

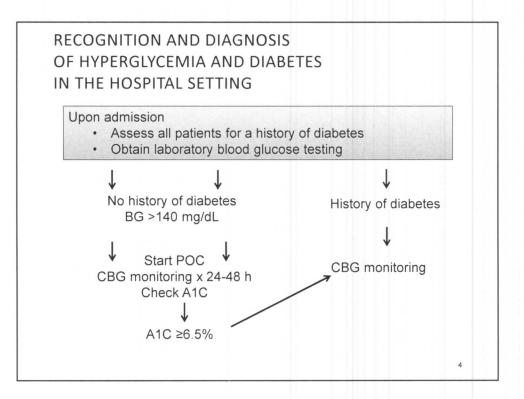
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RECOGNITION AND DIAGNOSIS OF HYPERGLYCEMIA AND DIABETES IN THE HOSPITAL SETTING

All patients

- Assess for history of diabetes
- Test BG (using laboratory method) on admission independent of prior diagnosis of diabetes
- Patients without a history of diabetes
 - BG >140 mg/dL: Monitor with POC testing for 24-48 h
 - BG >140 mg/dL: Ongoing POC testing
 - Patients receiving therapies associated with hyperglycemia (eg, corticosteroids): monitor with POC testing for 24-48 h
 - BG >140 mg/dL: continue POC testing for duration of hospital stay
- Patients with known diabetes or with hyperglycemia
 - Test A1C if no A1C value is available from past 2-3 months

BG, blood glucose; POC, point of care. Moghissi ES, et al. *Endocrine Pract.* 2009;15:353-369. Umpierrez GE, et al. *J Clin Endocrinol Metab.* 2012;97:16-38.



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A1C FOR DIAGNOSIS OF DIABETES IN THE HOSPITAL

- Implementation of A1C testing can be useful
 - Assist with differentiation of newly diagnosed diabetes from stress hyperglycemia
 - Assess glycemic control prior to admission
 - Facilitate design of an optimal regimen at the time of discharge
- A1C <a>6.5% indicates diabetes

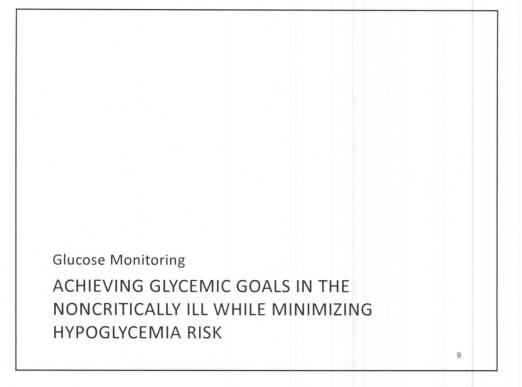
Moghissi ES, et al. *Endocrine Pract.* 2009;15:353-369. Umpierrez GE, et al. *J Clin Endocrinol Metab.* 2012;97:16-38.

GLYCEMIC GOALS FOR NONCRITICALLY ILL PATIENTS

INPATIENT GLYCEMIC MANAGEMENT: DEFINITION OF TERMS

Hospital hyperglycemia	Any BG >140 mg/dL	
Stress hyperglycemia	Elevations in blood glucose levels that occur in patients with no prior history of diabetes and A1C levels that are not significantly elevated (<6.5%)	
A1C value >6.5%	Suggestive of prior history of diabetes	
Hypoglycemia	ia Any BG <70 mg/dL	
Severe hypoglycemia	Any BG <40 mg/dL	

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MONITORING GLYCEMIA IN THE NONCRITICAL CARE SETTING

POC testing

- Preferred method for guiding ongoing glycemic management of individual patients
- Timing of glucose measures should match patient's nutritional intake and medication regimen
- Recommended schedules for POC testing
 - Before meals and at bedtime in patients who are eating
 - Every 4-6 h in patients who are NPO or receiving continuous enteral or parenteral nutrition

