

# Insulin and More Insulins, Oh My!

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an FQHC

# Learning Objectives

- Identify situations where insulin is the recommended treatment for diabetes management.
- Explain insulin using patient friendly concepts
- Describe protocols on how to dose insulin and convert patients to different types of insulin.
- Examine the variety of biosimilar insulins and compare and contrast differences.

# Common Diabetes Terminology

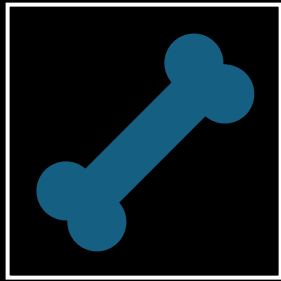
A1C

Insulin  
resistance

Glucose  
toxicity

Catabolism

# Common “Misterminology”



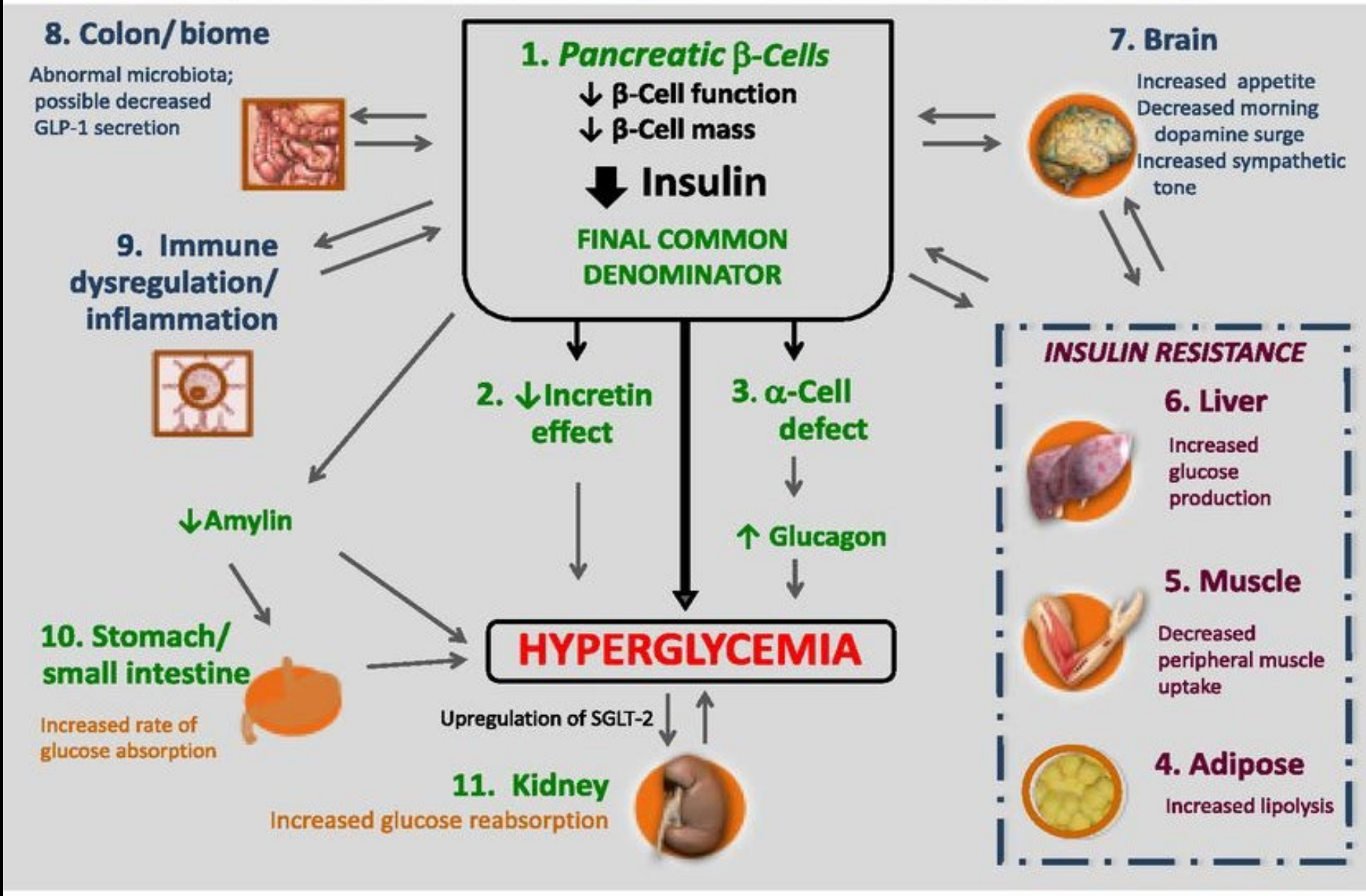
”Brittle Diabetic”



“Insulin dependent  
diabetes”



“A little too sweet”

**A** **$\beta$ -Cell-Centric Construct: Egregious Eleven****The  $\beta$ -Cell is the FINAL COMMON DENOMINATOR of  $\beta$ -Cell Damage**

# Insulin Timeline

- 1889 – dogs developed diabetes and died when pancreas removed
- 1910 – “insula” was missing in people with diabetes
- 1921 - insulin “muck” injections kept a dog with diabetes alive
- 1922 – Leonard Thompson kept alive on insulin injections
- 1923 - Banting, Macleod, Best, and Collip -Nobel Prize in Medicine
- 1924 – first large scale production of animal insulins for human use
- 1936 - first slower acting animal insulin for human use
- 1978 – first genetically engineered insulin
- 1982 – first marketing of genetically engineered insulin
- 2025 – so many, but many the same...

# Shout Out!

What are some reasons medical providers may be hesitant to use insulin?

