IT'S MORETHAN NUMBERS ON A GLUCOMETER

CURRENT PRIORITIES OF DIABETES MANAGEMENT



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DISCLOSURES

- ➤ Novo Nordisk: Advisor/Speaker (obesity, T2DM);
- ➤ Acella: Speaker: Hypothyroidism, desiccated thyroid extract (DTE)

All financial disclosures have been mitigated

OBJECTIVES

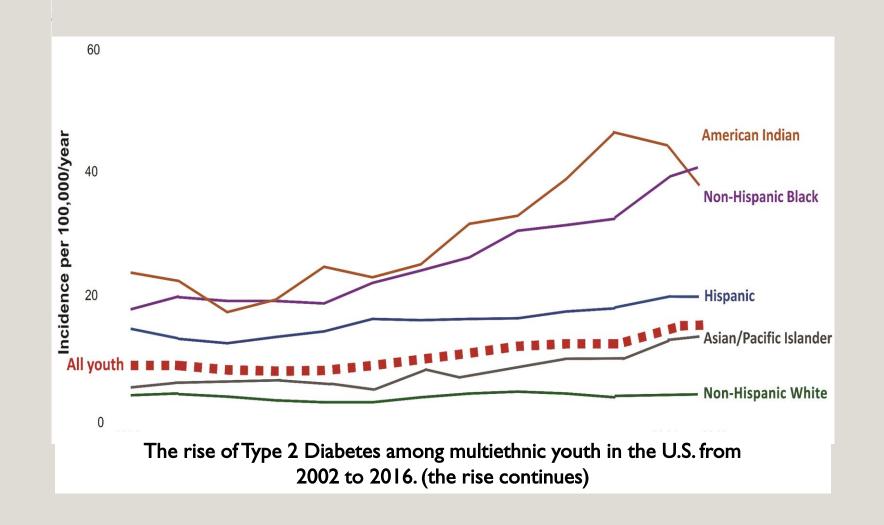
- List the FIVE categories of diabetes and clues to their presentation
- Identify those at risk for diabetes and/or insulin resistance
- Review the pathogenesis of Type 2 diabetes including at least 8 contributing metabolic impairments leading to dysglycemia
- Discuss laboratory findings in diagnosing prediabetes, Type 1 Diabetes and Type 2 diabetes
- Discuss how to obtain useful fingerstick blood glucose data
- Review risks and treatment of hypoglycemia
- Identify underlying neuropathy, retinopathy, and nephropathy using appropriate assessment tools

THE DIABETES EPIDEMIC THE SWEETEST PEOPLE IN THE WORLD

DIABETES

- Total: 37.3 million (11.3%) of the US population have diabetes
- Diagnosed: 28.7 million
- Undiagnosed: 8.5 million people
- PREDIABETES
- Total: 96 million (38.0%) adults (over 18 yrs) have prediabetes
- 65 years or older: 26.4 million (48.8%) have prediabetes!
- 1 in 3 people estimated to have diabetes by 2050
- Every 8 seconds someone dies from something related to

Youth-onset T2DM





Expected 400% increase within next 25 years

What types of diabetes are there?

- Autoimmune Type 1 (leads to absolute insulin deficiency)
 - Type 1 diabetes (more rapid development)
 - Latent autoimmune diabetes in adults (LADA) (slow development)
 - An "event" will trigger the onset
- Type 2 diabetes (most common) (non-autoimmune)
 - progressive loss of β-cell function & relative insulin deficiency
 - Associated with IR and metabolic syndrome (now in younger pts)

Gestational diabetes

Found in 2nd or 3rd trimester (unmasks underlying DM presence or risk)

MODY

- 4 main types (1-2% of those with DM—usually misdiagnosed)
- HNF1-alpha; HNF1-beta; HNF4-alpha; glucokinase
- Exocrine pancreas disease/impairment
 - Pancreatitis, cystic fibrose, adiposity
 - Drug-induced

 glucocorticoid use

Suspicions based on:

Family History (genes)