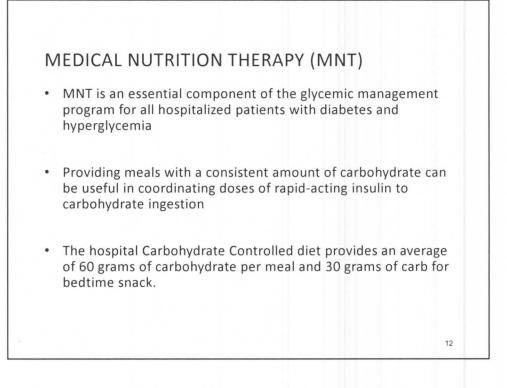
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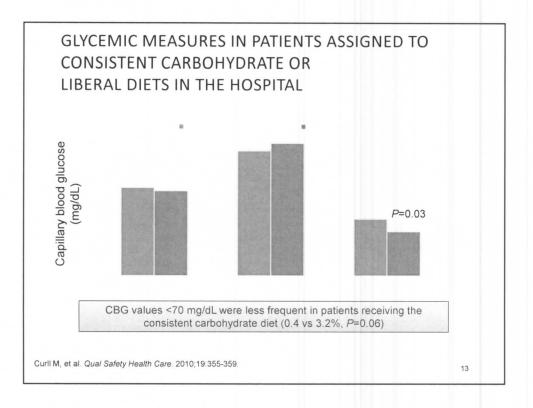
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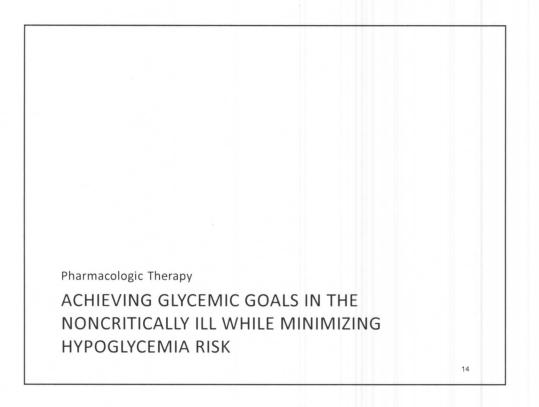
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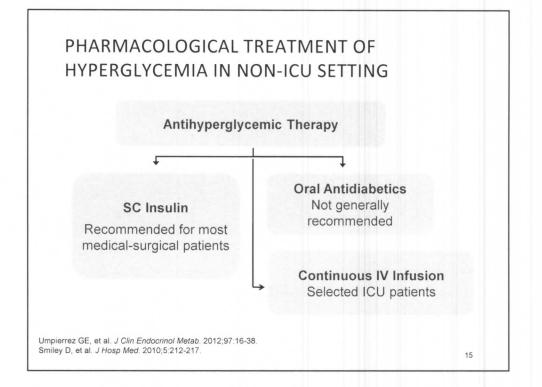
Hospital Diet

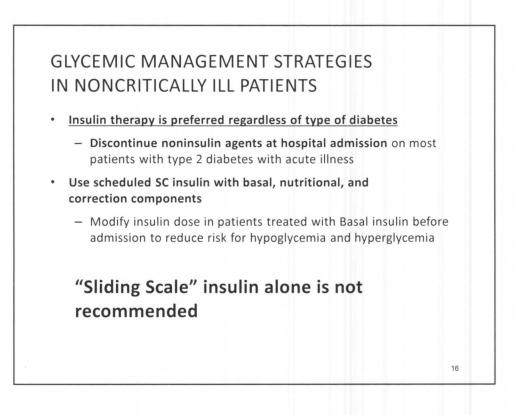
ACHIEVING GLYCEMIC GOALS IN THE NONCRITICALLY ILL WHILE MINIMIZING HYPOGLYCEMIA RISK











INPATIENT MANAGEMENT OF HYPERGLYCEMIA: MANAGING SAFETY CONCERNS

- Both undertreatment and overtreatment of hyperglycemia create safety concerns
- Areas of risk
 - Changes in carbohydrate or food intake
 - Changes in clinical status or medications
 - Failure to adjust therapy based on BG patterns
 - Prolonged use of SSI as monotherapy
 - Poor coordination of BG testing with insulin administration and meal delivery
 - Poor communication during patient transfers
 - Errors in order writing and transcription

NONINSULIN THERAPIES IN THE HOSPITAL Time-action profiles of oral agents can result in delayed ٠ achievement of target glucose ranges in hospitalized patients Sulfonylureas are a major cause of prolonged hypoglycemia Metformin is contraindicated in patients with decreased renal ٠ function, use of iodinated contrast dye, and any state associated with poor tissue perfusion (CHF, sepsis) Thiazolidinediones are associated with edema and CHF . α -Glucosidase inhibitors are weak glucose-lowering agents ٠ · Pramlintide and GLP-1 receptor agonists can cause nausea and exert a greater effect on postprandial glucose Insulin therapy is the preferred approach