# Provider barriers to insulin

- Doses are not standardized
- We are familiar with the serious potential side effects
- There are so many insulins to choose from
- We underestimate our patients
- These discussions take time and can be very emotional

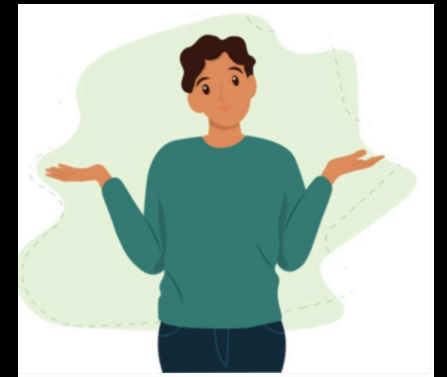
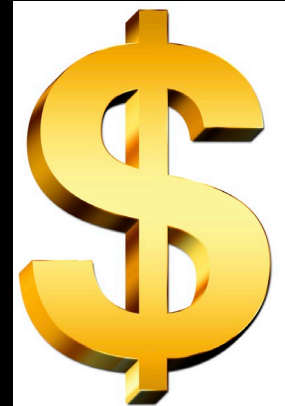


# Shout Out!

What are some reasons patients are afraid of insulin?

# Barriers to Optimal Insulin Therapy

- Fear of taking insulin (i.e. needle phobia, side effects)
- Health beliefs regarding insulin
- Complexity of regimen
- Cost
- Inadequate follow-up and support
  - Dose titrations (i.e. doesn't work)
  - Side effects (i.e. hypoglycemia and doses not adjusted)
  - Not shown how to use devices

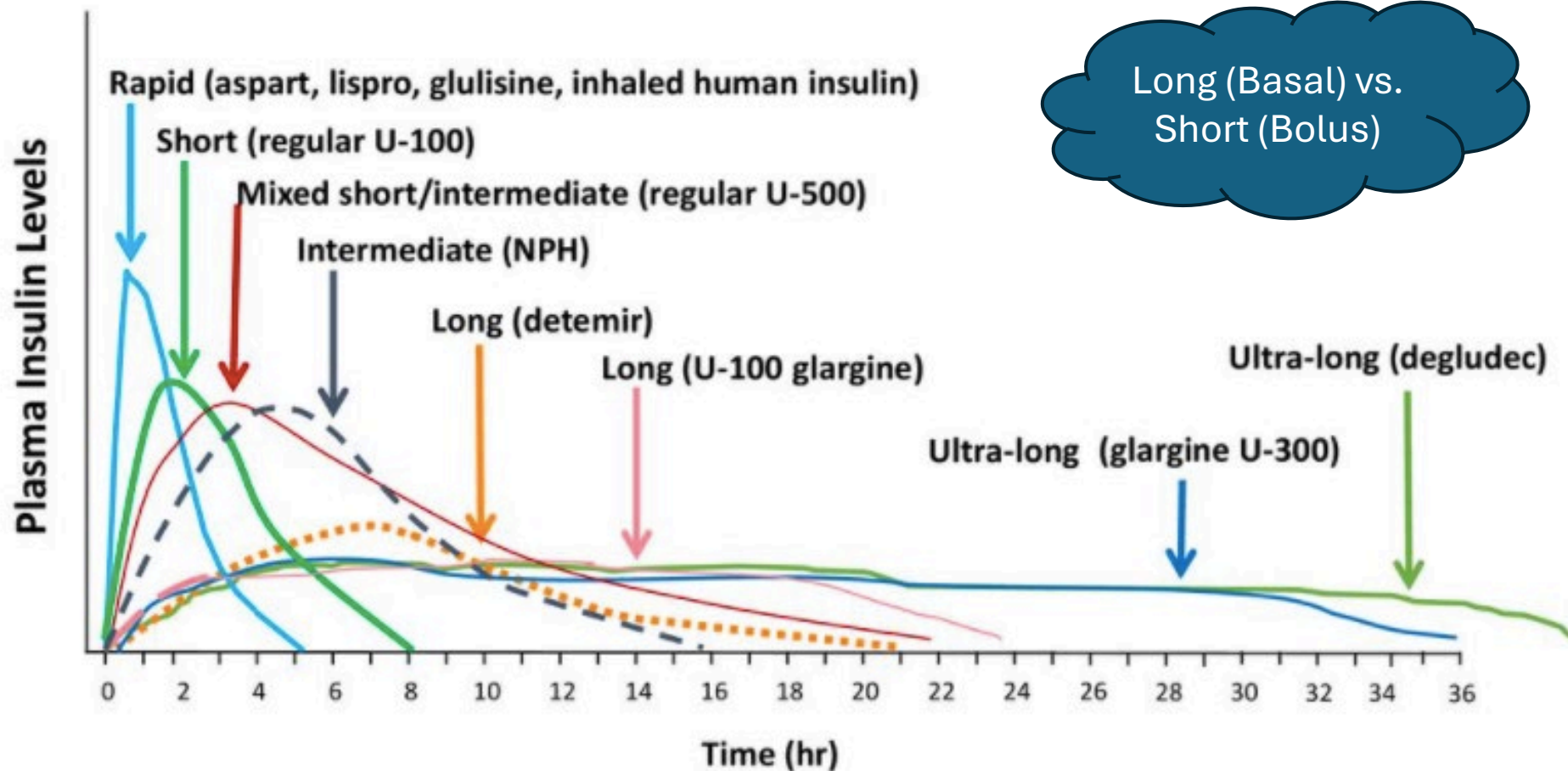


Shout Out!

Who is on Insulin?



# Pharmacokinetic Profile of Currently Available Single Insulin Products



# Sample Patient Case

46 year old male. A1c 8 months ago was 5.9%.

Presents to ED feeling very ill. Unintentional weight loss of 12 pounds in past 2 months and appears underweight today. Sunken facial features and polydipsia/polyuria. A1c today is 12.8% and Blood sugar 550.