Age

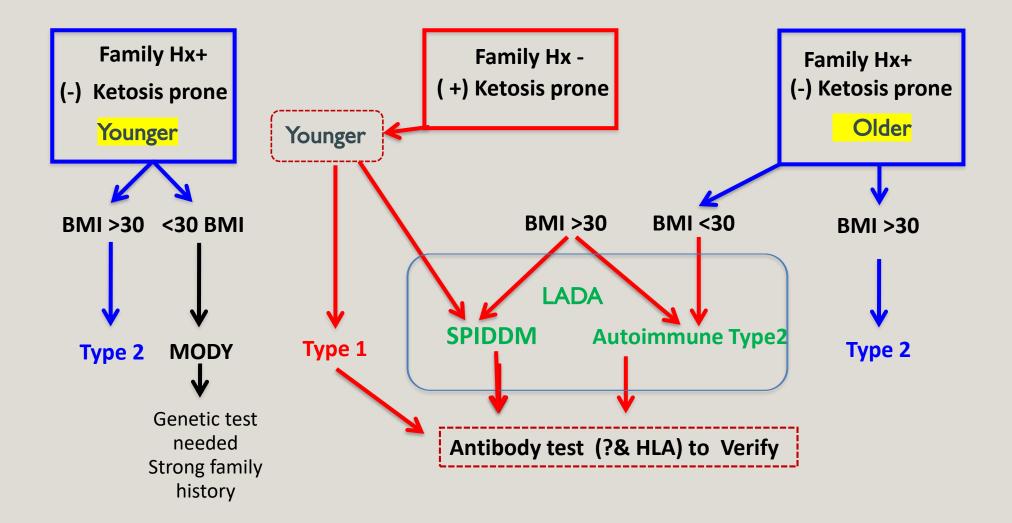
Ketosis prone

BMI

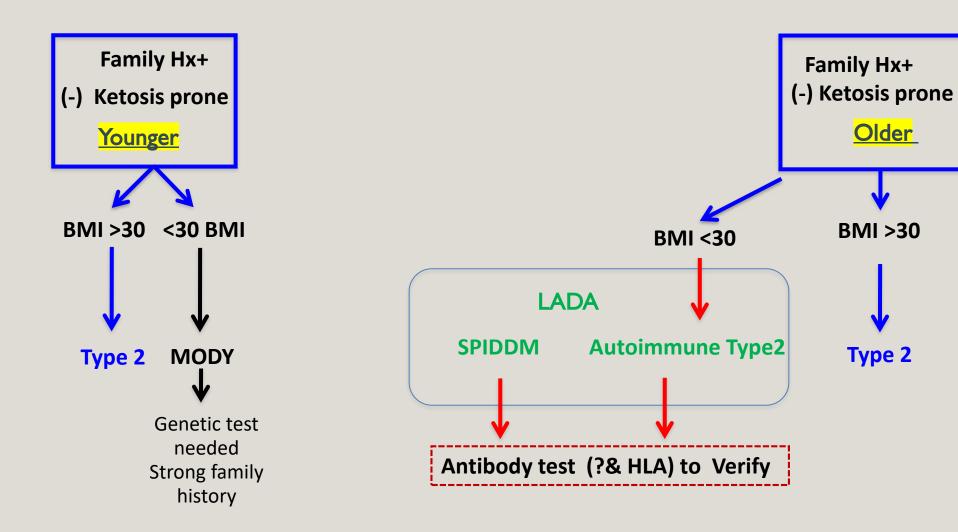
Antibodies

Greater risk in those with underlying or family history of autoimmune disease

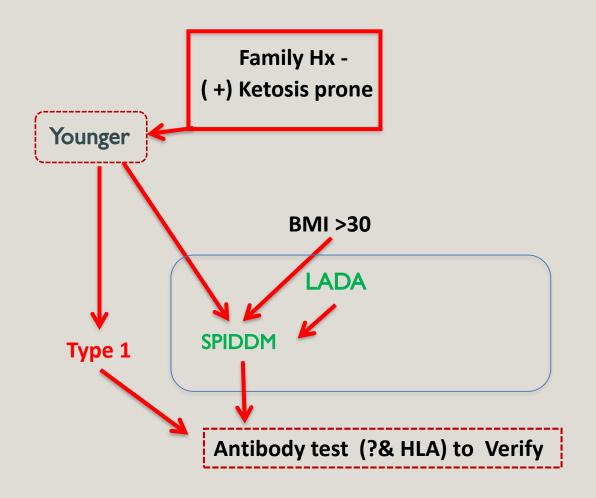
TIPS TO TYPES OF DIABETES



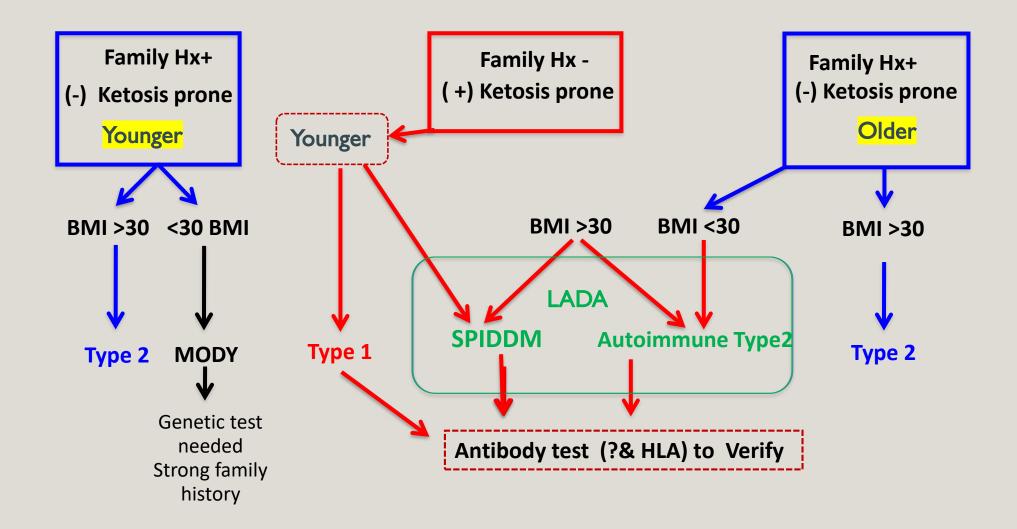
TIPS TO TYPE OF DIABETES



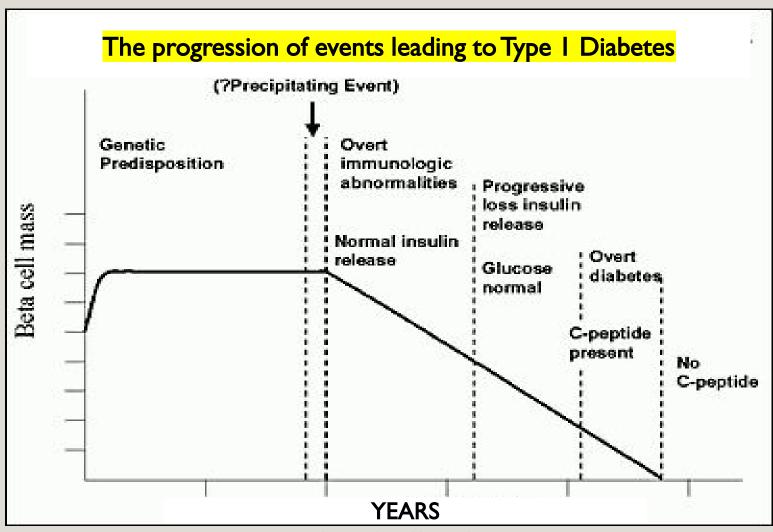
TIPS TO S OF DIABETES



TIPS TO TYPES OF DIABETES



Genetics-Environment-Biology



Examples of Triggers:

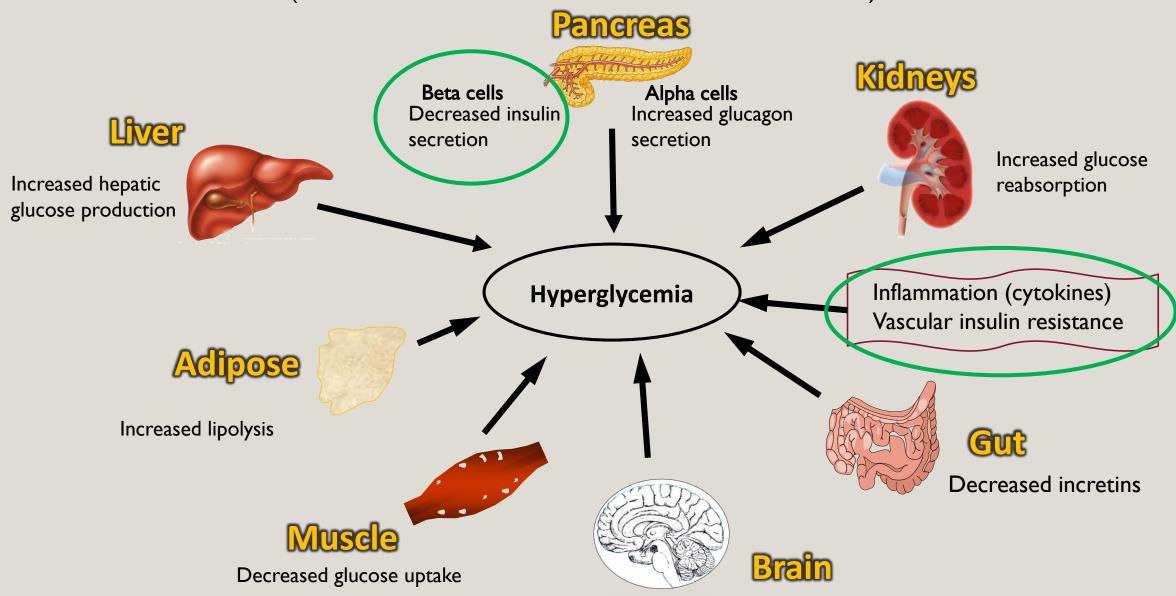
Endocrine disrupting chemical
Virus (esp norovirus,)
Certain foods (grains, bovine milk, etc
Vax reactions (yep)
Stress
PTSD
more

Image adapted by speaker

Type I Diabetes Stages

	Stage 1	Stage 2	Stage 3
	Autoimmunity	Autoimmunity	Autoimmunity
Characteristics	Normoglycemia	Dysglycemia	Overt hyperglycemia
	Presymptomatic	Presymptomatic	Symptomatic
Diagnostic criteria etes Care 2023;46(Supplement etes 2017;66:241–255 Open 2019;9:e031586	 Multiple islet autoantibodies No IGT or IFG 1):S19-S40 	 Islet autoantibodies (usually multiple) Dysglycemia: IFG and/or IGT FPG 100–125 mg/dL 2-h PG 140–199 mg/dL A1C 5.7–6.4% or ≥10% increase in A1C 	 Autoantibodies may become absent Diabetes by standard criteria

PATHOGENESIS OF DIABETES (INFLAMMATION & INSULIN RESISTANCE)



Neurotransmitter dysfunction

BETA CELL FAILURE & INSULIN REDUCTION IN T2DM



WHO'S AT GREATER RISK FOR DIABETES?