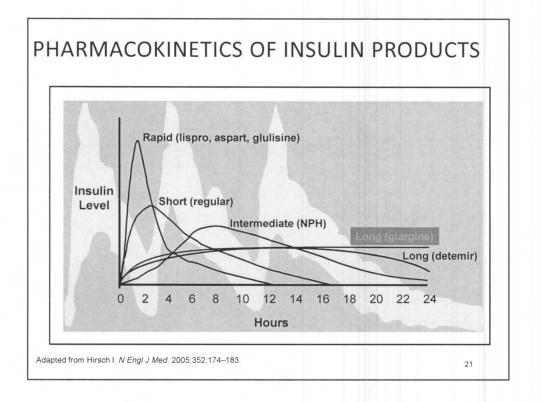
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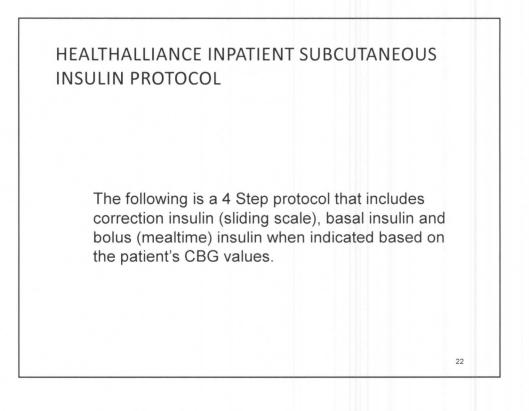
Basal insulin	•Detemir (Levemir), glargine (Lantus), NPH Blunts the rise in blood glucose following nutritional intake (meals, IV dextrose, enteral/parenteral nutrition)	
Nutritional (prandial) insulin		
Correction insulin (sliding scale)	Corrects hyperglycemia due to mismatch of nutritional intake and/or illness-related factors and scheduled insulin administration	

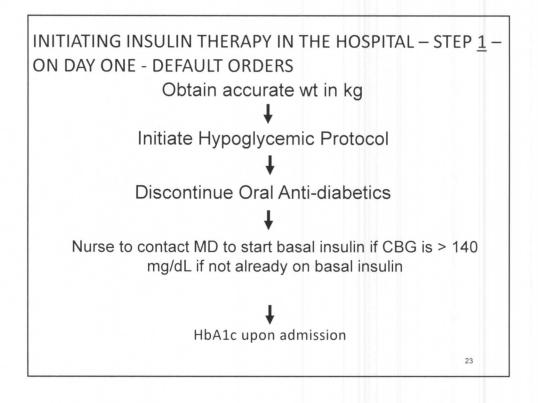
PHARMACOKINETICS OF INSULIN PREPARATIONS

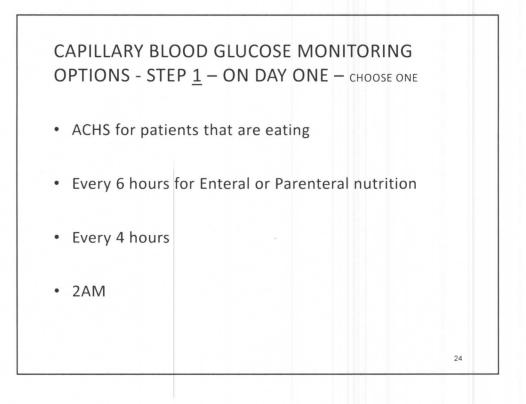
Insulin	Onset	Peak	Duration
Nutritional			
Rapid-acting analog (aspart, glulisine, lispro)	5-15 min	1-2 hours	4-6 hours
Regular	30-60 min	2-3 hours	6-10 hours
Basal			
Detemir	2 hours	Relatively peakless	16-24 hours
Glargine	2-4 hours	Relatively peakless	20-24 hours
NPH	2-4 hours	4-10 hours	12-18 hours

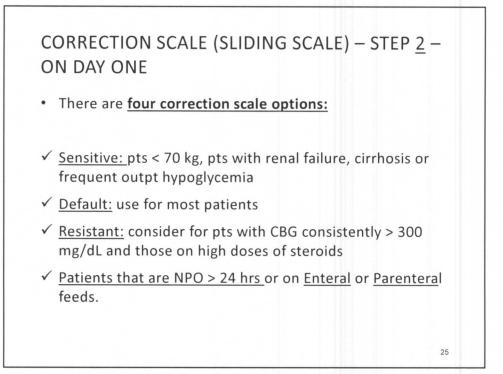
Hirsch I. *N Engl J Med.* 2005;352:174-183. Porcellati F, et al. *Diabetes Care*. 2007;30:2447-2552.