Considerations:

- DKA or HHS???
- Autoimmune vs. T2DM vs. underlying condition

# Diabetic Ketoacidosis Vs. Hyperosmolar Hyperglycemic State (DKA Vs . HHS)

A. DKA Diagnostic Criteria		
DKA	Diabetes/hyperglycemia	Glucose $\geq 200$ mg/dL (11.1 mmol/L) OR prior history of diabetes
	Ketosis	$\beta$ -Hydroxybutyrate concentration $\geq 3.0$ mmol/L OR urine ketone strip 2+ or greater
	Metabolic Acidosis	pH $< 7.3$ and/or bicarbonate concentration $< 18$ mmol/L
B. HHS Diagnostic Criteria		
HHS	Hyperglycemia	Plasma glucose $\geq 600$ mg/dL (33.3 mmol/L)
	Hyperosmolarity	Calculated effective serum osmolality $> 300$ mOsm/kg (calculated as $[2 \times \text{Na}^+ \text{ (mmol/L)} + \text{glucose (mmol/L)}]$ ), OR total serum osmolality $> 320$ mOsm/kg $[(2 \times \text{Na}^+ \text{ (mmol/L)} + \text{glucose (mmol/L)} + \text{urea (mmol/L)})]$
	Absence of significant ketonemia	$\beta$ -Hydroxybutyrate concentration $< 3.0$ mmol/L OR urine ketone strip less than 2+
	Absence of acidosis	pH $\geq 7.3$ and bicarbonate concentration $\geq 15$ mmol/L

# DKA vs. HHS

- Use a free online calculator such as [globalrph.com](http://globalrph.com)

ANION GAP	
Sodium level (Na+):	<input type="text"/> meq/L
Chloride level (Cl-):	<input type="text"/> meq/L
Bicarb level (HCO <sub>3</sub> -):	<input type="text"/> meq/L
<input type="button" value="Calculate Anion Gap"/> <input type="button" value="Reset"/>	
BACKGROUND	
Normal range: 8 to 16 meq/L	
Anion gap = [Na+] - [Cl-] - [HCO <sub>3</sub> -]	

OSMOLALITY CALCULATOR	
Osmolality = sodium x 2 + glucose/18 + bun/2.8 + Etoh/4.6 Normal range: 285-295 mOsm/kg	
Osmolality of blood increases with dehydration and decreases with overhydration. In normal people, increased osmolality in the blood will stimulate secretion of antidiuretic hormone (ADH). This will result in increased water reabsorption, more concentrated urine, and less concentrated blood plasma. A low serum osmolality will suppress the release of ADH, resulting in decreased water reabsorption and more concentrated plasma.	
Sodium:	<input type="text"/> mEq/L
Glucose:	<input type="text"/> mg/dL
Bun:	<input type="text"/> mg/dL
Etoh (optional):	<input type="text" value="0"/> mg/dL
<input type="button" value="Calculate"/> <input type="button" value="Reset"/>	

# Insulin as Primary Therapy

- Autoimmune Mediated
  - Type 1 Diabetes Mellitus
  - Latent Autoimmune Diabetes of Adulthood (LADA)
- Diabetic Ketoacidosis
- Hyperosmolar Hyperglycemic Syndrome
- Type 2 Diabetes Mellitus
  - Catabolism
  - Pancreatic insufficiency
    - Due to chronic pancreatitis , surgery, cystic fibrosis (underlying conditions)
    - Long standing T2DM, less than desirable A1c despite optimal other medications
  - Diabetes from “seemingly out of nowhere”



# Similarities between Autoimmune Diabetes (T1DM and LADA) **-AND-** Diabetes Due to Chronic Pancreatitis

- Insulin deficiency is primary disease process
- Body weight is typically lean to underweight
- Lack of insulin resistance
- C-peptide levels will be low even at high glucose levels


# Differences between Autoimmune Diabetes (T1DM and LADA) –AND– Diabetes Due to Chronic Pancreatitis

- Age of Onset
  - Type 1 typically young age
  - Latent Autoimmune Diabetes of **Adulthood**
- Presence of Autoimmune Antibodies (usually one is positive)
  - GAD-65
  - Islet cell Ab
  - Zinc Transporter
  - IA-2 (Insulinoma Associated Protein)
  - Others TBD???

# Diabetes is Like A Wildfire.....



# IHS Clinical Resources - Algorithms



**Indian Health Service**  
The Federal Health Program for American Indians and Alaska Natives

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About Us

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Clinical Training

Education Materials and Resources (Online Catalog)

**Clinical Resources**

- Diabetes Standards of Care and Resources for Clinician and Educators
- Diabetes Treatment Algorithms**
- Diabetes Education Lesson Plans
- Diabetes Educator Tools
- Kidney Health

Fact Sheets and Publications

IHS Diabetes Audit

Special Diabetes Program for Indians (SDPI)

## Clinical Resources

The IHS Division of Diabetes developed the resources below specifically for clinicians and educators working with American Indian and Alaska Native diabetes patients.

Search Division of Diabetes and SDPI


[Standards of Care](#) [Algorithms](#) [Diabetes Education Lesson Plans](#) [Diabetes Educator Tools](#) [Kidney Health](#)

### Diabetes Treatment Algorithms

These algorithms provide clinicians with a quick reference to specific diabetes care recommendations based on national guidelines and the [Diabetes Standards of Care and Resources for Clinicians and Educators](#). These tools are developed with Indian Health Service clinical consultants, pharmacists, and other professionals.

The algorithms provide:

- Basic information of diabetes-related conditions.
- Step-by-step management of common clinical problems.
- Dosing, common adverse reactions, and contraindications for medications on the IHS National Core Formulary.
- Treatment targets and recommended monitoring parameters.



The Diabetes Treatment Algorithms are intended to serve as a tool for providers who treat patients with type 2 diabetes. They are updated periodically but changes in national practice may occur more quickly—users are advised to stay abreast of current clinical practice recommendations.

# Initiation and Titration of Insulin

## Step 1: Start basal insulin therapy

- Start long-acting analog 10 units or 0.1-0.2 units/kg once a day (may advance to BID if necessary).
- Increase by 2-4 units or 10-15% every 3-4 days until fasting blood glucose (FBG) values fall within target range, generally 80-130 mg/dL, OR adjust dose following review of CGM profile.

▼ **If A1C\* not at goal but FBG at target and/or basal insulin dose >0.5 unit/kg/day**  
If not already on a GLP1-RA or tirzepatide, consider adding to basal insulin or starting long-acting insulin/GLP1-RA combination before adding mealtime insulin.

## Step 2: Add mealtime insulin before largest meal

- Start rapid-acting analog 2-4 units or 10% of basal insulin dose before largest meal.
- Increase insulin by 1-2 units or 10-15% until blood glucose (BG) within target range, generally 80-130 mg/dL premeal or <180 mg/dL 1-2 hours after meals, OR utilize CGM profile to guide dose adjustments (individualize).

