

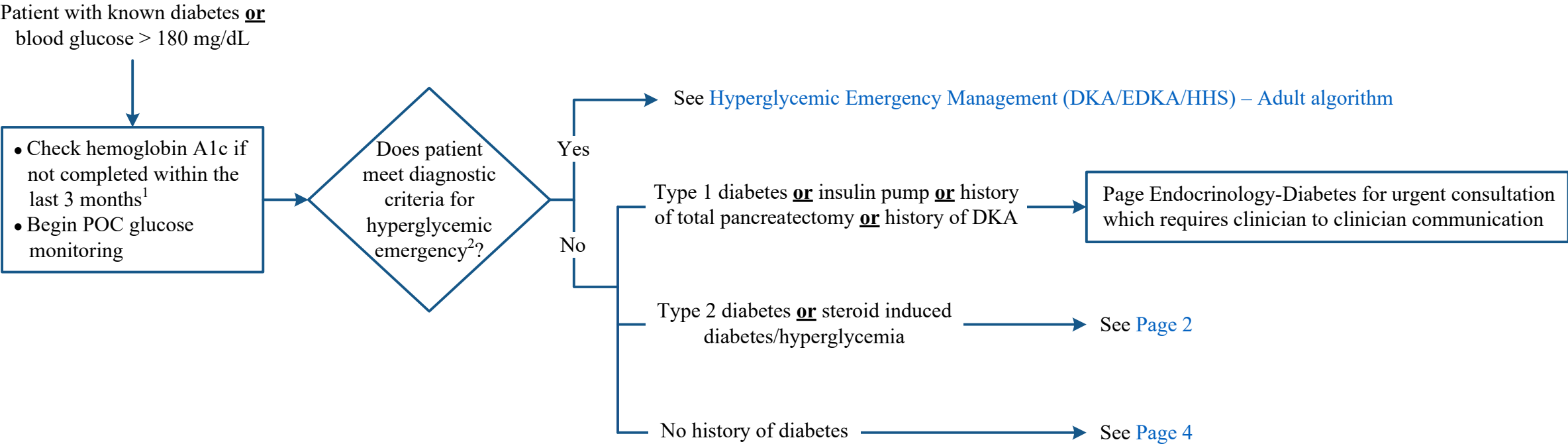
Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient’s care. This algorithm should not be used to treat pregnant women.

**Note:** Insulin dose adjustments should be made based on the individual patient’s glucoses. Refer to the [Hypoglycemia Management algorithm](#), as indicated.

PRESENTATION

INITIAL EVALUATION

TREATMENT



DKA = diabetic ketoacidosis  
EDKA = euglycemic diabetic ketoacidosis  
HHS = hyperosmolar hyperglycemic state  
POC = point of care

<sup>1</sup> Hemoglobin A1c may be inaccurate if recent blood transfusion within the past three months or severe anemia  
<sup>2</sup> Diagnostic criteria:  
DKA: blood glucose > 250 mg/dL, arterial or venous pH < 7.3, bicarbonate < 15 mEq/L, beta hydroxybutyrate (BHB) > 3 mmol/L, osmolality < 320 mosm/kg  
EDKA: blood glucose ≤ 250 mg/dL, arterial or venous pH < 7.3, bicarbonate < 15 mEq/L, BHB > 3 mmol/L, osmolality < 320 mosm/kg  
[**Note:** Blood glucose may be lower than expected in patients on SGLT-2 inhibitors (e.g., canagliflozin, dapagliflozin, empagliflozin, ertugliflozin)]  
DKA with HHS: blood glucose > 600 mg/dL, arterial or venous pH < 7.3, bicarbonate < 15 mEq/L, BHB > 3 mmol/L, osmolality ≥ 320 mosm/kg  
HHS: blood glucose > 600 mg/dL, arterial or venous pH ≥ 7.3, bicarbonate ≥ 15 mEq/L, BHB ≤ 3 mmol/L, osmolality ≥ 320 mosm/kg