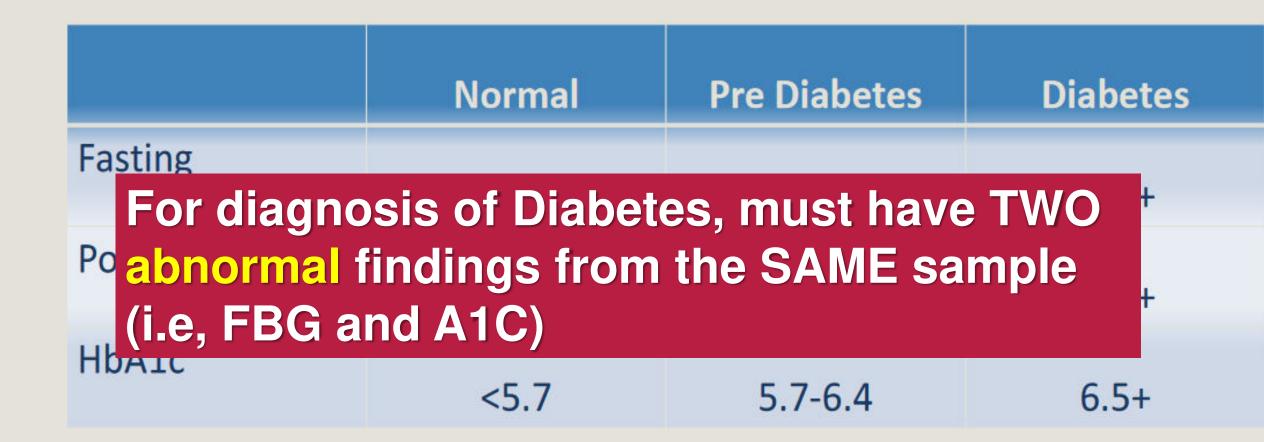
- Family history
- Hx of Autoimmune disease (139 of these)
- Age
- Gender or ethnic influences (African American, Latino, Native American, Asian American, Pacific Islander)
- Gestational diabetes
- Hx of carpal tunnel or Duprey's contracture?
- Morbid obesity
- New onset hyperglycemia in acute care setting?

Diagnosing diabetes

	Normal	Pre Diabetes	Diabetes
Fasting	< 100	100-125	126+
Postprandial	<140	140-199	200+
HbA1c	<5.7	5.7-6.4	6.5+

ElSayed, N. A., Aleppo, G., Aroda, V. R., Bannuru, R. R., Brown, F. M., Bruemmer, D., ... & Gabbay, R. A. (2023). 2. Classification and Diagnosis of Diabetes: Standards of Care in Diabetes—2023. *Diabetes Care*, 46(Supplement 1), S19-S40

Diagnosing diabetes



ElSayed, N. A., Aleppo, G., Aroda, V. R., Bannuru, R. R., Brown, F. M., Bruemmer, D., ... & Gabbay, R. A. (2023). 2. Classification and Diagnosis of Diabetes: Standards of Care in Diabetes—2023. *Diabetes Care*, 46(Supplement 1), S19-S40

POINTS ABOUT THE A I C

- Reflects AVERAGE blood glucose over a 3 month period
- It does not reflect blood glucose variations
- Its accuracy can be affected by the following:

False Increase:

Anemias (iron/Vit B12 deficiency)

Asplenia

Uremia

High triglycerides (>1750 mg/dl)

Chronic ETOH use

Chronic salicylate use

Chronic opioid use

HGB variants

Blood transfusion (high sugar in storage medium)

Vitamin C ingestion (depends on lab methods)

False Decrease:

Anemia (blood loss)

Splenomegaly

Pregnancy (1st & 2nd trimester)

Ribavirin & interferon-alpha

Vitamin E use

Blood transfusion (dilution effect)

Vitamin C ingestion (depends on lab methods)

Are there other options?

MOST COMMON ANTIBODY TESTS FOR AUTOIMMUNE DIABETES

- Positive rate in new-onset type 1 diabetes patients (combined analysis with 98% detection rate):
- ✓ GAD-65 antibodies
- ✓ ICA 512 antibodies
- ✓ Insulin antibodies
- ✓ ZnT8 antibodies
- The higher the # of antibodies the higher the certainty of T1DM
- 2% of patients with T2DM have positive antibodies

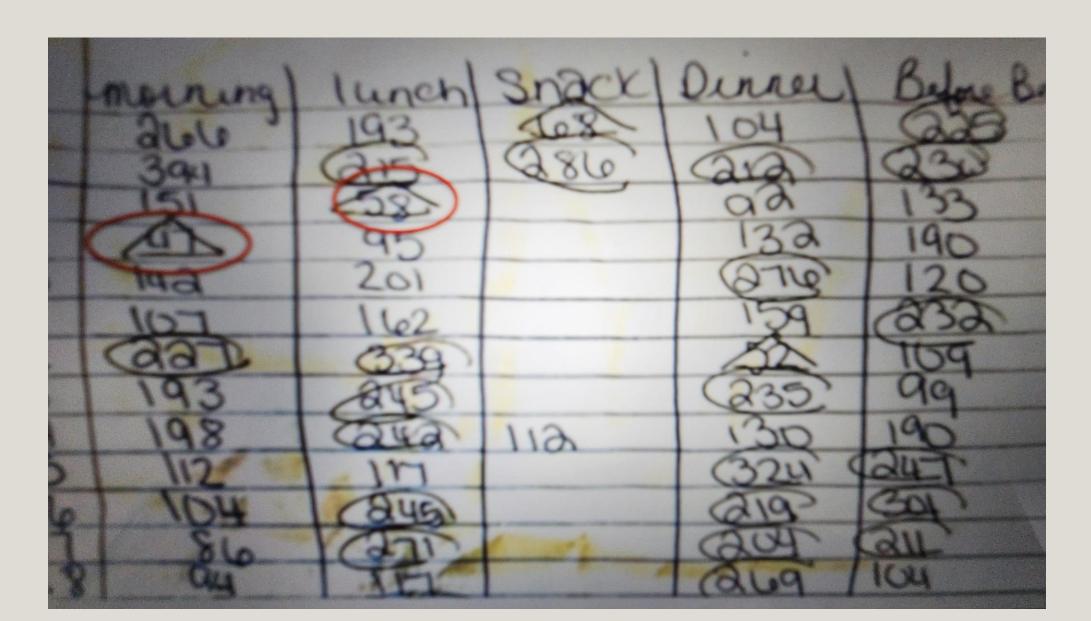
Self-monitoring blood glucose levels



Pixabay.com

Seeing is believing?