▼ **If A1C\* not within target range**

## Step 3: Add mealtime insulin before other meals

- Start additional rapid-acting analog 2-4 units or 10% of basal insulin dose before other meals (focus on one meal at a time).
- Increase insulin by 1-2 units or 10-15% every 3-7 days until BG falls within target range, generally 80-130 mg/dL premeal or <180 mg/dL 1-2 hours after meals, OR utilize CGM profile to guide dose adjustments (individualize).

# KK's Initiation of Basal and Initial Titration

- 10 units once daily – (6 units if normal or underweight)
  - Focus on AM fasting sugars
  - Initial AM fasting goal – get into the 100's
  - \*\*weekly calls if able and increase by 10 units if >200, smaller increase if already in upper 100's.
- 
- Once AM fastings in the 100's, then change focus and start checking 1 to 2 hours after dinner sugars
  - Address post prandial sugars with additional meds if necessary

# Our Patient



- Recap: adult aged, sudden onset, losing weight, underweight, and acutely ill
- Is there a history of pancreatitis?
- Insulin 6 units once a day, check AM fastings
  - Do initial dose calculation with patient ( $60\text{kg} \times 0.2 = 12$  units)
- If the patient absolutely refuses insulin shots  $\Rightarrow$  glipizide??
  - Challenge to prove sugars are coming down
- Close follow up

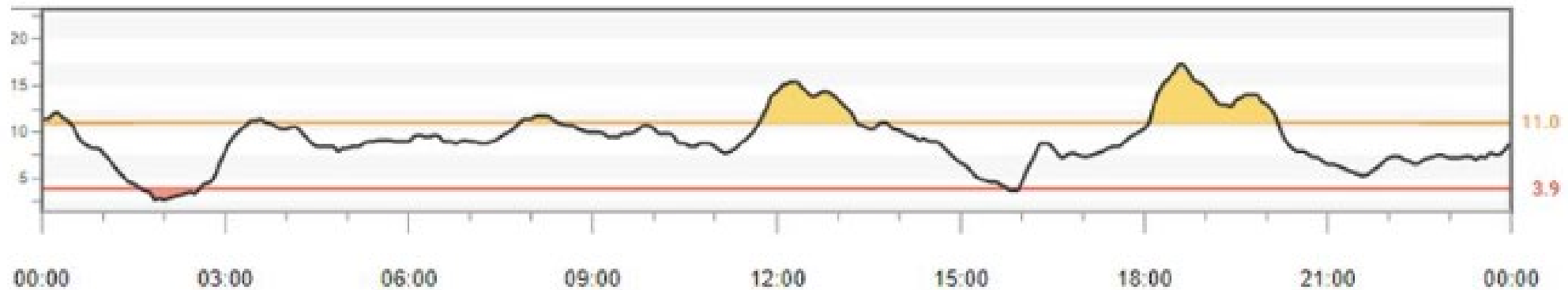
# Patient Managed - Super Simple Insulin Titration

- Especially effective if weekly follow up not possible
- Engages patient in checking and dosing
- Check AM fasting sugar
- If AM fasting sugar  $>200$ , then increase basal insulin dose by 1 unit that day
- If most AM fasting sugars are  $<200$ , then stay at that basal dose until seen again by provider
- If AM fasting sugar  $<80$ , then decrease basal dose by 5 units daily

\*can take patients VERY long to get to optimal dose if insulin resistant

# Adequacy of Basal Insulin Dose

- Indications of “over basalization”
  - Basal dose  $> 0.5$ units/kg/day, and not yet on prandial insulin
  - HS to AM drop of more than 50 points
  - High variability, lots of lows when not eating
  - Large post-prandial to pre-prandial differentials

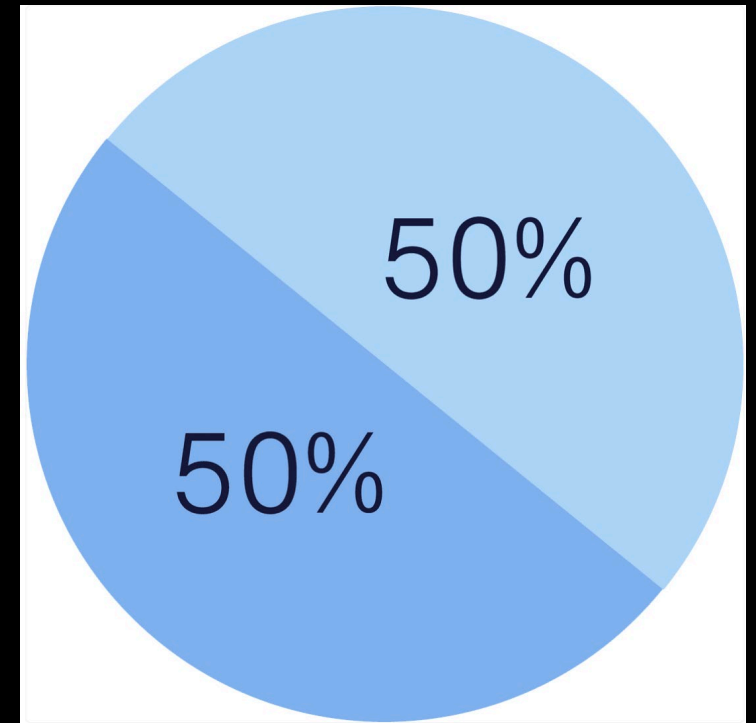


# Consider insulin over other post prandial medicines when:

- Already on GLP-1
- Not able to use GLP-1
  - T1DM
  - Intolerance
  - Gastroparesis
  - chronic pancreatitis
  - weight loss not desirable
- Post prandial sugars VERY high
- Pancreatic insufficiency patients