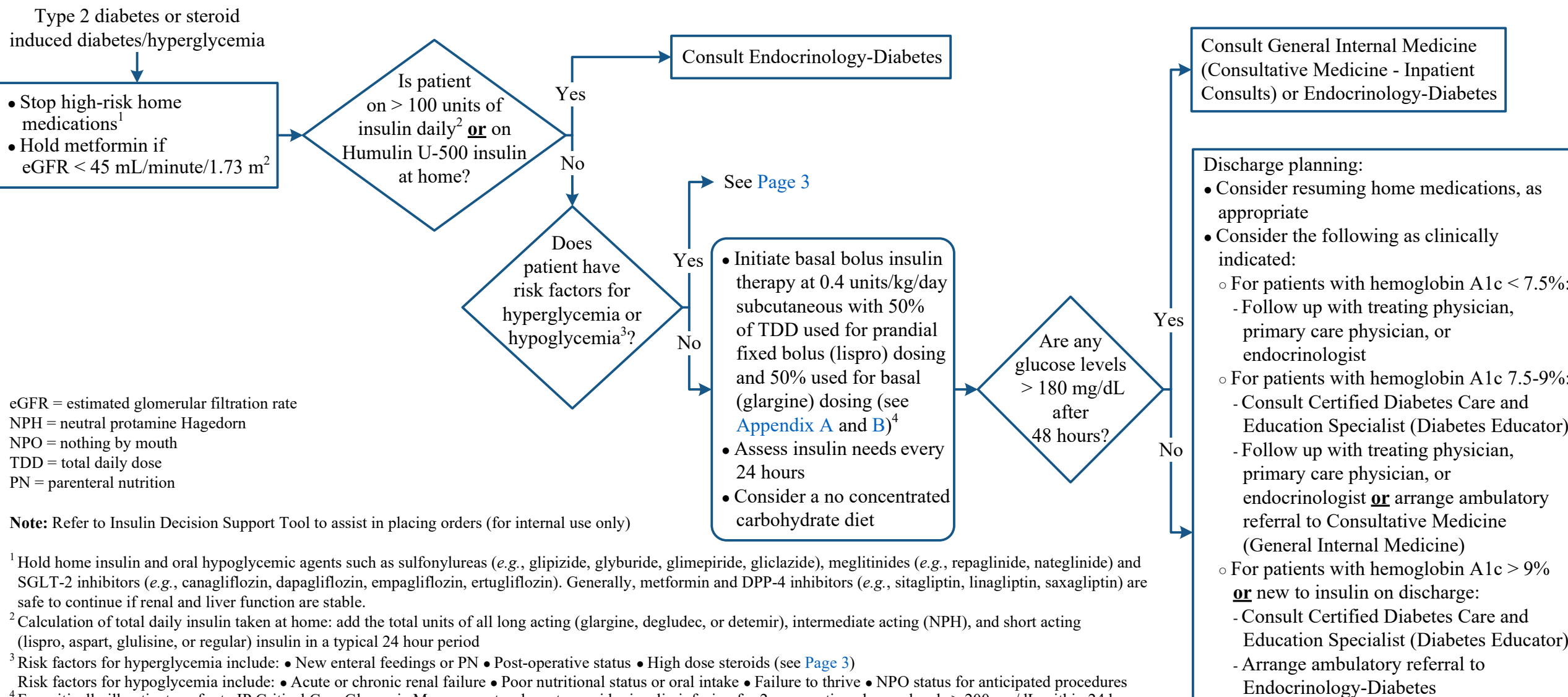
Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

**Note:** Insulin dose adjustments should be made based on the individual patient's glucoses. Refer to the [Hypoglycemia Management algorithm](#), as indicated.

## PRESENTATION

## ASSESSMENT

## TREATMENT



eGFR = estimated glomerular filtration rate  
NPH = neutral protamine Hagedorn  
NPO = nothing by mouth  
TDD = total daily dose  
PN = parenteral nutrition

**Note:** Refer to Insulin Decision Support Tool to assist in placing orders (for internal use only)

<sup>1</sup> Hold home insulin and oral hypoglycemic agents such as sulfonylureas (e.g., glipizide, glyburide, glimepiride, gliclazide), meglitinides (e.g., repaglinide, nateglinide) and SGLT-2 inhibitors (e.g., canagliflozin, dapagliflozin, empagliflozin, ertugliflozin). Generally, metformin and DPP-4 inhibitors (e.g., sitagliptin, linagliptin, saxagliptin) are safe to continue if renal and liver function are stable.