What I used to see all the time



Tips on how to use SMBG finger-sticks to get a picture of what is going on.

For the visit

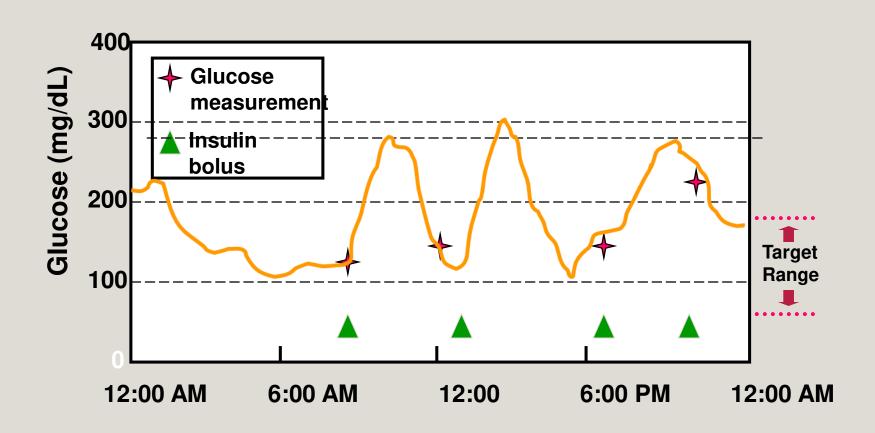
Paired tests—or 7 check in a week

	Fasting	Post Break-fast	Pre Lunch	Post Lunch	Pre Supper	Post Supper	Bed
Monday	X	X					
Tuesday							
Wed.			X	X			
Thurs.							
Friday					X	X	
Sat.							
Sunday							X

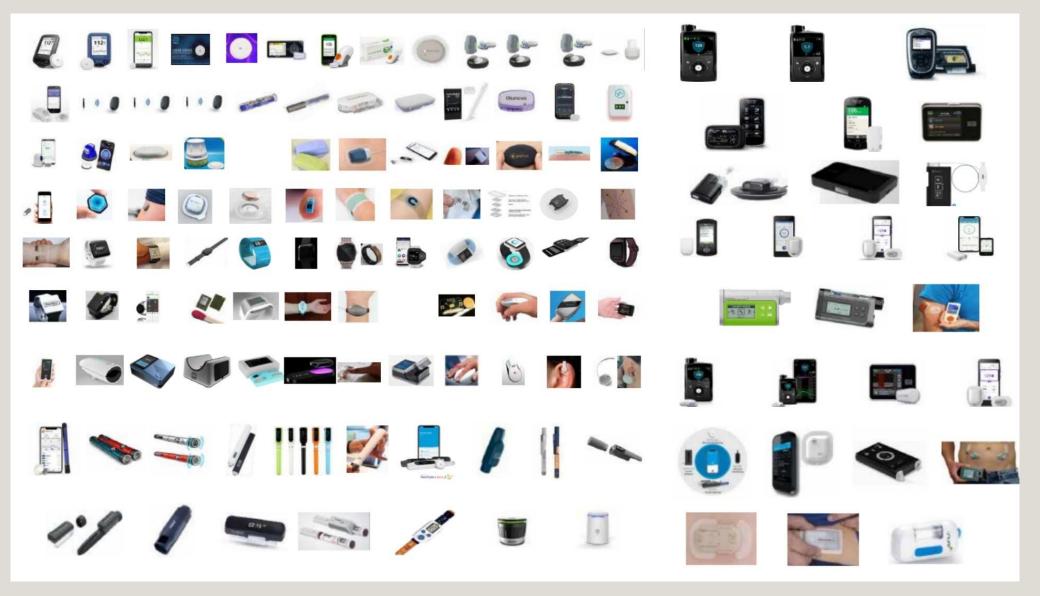
Testing 7 checks a week

	Fasting	Post Break-fast	Pre Lunch	Post Lunch	Pre Supper	Post Supper	Bed
Monday	X						
Tuesday		X					
Wed.			X				
Thurs.				X			
Friday					X		
Sat.						X	
Sunday							X

Fingerstick Blood Glucoses



Diabetes Technology is here to STAY



It's time to medically manage the patient

What do the guidelines say?

DECISION CYCLE FOR PERSON-CENTERED GLYCEMIC MANAGEMENT IN TYPE 2 DIABETES

REVIEW AND AGREE ON MANAGEMENT PLAN

- · Review management plan
- Mutually agree on changes
- Ensure agreed modification of therapy is implemented in a timely fashion to avoid therapeutic inertia
- Undertake decision cycle regularly (at least once/twice a year)
- Operate in an integrated system of care

PROVIDE ONGOING SUPPORT AND MONITORING OF:

- Emotional well-being
- · Lifestyle and health behaviors
- · Tolerability of medications
- Biofeedback including BGM/CGM, weight, step count, A1C, BP, lipids

IMPLEMENT MANAGEMENT PLAN

 Ensure there is regular review; more frequent contact initially is often desirable for DSMES

ASSESS KEY PERSON CHARACTERISTICS

- . The individual's priorities
- · Current lifestyle and health behaviors
- Comorbidities (i.e., CVD, CKD, HF)
- · Clinical characteristics (i.e., age, A1C, weight)
- Issues such as motivation, depression, cognition
- Social determinants of health