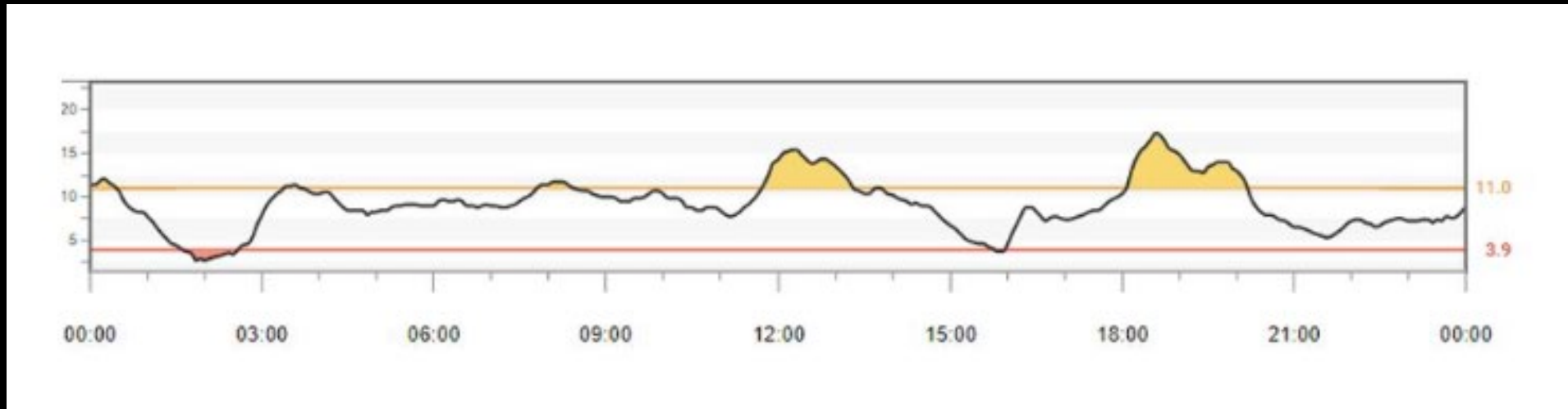
# Moving from Over Basalized to Balanced

- Calculate a total daily dose
- Compare it physiologic insulin ratios
  - 50/50 – heavier carb eaters
  - 70/30 – more carb balanced eaters
  - Evaluate where to go.....
    - Simply add on bolus doses?
    - Keep daily dose the same, but adjust ratios
  - Discuss your rationale with the patient – this gives you a second check



# Test Your Knowledge

- Pt on 100 units of glargine U300 once a day & metformin 2000mg daily



- Use a 70/30 calculation to decide separate basal and bolus doses
- $100 \text{ units} \times 0.7 = \text{long}$  (70 units glargine once a day)
- $100 \text{ units} \times 0.3 = \text{short}$  (30 units TTD = 15 units BID or 10 units TID, with meals – lispro or aspart)

# Insulin Detemir

- No longer produced
- Considered a basal insulin
  - Lower doses < 20 hours duration of action
  - Higher doses >20 hours duration of action
- Dose conversions
  - Calculate total daily dose of detemir
  - If at A1c goal or hypoglycemia: convert at 80%
  - If above A1c goal and no hypoglycemia: convert one to one

\*\*in general, patients required a slightly higher dose of detemir to reach the same lower effect of insulins such as glargine.



# Test Your Knowledge

- Pt on detemir 25 units BID, but no longer available (50 units daily)
  - Patient's insurance covers insulin glargine (and degludec)
    - Patient with A1c close to goal and felt one episode of low sugar (into the low 80's) overnight last week .
    - What would you recommend?
      - Glargine (or degludec) at 40 units once daily (new dose 80% of old 50 units)
    - Patient with A1c of 12%, lowest sugar in past 3 weeks was 180.
    - What would you recommend?
      - Glargine (or degludec) at 50 units daily (one to one conversion)
- \*\*concentrated glargine or concentrated degludec might be preferable if patient is on higher doses of basal insulin

# Combo Insulins to Separate Basal/Bolus

- Combo pens usually limited to 60 units per injection
- 70/30 - 70% long, 30% short
- 75/25- 75% long, 25% short
- 50/50 – 50% long, 50% short
- Conversion (example 70/30 to glargine/aspart)
  - Calculate total daily dose
  - 70% total daily dose to be given as once daily basal
  - 30% total daily dose DIVIDED and given with 2-3 largest meals
  - This is a starting place and can be adjusted individually to move toward needs of patient evidenced by blood glucose or symptoms.

# Test Your Knowledge

- Pt on 50 units BID of Humulin 75/25 = 100 units TTD
  - Patient with A1c close to goal and felt one episode of low sugar (into the low 80's) overnight last week .
  - What would you recommend?
    - Glargine (or degludec) at 60 units once daily (new dose 80% of old basal 75)
    - Aspart or lispro 20 units split between 2 or 3 meals. (80% of old bolus 25)
  - Patient with A1c of 12%, lowest sugar in past 3 weeks was 180.
  - What would you recommend?
    - Glargine (or degludec) at 75 units daily (75% long, one to one conversion)
    - Aspart or lispro 25 units split between 2 or 3 meals. (100% short, one to one)

# Insulin Therapy in Diabetes Treatment

The tables below provide estimates of insulin pharmacokinetic profiles of various preparations. Patients with type 2 diabetes may require high doses due to insulin resistance. Injection of large insulin boluses affects insulin absorption and activity.

	Administration/ Timing with Meals	Peak (hrs)	Duration (hrs)	Max Pen Dose (units)
<b>Long Acting Insulin</b>				
Detemir U-100 ( <i>Levemir</i> )	Usually at bedtime	Slow or no pronounced peak	8-24	80
<b>Glargine U-100</b> ( <i>Lantus</i> , <i>Semglee*</i> <i>Basaglar*</i> , <i>Rezvoglar*</i> )			Up to 24	80
<b>Glargine U-300</b> ( <i>Toujeo</i> )			24-36	80 <i>Toujeo</i> / 160 <i>Toujeo Max</i>
Degludec U-100, U-200 ( <i>Tresiba</i> )			Up to 42	80 (U-100)/ 160 (U-200)
<b>Intermediate Acting Insulin</b>				
NPH U-100 ( <b><i>Novolin N</i></b> )	Usually at bedtime (onset 1-2 hrs)	2-8	14-24	60
<b>Short Acting Insulin</b>				
Regular U-100 ( <b><i>Novolin R</i></b> , <i>Humulin R</i> )	30 min before	2-4	6-12	60
<b>Rapid Acting Insulin</b>				
Aspart U-100 ( <b><i>Novolog</i></b> , <i>Kirsty+</i> )	Within 5-10 min before	~1-2	5-7	60
Aspart U-100 ( <i>Fiasp</i> )	At the start or within 20 min after start	1	3-5	80
Lispro ( <i>Humalog U-100</i> , <i>U-200</i> ; <i>Admelog* U-100</i> )	Within 15 min or immediately after	1-2	3-5	60 - <i>Humalog</i> 80 - <i>Admelog</i>
Lispro-aabc ( <i>Lyumjev U-100</i> , <i>U-200</i> )	At the start or within 20 min after start	1	2-4	60 - <i>Lyumjev</i> <i>U-100</i> , <i>U-200</i>
Glulisine U-100 ( <i>Apidra</i> )	Within 15 min before or 20 min after start	1-2	3-6	80