<sup>2</sup> Calculation of total daily insulin taken at home: add the total units of all long acting (glargine, degludec, or detemir), intermediate acting (NPH), and short acting (lispro, aspart, glulisine, or regular) insulin in a typical 24 hour period

<sup>3</sup> Risk factors for hyperglycemia include: • New enteral feedings or PN • Post-operative status • High dose steroids (see [Page 3](#))

Risk factors for hypoglycemia include: • Acute or chronic renal failure • Poor nutritional status or oral intake • Failure to thrive • NPO status for anticipated procedures

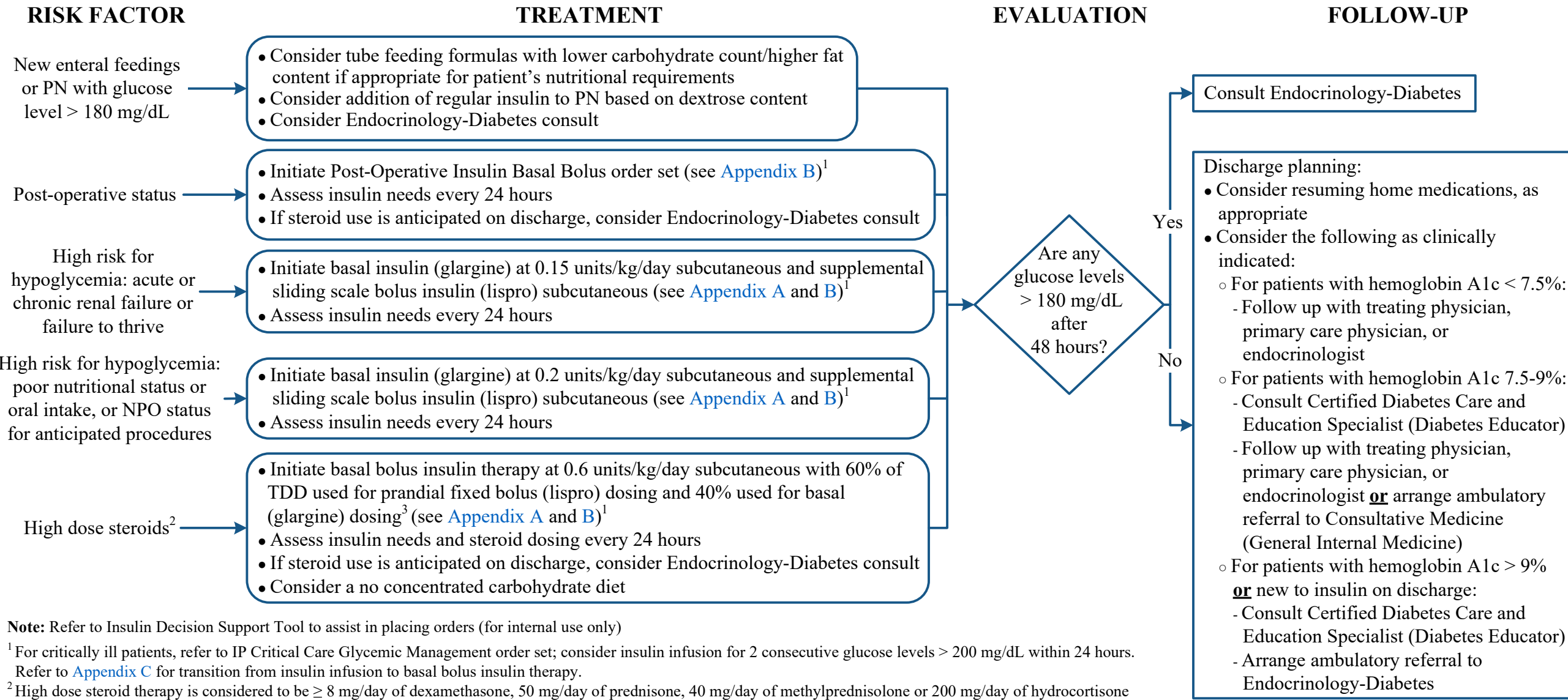
<sup>4</sup> For critically ill patients, refer to IP Critical Care Glycemic Management order set; consider insulin infusion for 2 consecutive glucose levels > 200 mg/dL within 24 hours.

