BG <70 mg/dL: 10% 20%
- BG <40 mg/dL: 20% 40%

Consider discontinuing or reducing sulfonylurea after starting basal insulin (basal analogs preferred to NPH)

*Glycemic Goal:

- <7% for most patients with T2D; fasting and premeal BG <110 mg/dL; absence of hypoglycemia
- A1C and FBG targets may be adjusted based on patient's age, duration of diabetes, presence of comorbidities, diabetic complications, and hypoglycemia risk

INTENSIFY (Prandial Control)

Add GLP1-RA

Or SGLT2i Or DPP4i

Add Prandial Insulin



Basal Plus 1, Plus 2, Plus 3

- Begin prandial insulin before largest meal
- · If not at goal, progress to injections before 2 or 3 meals
- Start: 10% of basal dose or 5 units

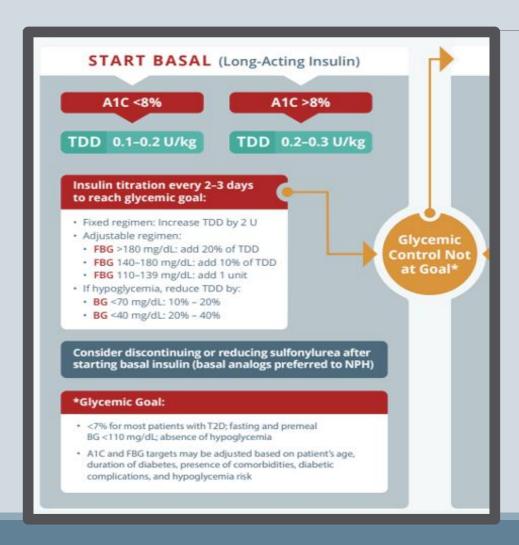
Basal Bolus

- Begin prandial insulin before each meal
- 50% Basal / 50% Prandial TDD 0.3-0.5 U/kg
- Start: 50% of TDD in three doses before meals

Insulin titration every 2-3 days to reach glycemic goal:

- . Increase prandial dose by 10% or 1-2 units if 2-h postprandial or next premeal glucose consistently >140 mg/dL
- · If hypoglycemia, reduce TDD basal and/or prandial insulin by:
- BG consistently <70 mg/dL: 10% 20%
- · Severe hypoglycemia (requiring assistance from another person) or BG <40 mg/dL: 20% - 40%

STARTING INSULIN



BASAL

- Lower incidence of hypoglycemia
- Glargine
 - U-100: Lantus or Basaglar
 - U-300: Toujeo concentrated
- Detemir: Levemir, usually BID
- Degludec (Tresiba)
 - U-100
 - U-200 concentrated , lasts 40+hrs = steady state
- Titration
 - Fixed: increase by 2U if glucose over goal for 2-3 days straight
 - Sliding: increase insulin dose by10%-20% of the TDD if over 180 or 140 respectively
 - Decrease insulin if hypoglycemia is noted or FBG<140 x 3 days

BOLUS (aka: prandial, mealtime)

- Timing is KEY to prevent hypoglycemia.
- Administer 20-30 mins before meal
- Eat before the peak

Basal + Bolus with the Biggest meal (BBB)

- Fewer sticks a day (ie: insulins + glucose monitoring)
- May help patients get past fear of needles

Start both Basal + Bolus together

- Give half of the TDD in Basal dose at night
- Give the other half in Bolus with meals
- SLE pt 120kg pt @ 0.3U/kg = 36 Units TDD
 - Basal: 18U at night
 - Bolus: 6U x 3 meals for remainder18U



Add GLP1-RA

Or SGLT2i Or DPP4i

Glycemic

Control Not

at Goal*

Add Prandial Insulin

Basal Plus 1, Plus 2, Plus 3

 Begin prandial insulin before largest meal

- If not at goal, progress to injections before 2 or 3 meals
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Insulin Pearls