# Insulin Therapy in Diabetes Treatment

	Administration/ Timing with Meals	Peak (hrs)	Duration (hrs)	Max Pen Dose (units)
<b>Premixed Insulin</b>				
NPH/Regular U-100 ( <i>Novolin 70/30</i> )	Use guidance for short- acting or rapid-acting insulin component	2-4	18-24	60
NPH/Aspart U- 100( <i>Novolog Mix 70/30</i> )		1-2	12-24	60
NPH/Lispro U-100 ( <i>Humalog Mix 75/25</i> and <i>Mix 50/50</i> )		1-2	13-22	60
<b>Concentrated Regular Insulin</b>				
Regular U-500 ( <i>Humulin R</i> <i>U-500 Kwikpen</i> )	30 min before	0.5-8	13-24	300
Patients with severe insulin resistance requiring >200 units per day of insulin are candidates for Regular U-500. Total daily dosing may be started BID split 50-50 with meals or TID split 40-30-30 with breakfast, lunch, and dinner meals respectively.				
<b>Inhaled Insulin</b>				
Inhaled Regular Insulin ( <i>Affreza</i> )	At the start	0.5-0.9	1.5-3	12 unit cartridge
<b>Long-Acting Insulin/Glucagon-like Peptide-1 Receptor Agonist Combinations</b>				
Insulin degludec/ liraglutide ( <i>Xultrophy</i> <i>100/3.6</i> )	30 min before	No peak	24	50 units/ 1.8 mg
Insulin glargine/ lixisenatide ( <i>Soliqua</i> <i>100/33</i> )	Within 1 hour prior to first meal of the day	No peak	20-24	60 units/ 20 mcg

## Biosimilar and Interchangeable Insulins

\* **Semglee** (glargine-yfgn), **Rezvoglar** (glargine-aglr), and **Basaglar** are biosimilar and do not differ clinically from **Lantus** (insulin glargine).

+ **Admelog** (lispro) is biosimilar to **Humalog**, and **Kirsty** (aspart) is biosimilar to **Novolog**.

**Semglee and Rezvoglar** are also interchangeable with **Lantus**, meaning that the pharmacist can substitute without notifying the provider.

Medications on the [IHS National Core Formulary](#) are in **BOLD** above. Please consult a complete prescribing reference for more detailed information. No endorsement of specific products is implied.

## Network meta-analysis estimates for change in HbA1c level (green) and incidence of nocturnal hypoglycemia (white) for each comparison of basal insulin analogues

Detemir	1.03 (0.88 to 1.21)	1.39 (1.12 to 1.71)	0.91 (0.70 to 1.20)	-	-	0.64 (0.50 to 0.82)
0.15 (0.04 to 0.25)	Glar-100	1.34 (1.17 to 1.51)	0.89 (0.71 to 1.11)	-	-	0.62 (0.49 to 0.78)
0.20 (0.05 to 0.35)	0.05 (-0.05 to 0.16)	Glar-300	0.66 (0.51 to 0.86)	-	-	0.46 (0.35 to 0.61)
0.15 (-0.02 to 0.33)	0.00 (-0.14 to 0.14)	-0.05 (-0.23 to 0.12)	<b>Basaglar</b>	-	-	0.70 (0.51 to 0.97)
0.13 (-0.15 to 0.41)	-0.02 (-0.28 to 0.24)	-0.07 (-0.35 to 0.20)	-0.02 (-0.32 to 0.27)	MK-1293	-	-
0.09 (-0.24 to 0.42)	-0.06 (-0.37 to 0.25)	-0.11 (-0.44 to 0.22)	-0.06 (-0.41 to 0.28)	-0.04 (-0.44 to 0.36)	<b>Semglee</b>	-
0.09 (-0.07 to 0.24)	-0.06 (-0.20 to 0.07)	-0.12 (-0.29 to 0.05)	-0.07 (-0.26 to 0.13)	-0.04 (-0.33 to 0.25)	0.00 (-0.34 to 0.34)	<b>NPH</b>

Favors glargine 100, glargine 300, A1C lowering

Favors glargine 300 for nocturnal hypoglycemia

Favors long-acting analogs over NPH for nocturnal hypoglycemia

## REFERENCE PRODUCT



Original FDA-approved biological product.

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Prescribed by a provider.

## BIOSIMILAR



Highly similar and with no clinically meaningful differences when compared to the reference product.

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Prescribed by a provider.

## INTERCHANGEABLE PRODUCT

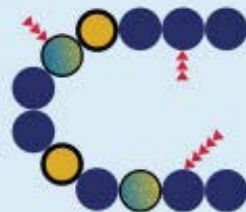


Highly similar and with no clinically meaningful differences when compared to the reference product.

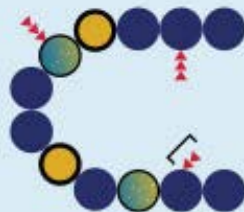
The application includes additional data and information about the impact of switching or alternating between the product and the reference product.

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May be substituted at the pharmacy without the intervention of the prescribing provider.