Refer to [Appendix C](#) for transition from insulin infusion to basal bolus insulin therapy.

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient’s care. This algorithm should not be used to treat pregnant women.

**Note:** Insulin dose adjustments should be made based on the individual patient’s glucoses. Refer to the [Hypoglycemia Management algorithm](#), as indicated.

## TYPE 2 DIABETES OR STEROID INDUCED DIABETES/HYPERGLYCEMIA



**Note:** Refer to Insulin Decision Support Tool to assist in placing orders (for internal use only)

<sup>1</sup> For critically ill patients, refer to IP Critical Care Glycemic Management order set; consider insulin infusion for 2 consecutive glucose levels > 200 mg/dL within 24 hours. Refer to [Appendix C](#) for transition from insulin infusion to basal bolus insulin therapy.

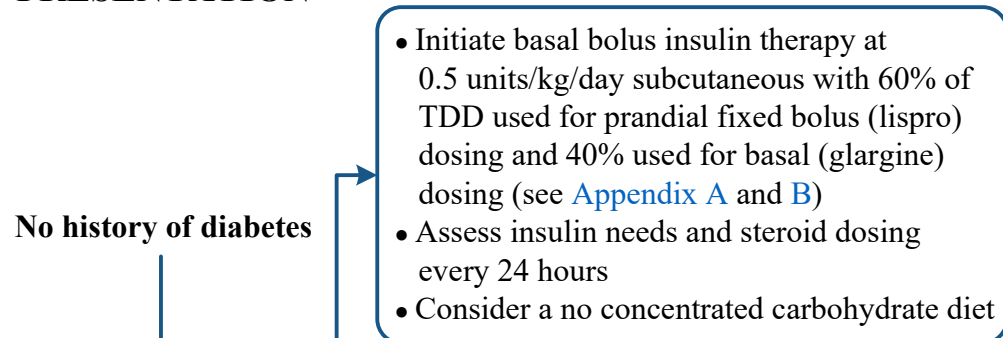
<sup>2</sup> High dose steroid therapy is considered to be ≥ 8 mg/day of dexamethasone, 50 mg/day of prednisone, 40 mg/day of methylprednisolone or 200 mg/day of hydrocortisone

<sup>3</sup> In recurrent admissions for chemotherapy containing steroids, please review last Endocrinology-Diabetes note for more specific insulin dose recommendations and use as indicated

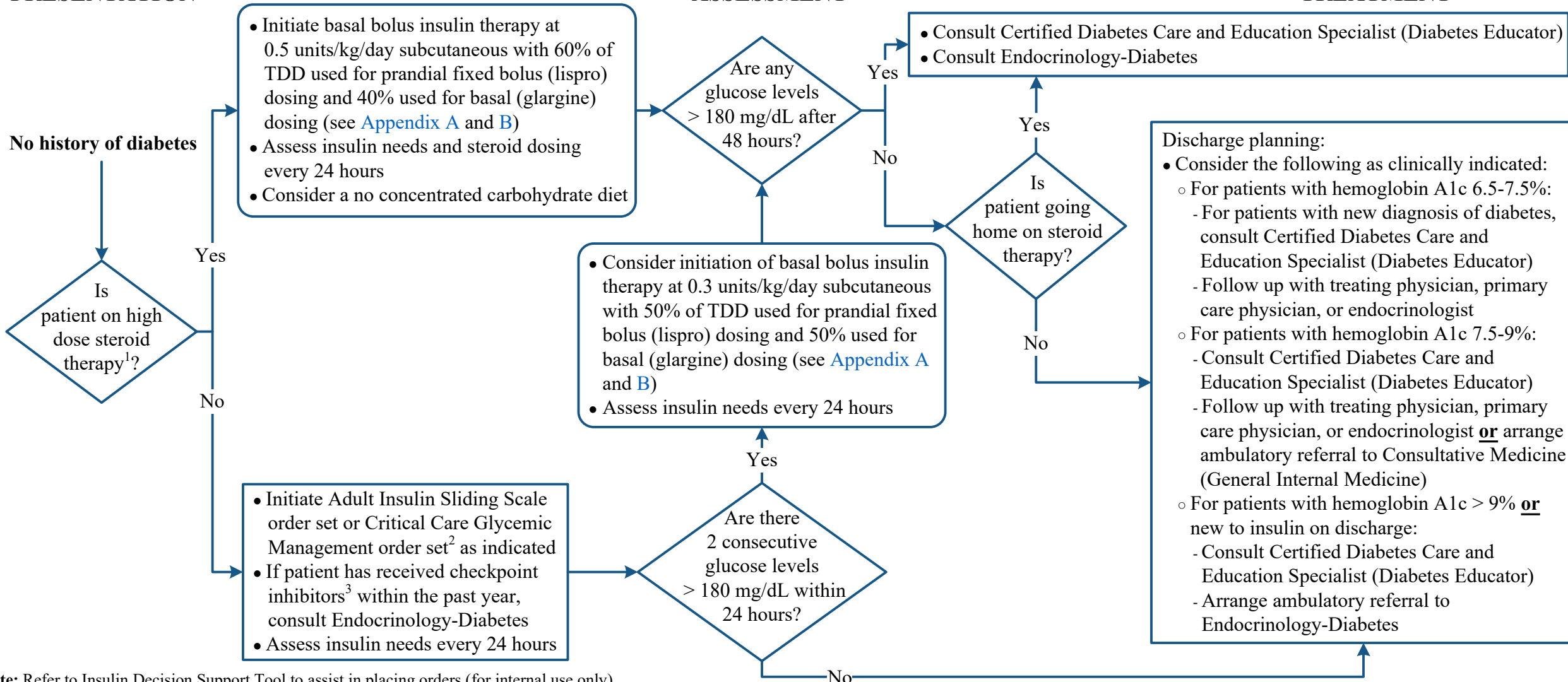
Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