GOALS OF CARE

- · Prevent complications
- . Optimize quality of life

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CONSIDER SPECIFIC FACTORS THAT IMPACT CHOICE OF TREATMENT

- · Individualized glycemic and weight goals
- · Impact on weight, hypoglycemia, and cardiorenal protection
- Underlying physiological factors
- Side effect profiles of medications
- · Complexity of regimen (i.e., frequency, mode of administration)
- Regimen choice to optimize medication use and reduce treatment discontinuation
- · Access, cost, and availability of medication

AGREE ON MANAGEMENT PLAN

- Specify SMART goals:
 - Specific
 - Measurable
 - Achievable
 - Realistic
 - Time limited

UTILIZE SHARED DECISION-MAKING TO CREATE A MANAGEMENT PLAN

- Ensure access to DSMES
- Involve an educated and informed person (and the individual's family/caregiver)
- · Explore personal preferences
- Language matters (include person-first, strengths-based, empowering language)
- Include motivational interviewing, goal setting, and shared decision-making

WHAT IS THE CURRENT FOCUS FOR DIABETES CARE?

- Improving glycemic control
- Preventing Beta Cell failure
- Cardiometabolic health (macrovascular)
 - Weight (adiposity) reduction
 - Reducing NAFLD
 - Lipid & BP stabilization
 - Monitor for PAD
 - Monitor for hypothyroidism & other CMD risk
- Screening for & treatment of the "opathies" (microvascular)
 - Retinopathy
 - Nephropathy
 - Neuropathy (peripheral & autonomic)

Real focus on lifestyle: dietary, activity, behaviors

Will better glycemic control help macro- & microvascular impairment?

GOALS FOR ADULTS WITH DIABETES

<u>Risks</u>	Goals Les	s stringent goals	
A1C	<7.0%	<8.0%	
Blood Pressure	<140/90 mmHg	<140/90 mmHg	
Cholesterol, non-HDL	<130 mg/dL	<160 mg/dL	
Smoking, current	Nonsmoker	Nonsmoker	
Percentage meeting all goals	18.2 (14.6–22.5)	35.8 (31.7–40.2)	

GUIDELINES FOR THE "OPATHIES"

RETINOPATHY

- Adults with type 1 diabetes should have an initial dilated and comprehensive eye examination by an ophthalmologist or optometrist within 5 years after the onset of diabetes.
- People with type 2 diabetes should have an initial dilated and comprehensive eye examination by an ophthalmologist or optometrist at the time of the diabetes diagnosis.
- If there is no evidence of retinopathy for one or more annual eye exams and glycemia is well controlled, then screening every 1–2 years may be considered

GUIDELINES FOR THE "OPATHIES"

Nephropathy

- DM & DKD doubles the risk of CV mortality
- DKD increases the risk of hypoglycemia
- Annual urinary albumin (spot urinary albumin-to-creatinine ratio UACR) and eGFR assessed in T1DM ≥5 years and in all people with T2DM regardless of treatment.
- In patients with established kidney disease do the above 1-4 x annually

What are the obvious & not so obvious problems with impaired renal function in DM?

Monitor nephropathy risk & progression

					Albuminuria categories Description and range	
CKD is classified based on: • Cause (C) • GFR (G) • Albuminuria (A)			A1	A2	А3	
			Normal to mildly increased	Moderately increased	Severely increased	
		<30 mg/g <3 mg/mmol	30-299 mg/g 3-29 mg/mmol	≥300 mg/g ≥30 mg/mmol		
GFR categories (mL/min/1.73 m²) Description and range	G1	Normal to high	≥90	1 if CKD	Treat 1	Refer* 2
	G2	Mildly decreased	60-89	1 if CKD	Treat 1	Refer*
	G3a	Mildly to moderately decreased	45-59	Treat 1	Treat 2	Refer 3
	G3b	Moderately to severely decreased	30-44	Treat 2	Treat 3	Refer 3
	G4	Severely decreased	15-29	Refer* 3	Refer*	Refer 4+
	G5	Kidney failure	<15	Refer 4+	Refer 4+	Refer 4+

GUIDELINES FOR THE "OPATHIES"

Neuropathy

- Everyone with DM should be assessed for peripheral neuropathy starting at diagnosis of T2DM & 5 years post diagnosis of T1DM and annually thereafter
- Assess for distal symmetric polyneuropathy: take careful history & assessment of either temperature or pinprick sensation (small-fiber function) and vibration sensation using a 128-Hz tuning fork (for large-fiber function).
- If NO sensation—monitor for foot ulcers and fall risk.
- Monitor for signs of autonomic neuropathy as this will indicated an increased risk
 CVD and mortality

Tips for assessing peripheral neuropathy

Touch
Vibration
Temperature
Reflexes



COMPARING DISTAL DIABETES NEUROPATHIES

Large fiber

- Deep seated pain
- Wasting & weakness
- Ataxia
- Numbness, tingling, pins & needles
- Impaired vibration sense
- Loss of proprioception
- Impaired nerve conduction, reflexes
- Risks of falls & injury

Small fiber

- Superficial (C-type) pain
- Electric shocks, burning sensation
- Autonomic dysfunction
- Thermal imperception
- Normal strength & reflexes
- Highly symptomatic—miserable
- Associated with increased morbidity & mortality

What's the worst finding?