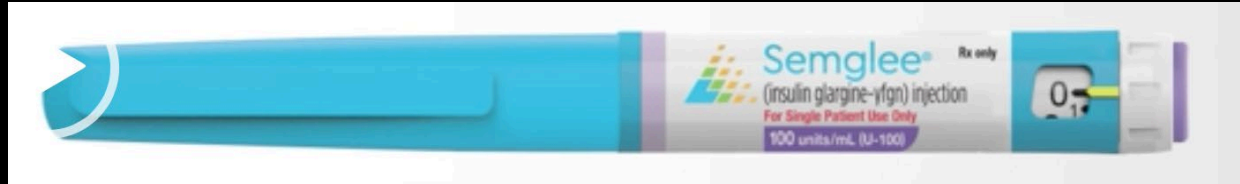
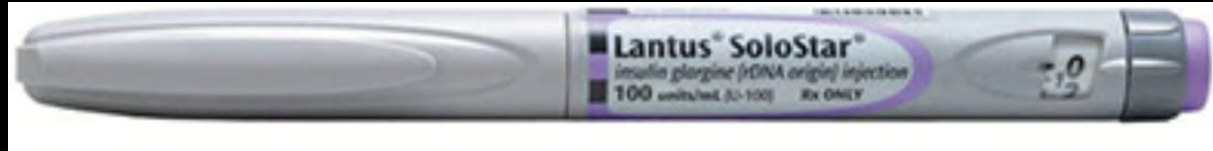


Reference product



Biosimilar product

# Glargine U100 Biosimilar AND Interchangeable

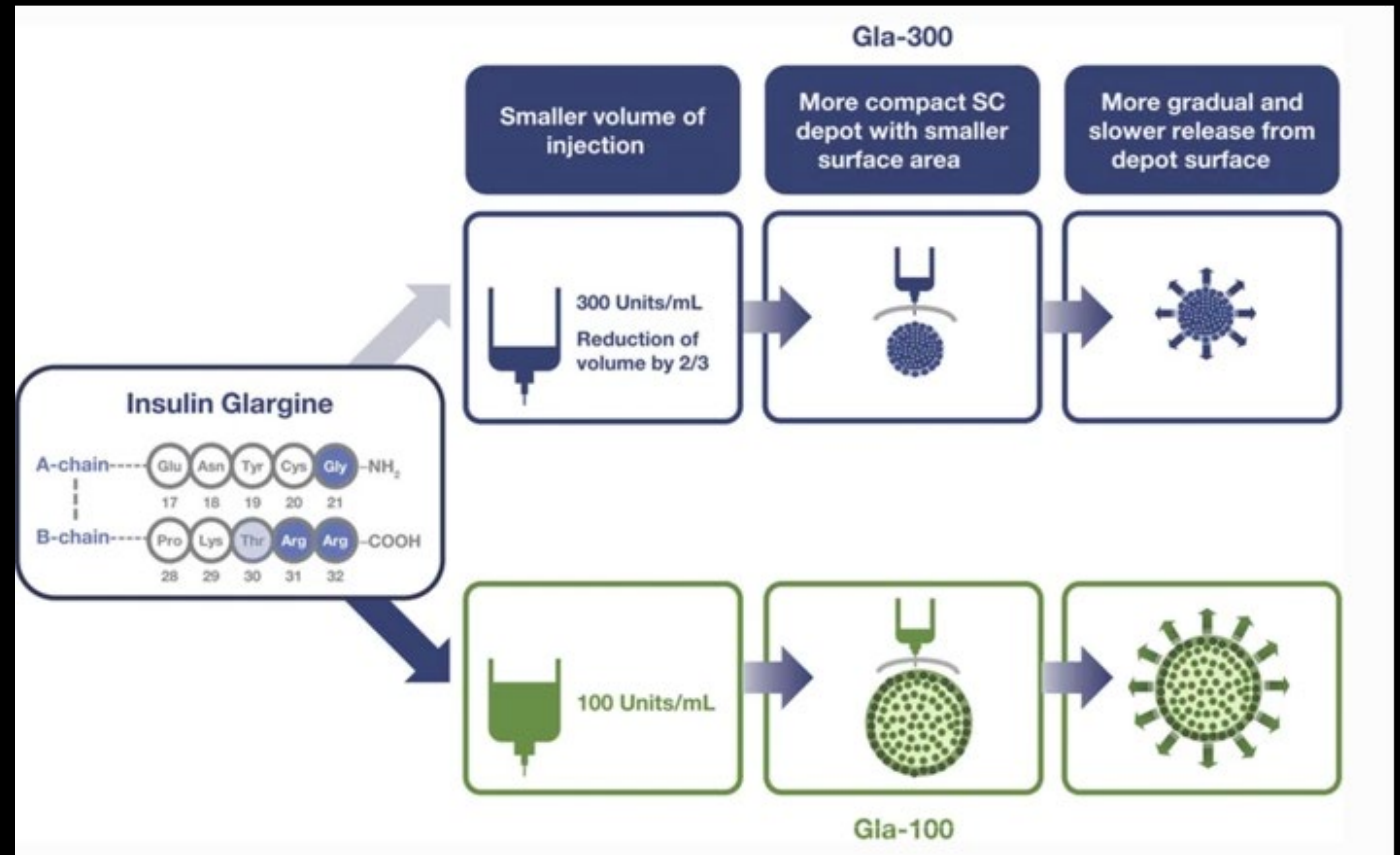


# Glargine U100 Biosimilar NOT Interchangable



# High Dose Basal Options

- Consider when requiring higher daily doses of basal insulin (esp >80 units/day)
- Up to 160 units per injection
- Insulin degludec 200units/ml (Tresiba®)
- Insulin glargine 300 units/ml (Toujeo Max SoloStar®)