**Note:** Insulin dose adjustments should be made based on the individual patient's glucoses. Refer to the [Hypoglycemia Management algorithm](#), as indicated.

## PRESENTATION



## ASSESSMENT



**Note:** Refer to Insulin Decision Support Tool to assist in placing orders (for internal use only)

<sup>1</sup> High dose steroid therapy is considered to be ≥ 8 mg of dexamethasone, 50 mg of prednisone, 40 mg of methylprednisolone or 200 mg of hydrocortisone per day

<sup>2</sup> For critically ill patients, consider insulin infusion for 2 consecutive glucose levels > 200 mg/dL within 24 hours. Refer to [Appendix C](#) for transition from insulin infusion to basal bolus insulin therapy.

<sup>3</sup> Checkpoint inhibitors: nivolumab, pembrolizumab, durvalumab, atezolizumab, and related drugs. Patients with recent exposure to checkpoint inhibitors are at risk for DKA and should be evaluated for new onset type 1 diabetes mellitus.

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

## APPENDIX A: Common Insulin Types and Frequency

Fast Acting Insulin	Dose Frequency
Lispro (Humalog® or Lyumjev® <sup>1</sup> )	Before meals or every 4 hours
Aspart (Novolog® or Fiasp®) <sup>1</sup>	Before meals or every 4 hours
Glulisine (Apidra®) <sup>1</sup>	Before meals or every 4 hours
Regular insulin (Novolin®-R/Humulin®-R)	Before meals or every 6 hours
Long Acting Insulin	
Glargine (Lantus®/Basaglar®/Toujeo®/Semglee® <sup>1</sup> )	Daily or every 12 hours
Detemir (Levemir®)	Daily or every 12 hours
Degludec (Tresiba®) <sup>1</sup>	Daily
Intermediate Acting Insulin	
NPH (Novolin®-N/Humulin®-N)	Every 12 hours
Mixed Insulin	
70/30, 75/25 <sup>1</sup> , 50/50 <sup>1</sup> (mixes of NPH and a fast acting insulin)	Every 12 hours or every 6 hours with continuous tube feedings

<sup>1</sup> Not currently on MD Anderson Formulary

## APPENDIX B: Basal Bolus Insulin Terms

- **Bolus** insulin refers to a dose of fast acting insulin. This is typically comprised of **prandial** insulin which is scheduled to compensate for the carbohydrate content of a meal and **supplemental** (or sliding scale) insulin to correct hyperglycemia. Bolus insulin is most effective when given before meals, but supplemental insulin alone can be scheduled for patients who are not eating or are high risk for hypoglycemia.
- **Basal** insulin refers to a dose of long acting insulin given 1 or 2 times daily. These insulins absorb slowly to help maintain stable glucose levels.
- **Supplemental** insulin is dosed based on either weight or total daily insulin requirement
- A **basal/bolus** insulin regimen uses both types of insulin to recreate a physiologic pattern of insulin release. This regimen is more effective for most patients than sliding scale supplemental insulin only. Most patients need about half of their insulin as basal and half as bolus. Patients on high doses of steroids will often need more bolus insulin.



Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

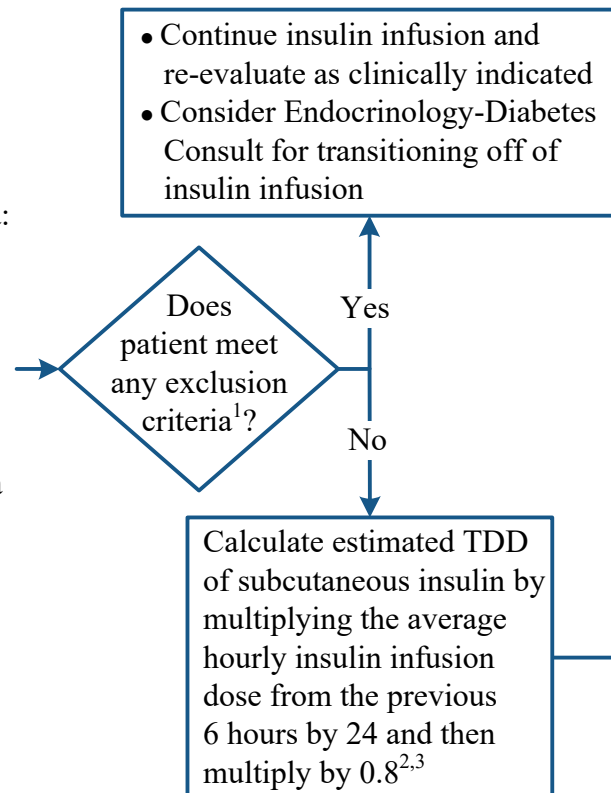
## APPENDIX C: Critical Care Insulin Infusion Transition

**Note:** This does NOT apply to patients with DKA, EDKA, or HHS  
(see [Hyperglycemic Emergency Management \(DKA/EDKA/HHS\) – Adult algorithm](#))

Clinical judgement may supersede exact calculation of total daily dose if patient's clinical status has rapidly changed during transition period.

Patient on ICU insulin infusion **and** meets **ALL** of the following criteria:

- Not in DKA/EDKA/HHS
- Maintained on insulin infusion  $\geq 6$  hours
- Controlled blood glucose ( $\geq 3$  blood glucoses  $< 180$  mg/dL in past 6 hours)
- Steady insulin infusion doses for a minimum of 6 hours (not varying by  $> 2$  units/hour)



<sup>1</sup> Exclusion criteria include: Currently receiving vasopressor therapy or parenteral nutrition (PN) **or** steroid dose fluctuating  $> 20\%$  in the past 24 hours **or** insulin infusion dose  $> 5$  units/hour **or** enteral feedings not at goal rate

<sup>2</sup> Example: Average hourly infusion dose from the previous 6 hours = 3 units/hour; multiply by 24 = 72 units; multiply by 0.8 = 57.6 units, rounded up to 58 units = TDD; see [Appendix D](#) for basal/bolus insulin calculation tables

<sup>3</sup> Consider decreasing the TDD by 20% for patients with renal dysfunction; Example: TDD = 58 units decreased by 20% = 46 units

**Note:** After transitioning off of insulin infusion, continue to evaluate glycemic control and adjust management as clinically indicated

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

## APPENDIX D: Basal/Bolus Insulin Dosing Calculation Tables

**Note:** • Clinical judgement may supersede exact calculation of total daily dose if patient’s clinical status has rapidly changed during transition period  
• Consider dose reduction in patients with renal dysfunction