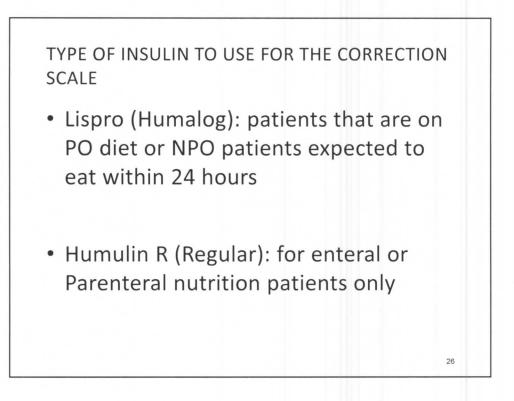


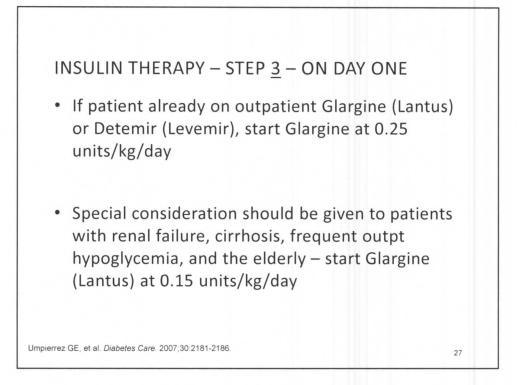


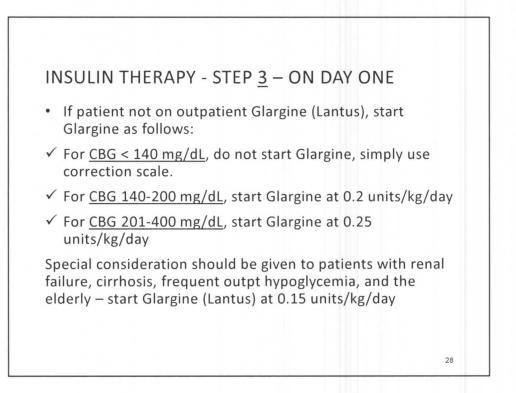












INSULIN THERAPY - STEP <u>4</u> – ON DAY TWO (OR EARLIER IF 3 CONSECUTIVE CBG VALUES ARE > 200) FOR PATIENTS THAT ARE EATING

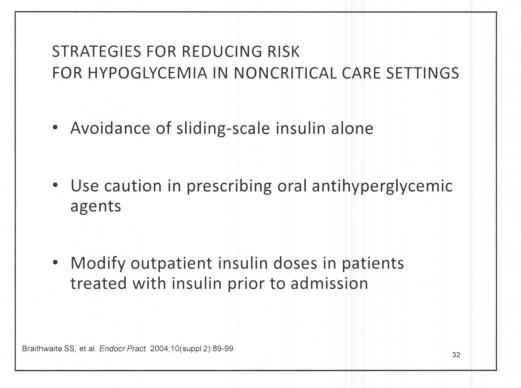
- For patients receiving Basal Insulin and Correction (sliding scale) insulin and the <u>mean daily</u> CBG > 200 mg/dL or <u>three consecutive</u> CBG are > 200 mg/dL, add bolus (mealtime) insulin.
- This is given with food <u>in addition</u> to correction Insulin

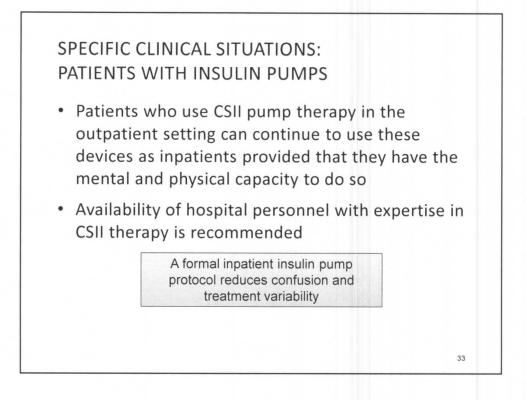
INSULIN THERAPY - STEP $\underline{4}$ – ON DAY TWO (OR EARLIER IF 3 CONSECUTIVE CBG VALUES ARE > 200 MG/DL) FOR PATIENTS THAT ARE EATING