Treatment options for peripheral neuropathy

Therapy Class	Drug Name	Precautions or contraindications
Gabapentinoids	Pregabalin, Gabapentin	Dizziness, somnolence, edema
Serotonin/Norepinephrine Reuptake Inhibitor	Duloxetine, Venlafaxine	Exacerbate restless leg syndrome, sexual dysfunction, dizziness, nausea, avoid with other serotonergic drugs
Central Acting Opioid Analgesics	Tapentadol & Tramadol	Beers Criteria,
Tricyclic Antidepressants	Amitriptyline, nortriptyline	Urinary retention, cardiac disease, avoid with other serotonergic drugs, Beers criteria
Sodium Channel Blockers	Lamotrigine, Lacosamide, Oxcarbazepine, Valproic acid	Blood dyscrasias, CNS effects, liver disease
Topical Analgesic	Capsaicin, Lidocaine patches	Local burning & irritation
Anti-seizure, antimanic	Carbamazepine, Oxcarbazepine	Liver disease, Beers criteria, Hematologic effects, cardiac disease
Nutritional Supplement, Antioxidant	Alpha Lipoic Acid	None listed

Autonomic Neuropathy— LOOK for it

Cardiovascular

Resting tachycardia, orthostatic hypotension

Cardiovascular autonomic testing

Gastrointestinal

Diarrhea, constipation, gastroparesis, esophageal dysmotility, fecal incontinence

Endoscopy, colonoscopy, gastric emptying study

Genitourinary

Erectile dysfunction, retrograde ejaculation, decreased libido, painful intercourse, bladder dysfunction

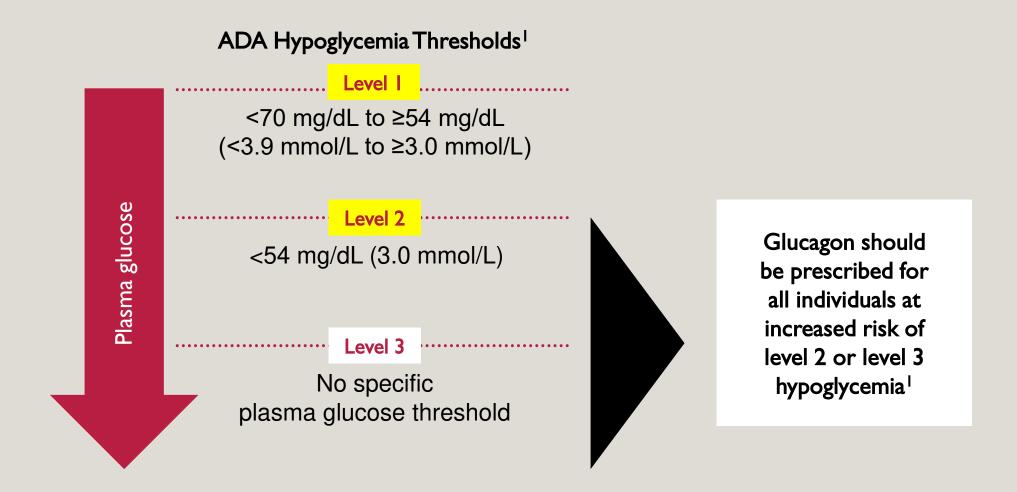
Urodynamic studies

WHAT ABOUT HYPOGLYCEMIA RISK?

When sweet goes sour

Is your patient aware and prepared?

HYPOGLYCEMIA DIAGNOSTIC CRITERIA (PER ADA)



OCCURRENCES OF HYPOGLYCEMIA

T1DM: Pediatric incidence of symptomatic hypoglycemia (SH)

- Pediatric and adult patients receiving intensive insulin therapy have 3-fold higher risk
 - Pediatric pts average of 2 episodes of SH per week
 - 30% to 40% of adults experience SH, at an average of 1-3 episodes/year

T2DM adult patients:

- 21% of adult patients on insulin experience SH once annually
 - 50% have mild/moderate episodes, with 23 events per person-year

WHY IS HYPOGLYCEMIA BAD?!!

Hypoglycemia is associated with adverse cardiovascular outcomes:

- Increased myocardial contractility, (increased O2 requirements)
- Prolonged QT interval with dysrhythmias (due to a rapid drop in serum potassium from increased circulating epinephrine and norepinephrine?), Ischemic electrocardiogram changes and repolarization abnormalities,
- Angina, arrhythmias, increased inflammation, and sudden death
- Mechanisms for the poor outcome are not completely understood,
 - Likely d/ tincreases in pro-inflammatory cytokines (TNFα, IL-1β, IL-6, and IL-8), markers of lipid peroxidation,
 - And acute endothelial dysfunction with associated vasoconstriction, increased blood coagulability, cellular adhesion, and oxidative stress

THE SIGNIFICANCE OF SEVERE HYPOGLYCEMIA

Morbidity and mortality of SH are significant and can be barriers to adherence and intensification of diabetes treatments in patients with T1DM and T2DM.



hypoglycemia¹

"Dead-in-bed" syndrome (up to 6% of deaths in patients with T1DM <40 years of age)⁴



Additional clinical consequences

- Increased risk for cardiovascular events
- Loss of consciousness or coma
- Seizures
- Impaired cognitive function
- Increased risk for more SH episodes
- Fear, depression & quality of life impairment

SH=severe hypoglycemia; T1DM=Type 1 diabetes mellitus; T2DM=Type 2 diabetes mellitus.

HYPOGLYCEMIA RISK MEDICATIONS & CONDITIONS

- warfarin
- quinine
- salicylates
- fibrates
- Sulfonamides (including co-trimoxazole)
- monoamine oxidase inhibitors
- NSAIDs
- probenecid
- somatostatin analogues
- selective serotonin reuptake inhibitors.
- <u>loss of counter-regulatory hormone function</u>, (e.g., Addison's disease, growth hormone deficiency, hypothyroidism, or hypopituitarism.)