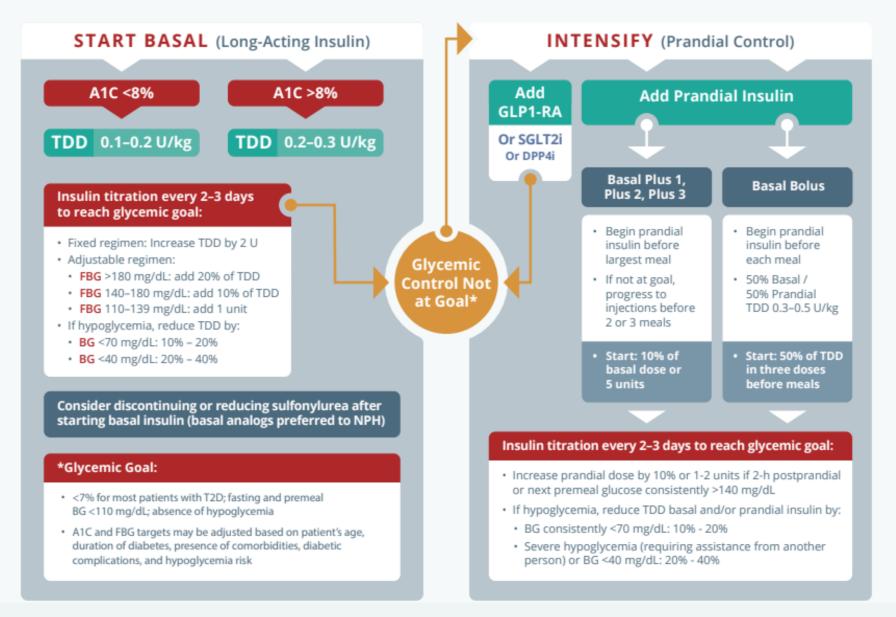
AACE

- If insulin is to be used, start with a BASAL insulin
- Basal insulin (or GLP1-RA) should be considered if a patient is on two oral agents and has an A1c >8.0%

ADA

- Start insulin early if the patients has weight loss, hyperglycemia symptoms, or when A1c or glucose are high (>10% or >300mg/dL respectively)
- Continue metformin if insulin is started (unless there are contraindications)

ALGORITHM FOR ADDING/INTENSIFYING INSULIN



Blood Glucose Targets Goals (for INSULIN)

in Nonpregnant Adults With Diabetes

Pre-prandial Blood Glucose

80 - 130 mg/dL

- •Variable based on comorbidities, age, risk of hypoglycemia
- •Readings used to determine bolus insulin dose

Post-prandial Blood Glucose

<180 mg/dL





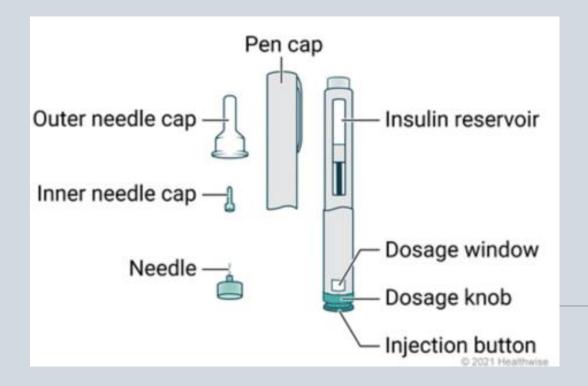


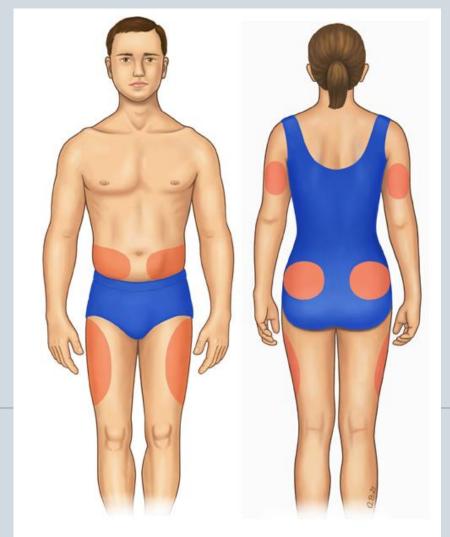




CbGM:

Continuous
Blood Glucose
Monitor





The shaded areas may be used for insulin injections. Injection sites should be rotated. Insulin is absorbed more rapidly when injected into the abdomen as compared with the arms or legs.