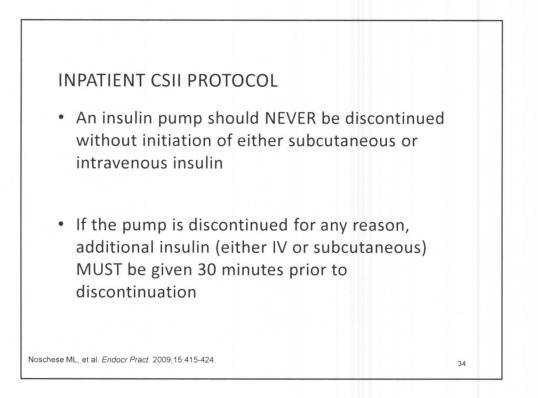
- For CBG 201 400 mg/dL, start Lispro (Humalog) at 0.25 units/kg/day <u>divided into 3 mealtime</u> boluses.
- Special consideration should be given to patients with renal failure, cirrhosis, frequent outpt hypoglycemia, and the elderly – start Lispro (Humalog) at 0.15 units/kg/day <u>divided into 3</u> <u>mealtime</u> boluses.

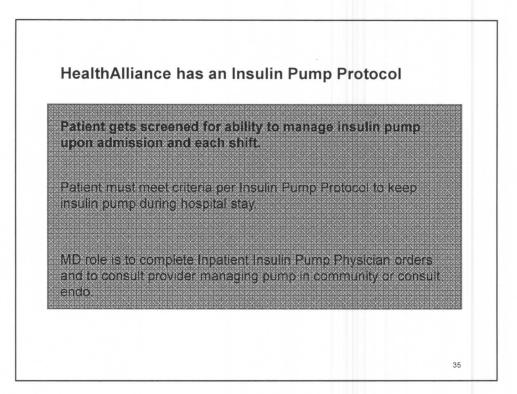
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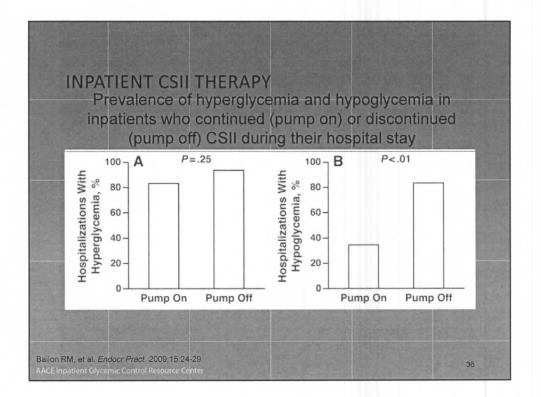
to de bla	<i>P</i> value		
/ariable	Univariate Analysis	Multivariate Analysis*	
Age	<0.001	<0.001	
GFR <60 mL/s	0.005	0.11	
rDD ≥0.5 U/kg	0.006	0.31	
Previous insulin use	<0.001	0.02	
nsulin regimen basal-bolus vs SSI)	<0.001	0.001	

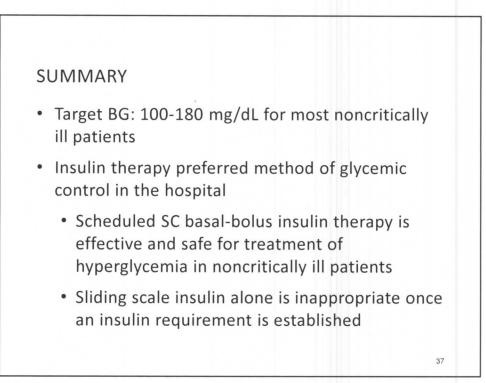














MANAGEMENT OF HYPERGLYCEMIA IN THE NONCRITICAL CARE SETTING

I acknowledge that I have read the training module (PowerPoint under "Reference Material" on the Management of Hyperglycemia in the Noncritical Care Setting.

Print Name:

Signature: _____ Date:

Please return this page with your application.