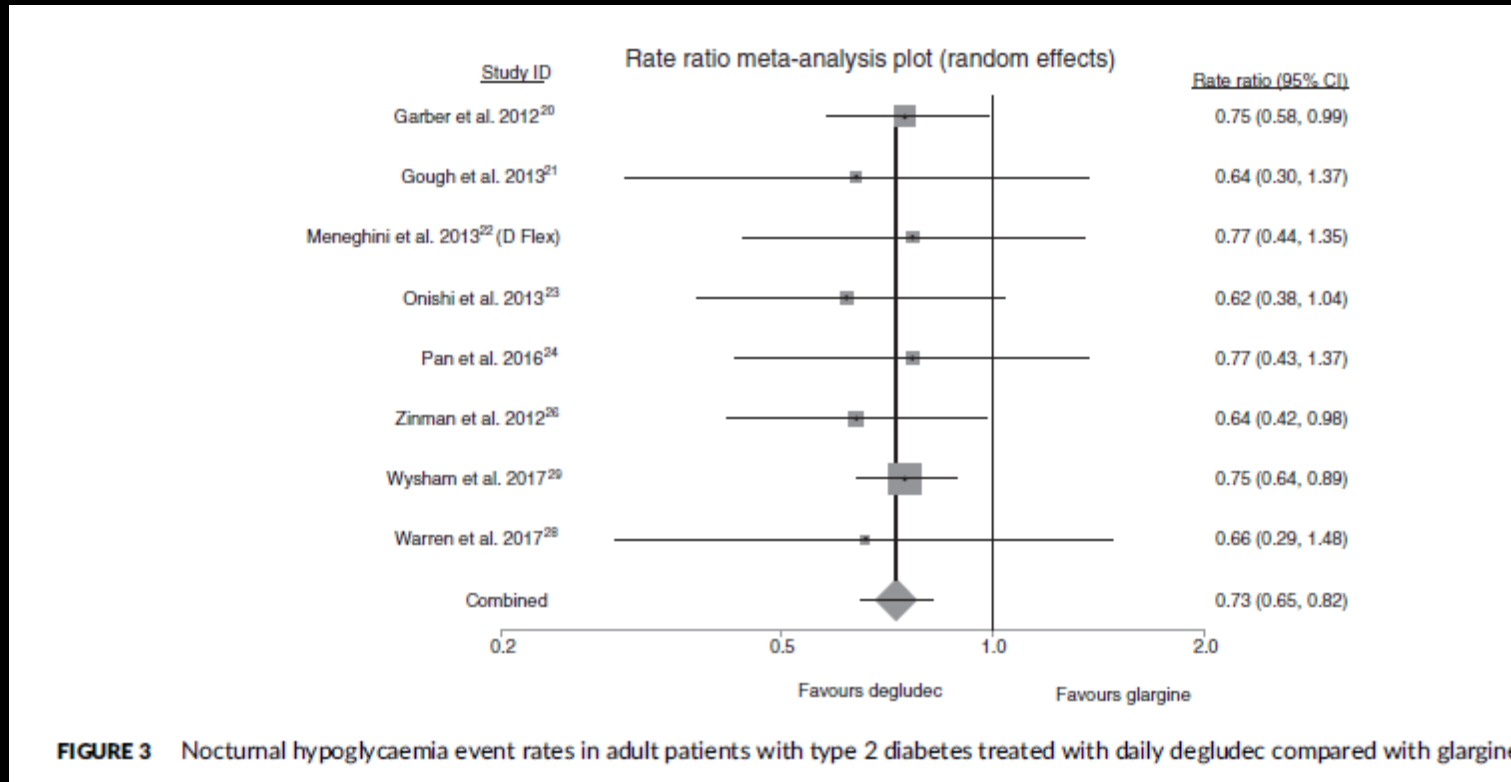


# Glargine U300



- >24 hour action
- MAX pen
  - doses up to 160 units per shot
  - 2 units increments, but only the 4 unit multiples shown on dial
- IHS National formulary

# Nocturnal Hypo – Degludec vs. Glargine U100



# Insulin Degludec U100 & U200



- FlexTouch device does not dial out, may be easier for arthritic hands
- >24 hour duration
- U100 pen
  - Max dose 80 units
  - Shows 2 unit increments
- U200 pen
  - Shows 4 unit increments
  - Max dose 160 units

# Aspart U100



Insulin aspart-szjj - not interchangeable

# Lispro U100 & U200

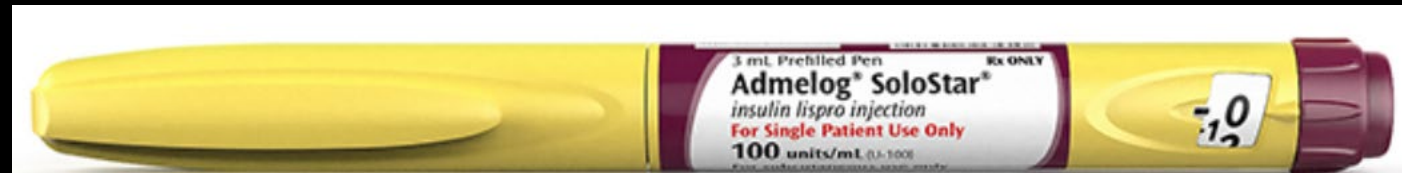


- Both only go to 60 units per dose

# Lispro Biosimilars NOT Interchangeable



Lyumejev – aspart aabc, U100 and U200



# Ultra-Rapid Insulins



Any clinically important advantages? – Likely not

# Highly Concentrated Insulin (U500)



- Indicated if total daily dose of insulin is over 200 units per day
- This is a stand alone insulin – it provides both bolus AND basal insulin
- It is dosed twice or three times daily
- Pen dials to increments of 5 units
- **MUST BE SURE** that patient is actually taking insulin

# Combo Pen Examples



# Insulin Considerations

- Pens vs. vials
- Arthritic hands – fatter pens?
- Site rotation
- When the numbers don't match the prescription and story
  - Proper storage
  - Proper insulin technique - twisting vs. pushing?
  - How many doses are being missed?
  - Is it even being taken?

# Test Your Knowledge

- Patient on 100 units of glargine U300 insulin once a day
  - Blood sugar is rarely below 250
  - A1c = 14.2%
- 
- This may be a patient that you take all the way back to the beginning..... trust your gut.....

# Continuous Glucose Monitoring



# Insulin Pumps



# IHS Food for Thought....

- OK to have more than one “version” on local formulary
- Make P&T Committee allowances for pharmacists to interchange biosimilars (with or with-interchangeable status) based on formulary & Point of Sale reimbursements
- When interchanging insulins – there is a chance of the patient taking BOTH. Patient education is of utmost importance.
- MUST keep up to date on new forms available and occasionally check pricing as these are EVER changing.

# And now...your questions...

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