NPO <u>or</u> Eating < 50% of Meals				
Average Hourly Dose of Insulin Infusion	First Dose of Basal Insulin (Glargine)	Maintenance Dose of Basal Insulin (Glargine) <sup>1</sup>	Dose of Prandial (Pre-Meal) Scheduled Short/Rapid Acting	Regular Insulin Supplemental Sliding Scale
Less than or equal to 1 unit/hour	9 units once	5 units every 12 hours	-	Low dose sliding scale every 6 hours
2 units/hour	19 units once	10 units every 12 hours	-	Low dose sliding scale every 6 hours
3 units/hour	28 units once	14 units every 12 hours	-	Medium dose sliding scale every 6 hours
4 units/hour	38 units once	19 units every 12 hours	-	Medium dose sliding scale every 6 hours
5 units/hour	48 units once	24 units every 12 hours	-	High dose sliding scale every 6 hours

Eating ≥ 50% of Meals <u>or</u> Intermittent Enteral Feedings				
Average Hourly Dose of Insulin Infusion	First Dose of Basal Insulin (Glargine)	Maintenance Dose of Basal Insulin (Glargine) <sup>1</sup>	Dose of Prandial (Pre-Meal) <sup>2</sup> Scheduled Lispro	Lispro Insulin Supplemental Sliding Scale
Less than or equal to 1 unit/hour	9 units once	5 units every 12 hours	3 units three times daily before meals	Low dose sliding scale three times daily before meals
2 units/hour	19 units once	10 units every 12 hours	6 units three times daily before meals	Low dose sliding scale three times daily before meals
3 units/hour	28 units once	14 units every 12 hours	9 units three times daily before meals	Medium dose sliding scale three times daily before meals
4 units/hour	38 units once	19 units every 12 hours	12 units three times daily before meals	Medium dose sliding scale three times daily before meals
5 units/hour	48 units once	24 units every 12 hours	16 units three times daily before meals	High dose sliding scale three times daily before meals

<sup>1</sup> Maintenance dose of basal insulin (glargine) should start no sooner than 12 hours AFTER first dose. Hold for glucose < 120 mg/dL.

<sup>2</sup> Hold for glucose < 100 mg/dL or if NPO

Continued on next page

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

## APPENDIX D: Basal/Bolus Insulin Dosing Calculation Tables - continued

**Note:** • Clinical judgement may supersede exact calculation of total daily dose if patient’s clinical status has rapidly changed during transition period  
• Consider dose reduction in patients with renal dysfunction

Continuous Enteral Feedings				
Average Hourly Dose of Insulin Infusion	First Dose of Basal Insulin (Glargine)	Maintenance Dose of Basal Insulin (Glargine) <sup>1</sup>	Dose of Scheduled Regular Insulin <sup>2</sup>	Regular Insulin Supplemental Sliding Scale
Less than or equal to 1 unit/hour	9 units once	5 units every 12 hours	3 units every 6 hours	Low dose sliding scale every 6 hours
2 units/hour	19 units once	10 units every 12 hours	4 units every 6 hours	Low dose sliding scale every 6 hours
3 units/hour	28 units once	14 units every 12 hours	7 units every 6 hours	Medium dose sliding scale every 6 hours
4 units/hour	38 units once	19 units every 12 hours	9 units every 6 hours	Medium dose sliding scale every 6 hours
5 units/hour	48 units once	24 units every 12 hours	12 units every 6 hours	High dose sliding scale every 6 hours

<sup>1</sup> Maintenance dose of basal insulin (glargine) should start no sooner than 12 hours AFTER first dose. Hold for glucose < 120 mg/dL.

<sup>2</sup> Hold for glucose < 100 mg/dL or if enteral feedings interrupted and notify ICU Team



Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

## SUGGESTED READINGS

- Chang, L. L., Umpierrez, G. E., & Inzucchi, S. E. (2022). Management of hyperglycemia in hospitalized, non-critically ill adults. *The New England Journal of Medicine*, 387(11), 1040-1042. <https://doi.org/10.1056/NEJMc1de2204691>
- Honarmand, K., Sirimatuross, M., Hirshberg, E. L., Bircher, N. G., Agus, M. S. D., Carpenter, D. L., . . . Jacobi, J. (2024). Society of Critical Care Medicine guidelines on glycemic control for critically ill children and adults 2024. *Critical Care Medicine*, 52(4), e161–e181. <https://doi.org/10.1097/CCM.00000000000006174>
- Kirk, J. K., & Oldham, E. C. (2010). Hyperglycemia management using insulin in the acute care setting: Therapies and strategies for care in the non-critically III patient. *Annals of Pharmacotherapy*, 44(7-8), 1222-1230. <https://doi.org/10.1345/aph.1M695>
- Kodner, C., Anderson, L., & Pohlgeers, K. (2017). Glucose management in hospitalized patients. *American Family Physician*, 96(10), 648-654. Retrieved from <https://www.clinicalkey.com/#!/content/playContent/1-s2.0-S0002838X17303660>
- Korytkowski, M. T., Muniyappa, R., Antinori-Lent, K., Donihi, A. C., Drincic, A. T., Hirsch, I. B., . . . Umpierrez, G. E. (2022). Management of hyperglycemia in hospitalized adult patients in non-critical care settings: An Endocrine Society clinical practice guideline. *The Journal of Clinical Endocrinology & Metabolism*, 107(8), 2101-2128. <https://doi.org/10.1210/clinem/dgac278>
- Lansang, M. C., & Umpierrez, G. E. (2016). Inpatient hyperglycemia management: A practical review for primary medical and surgical teams. *Cleveland Clinic Journal Of Medicine*, 83(suppl 1), S34-S43. <https://doi.org/10.3949/ccjm.83.s1.06>
- Maynard, G. A., Childers, D., Holdych, J., Kendall, H., Hoag, T., & Harrison, K. (2017). Improving glycemic control safely in non-critical care patients: A collaborative systems approach in nine hospitals. *The Joint Commission Journal on Quality and Patient Safety*, 43(4), 179-188. <https://doi.org/10.1016/j.jcjq.2017.01.003>
- Umpierrez, G. E., Smiley, D., Zisman, A., Prieto, L. M., Palacio, A., Ceron, M., . . . Mejia, R. (2007). Randomized study of basal-bolus insulin therapy in the inpatient management of patients with type 2 diabetes (RABBIT 2 trial). *Diabetes Care*, 30(9), 2181-2186. <https://doi.org/10.2337/dc07-0295>

Disclaimer: *This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson’s specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.*

## DEVELOPMENT CREDITS

This practice consensus statement is based on majority opinion of the Hyperglycemic Management experts at the University of Texas MD Anderson Cancer Center for the patient population. These experts included:

### Core Development Team Leads

Conor Best, MD (Endocrine Neoplasia and HD)  
Sonya Khan, MD (Endocrine Neoplasia and HD)  
Sonali Thosani, MD (Endocrine Neoplasia and HD)

### Workgroup Members

Veronica Brady, PhD, MSN, RN (Endocrine Neoplasia and HD)  
Vivian Crowder, MSN, APRN, FNP-C (Endocrine Neoplasia and HD)  
Lakeisha Day, DMSc, PA-C (Nocturnal Program)  
Carmen Escalante, MD (General Internal Medicine)  
Wendy Garcia, BS♦  
Abigayle Harrington, BSN, RN (Nursing Post Anesthesia Care Unit)  
Michelle Horng, PharmD (Pharmacy Clinical Programs)  
Kwame Koom-Dadzie, MD (Hospital Medicine)  
Victor Lavis, MD (Endocrine Neoplasia and HD)  
Celia Levesque, MS, APRN, FNP, CNS-BC (Endocrine Neoplasia and HD)  
Sally Mathews, MSN, APRN (Hospital Medicine)  
Michael Tanner Moser, PharmD (Pharmacy Clinical Programs)  
Hadeel Sahar, MBBCH (Hospital Medicine)  
Jolyn Taylor, MD (Gynecological Oncology & Reproductive Medicine)  
Jeena Varghese, MD (Endocrine Neoplasia and HD)  
Sigi Varghese, MS, APRN (Endocrine Neoplasia and HD)  
Khanh Vu, MD (General Internal Medicine)  
Mary Lou Warren, DNP, APRN, CNS-CC♦  
Jessica Williams, MS, PA-C (Endocrine Neoplasia and HD)  
Mara Wilson, MS, APRN (Endocrine Neoplasia and HD)

♦Clinical Effectiveness Development Team