MANIFESTATIONS OF HYPOGLYCEMIA

- Feeling shaky
- Being nervous or anxious
- Sweating, chills and clamminess
- Irritability or impatience
- Confusion
- Fast heartbeat
- Feeling lightheaded or dizzy
- Hunger
- Nausea
- Color draining from the skin (pallor)

- Feeling sleepy
- Feeling weak or having no energy
- Blurred/impaired vision
- Tingling or numbness in the lips, tongue or cheeks
- Headaches
- Coordination problems, clumsiness
- Nightmares or crying out during sleep
- Seizures

TREATMENT OF OUTPATIENT HYPOGLYCEMIA

- Prevention!
- Quick recognition of symptoms (neuronal starvation)
- Rapid-acting carbohydrate
- Glucagon

CARBOHYDRATE TREATMENT

- Patient must be conscious and able to swallow!!!
- The rule of 15: Administer approximately 15 g (or 20 g) of rapid-acting carbohydrate and check BG 10-15 minutes later
 - Repeat dose as needed x 3.
 - If sugars <u>still low---need glucagon</u>

FAST-ACTING CARBOHYDRATE POINTERS

Forms of rapid-acting carbs

- Sucrose & glucose best (tabs, syrup, drinks)
- Is orange juice best?
- What about chocolate or milk?

Amount of carbohydrates is best?

- Which is the most effective...10g, 15 g, or 20 g?
- Should it be weight-dosed?

Time to resolution of symptoms

- If not resolved quickly it will lead to an even slower response with next dose
- How soon should you really wait to recheck sugars....5 minutes?,10 minutes? Longer?

GLUCAGON ADMINISTRATION

- Glucagon kits are reconstituted with the provided solution to a 1 mg/mL clear, particulatefree solution. The entire 1 mg (1 mL) is administered SC or IM
- Other formulations of glucagon (premixed injectable solutions and nasal sprays) have recently been developed
- Reversal of hypoglycemia relies on sufficient hepatic glycogen stores and other factors. Patients normally respond within 15 minutes;
- Will Not work in those with alcohol toxicity (blunted glycogen response)
- ONLY USE ONCE—do not repeat (won't work)
- NEW glucagon pens and nasally-inhaled glucagon (Zegalogue, Gvoke, BAQSIMI)

So what pills, pokes and potions do we have for diabetes treatment?

ORAL MEDICATIONS FOR DIABETES

<u>Biguanides</u>

*metformin

Sulfonylureas

- *glipizide
- *glyburide
- *glimepiride

Thiazolidinediones (TZDs)

*pioglitazone (Actos) rosiglitazone (Avandia)

GLP-1 agonists

semaglutide (Rybelsus)

DPP-4 Inhibitors

sitagliptin (Januvia) saxagliptin (Onglyza) linagliptin (Tradjenta) alogliptin (Nesina)

SGLT 2 Inhibitors

dapagliflozin (Farxiga)
canagliflozin (Invokana)
empagliflozin (Jardiance)
ertugliflozin (Steglatro)
Bexagliflozin (Brenzavvy)

* = generic available

Meglinatides

- *nateglinide (Starlix)
- *repaglinide (Prandin)

Alpha-glucosidase inhibitors

- *acarbose (Precose)
- *miglitol (Glyset)

Bile acid sequestrants

*colesevelam (Welchol)

Dopamine agonist

bromocriptine (Cycloset ONLY)

INJECTABLE MEDICATIONS FOR DIABETES

GLP-1 agonist

liraglutide (Victoza) dulaglutide (Trulicity) exenatide ER (Bydureon, Bcise) exenatide (Byetta) Semaglutide (Ozempic)

GLP-1/GIP agonist

tirzapetide (Mounjaro)

Basal Insulin

insulin glargine u100*
(Lantus, Basaglar)
insulin detemir u100 (Levemir)
insulin glargine u300 (Toujeo)
insulin degludec u100 or u200
(Tresiba)

Rapid Acting Insulin

Regular Insulin

insulin regular human u100 (Humulin R, Novolin R, Relion R) Insulin regular human u500 (Humulin R u500)

Intermediate Insulin

Human insulin isophane, aka NPH (Humulin N, Novolin N, Relion N)

Split mixes

Humulin 70/30 (70% NPH, 30% regular)
Novolin 70/30 (70% NPH, 30% regular)
Relion 70/30 (70% NPH, 30% regular)
Humalog 50/50 (lispro /lispro protamine)
Humalog 75/25 (75% lispro protamine, 25% lispro

Mixed injectables - GLP1 and basal insulin

insulin degludec and liraglutide (Xultophy) insulin glargine and lixisenatide (Soliqua)

Amylin mimetic

Pramlinitide (Symlin)

*Indicates units per mL: u100 = 100 units/mL, u300 = 300 units/mL, etc.

MORE ON WHY, HOW AND WHEN TO USE THESE MEDICATIONS LATER!

WHAT O-T-C SUPPLEMENTS OR FOOD MIGHT YOUR PATIENTS BE TAKING FOR DIABETES PREVENTION OR CONTROL?

Do they help?

- Cinnamon
- Chromium
- Vitamin BI
- Bitter Melon
- Alpha lipoic acid
- Aloe Vera
- Green tea
- Resveratrol
- Magnesium
- Others.....



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Can we cure diabetes?



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Time for a brain break



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