



CULTIVATING A CULTURE OF CARING FOR OLDER ADULTS

1 in 4 Older adults have DIABETES!

Glycemic control goals should be individualized based on comorbid conditions, life expectancy and risks/benefits of treatment.

Ask patients about common geriatric syndromes (falls, incontinence, cognitive impairment, depression, etc.), as they are more common in patients with Diabetes.

Remember it takes approximately 5-10 years of poor glycemic control to develop microvascular complications.

Avoid hypoglycemia. Hypoglycemia symptoms include: irritability, confusion, tachycardia, impaired concentration, shakiness, etc.

Remember the Glycemic Control Goals for Older Adults

| Health Status | HbA1c Goal |
|---|------------|
| Healthy with few chronic conditions | <7.5% |
| Multiple coexisting conditions or limited life expectancy | <8.0% |
| End stage chronic disease or significant dementia | <8.5% |

Adapted from the American Diabetes Association. Standards of Medical Care in Diabetes 2017. Diabetes Care 2017; 40 (Suppl.1):S99-s104

START WITH MONOTHERAPY UNLESS:

- A1C is $\geq 9\%$, consider **DUAL THERAPY**
- A1C is $\geq 10\%$, blood glucose is $\geq 300\text{mg/dL}$, or patient is markedly symptomatic, consider **COMBINATION INJECTABLE THERAPY**.

MONOTHERAPY: METFORMIN LIFESTYLE MANAGEMENT

| | |
|---------------------|--------------------|
| Efficacy | High |
| Hypo Risk | Low Risk |
| Weight | Neutral/Loss |
| Side Effects | GI/Lactic Acidosis |
| Costs | Low |

If A1C target not achieved after approximately 3 months of monotherapy, proceed to 2-drug combination (order not meant to denote any specific preference – choice dependent on a variety of patient and disease-specific factors.)

DUAL THERAPY: METFORMIN + LIFESTYLE MANAGEMENT

| | Sulfonylurea | Thiazolidine-dione | DPP-4 Inhibitor | SGLT2 Inhibitor | GLP-1 Receptor Agonist | Insulin (Basal) |
|---------------------|---------------|--------------------|-----------------|----------------------|------------------------|-----------------|
| Efficacy | High | High | Intermediate | Intermediate | High | Highest |
| Hypo Risk | Moderate Risk | Low Risk | Low Risk | Low Risk | Low Risk | High Risk |
| Weight | Gain | Gain | Neutral | Loss | Loss | Gain |
| Side Effects | Hypoglycemia | Edema, HF, fxs | Rare | GU, dehydration, fxs | GI | Hypoglycemia |
| Costs | Low | Low | High | High | High | High |

If A1C target not achieved after approximately 3 months of monotherapy, proceed to 2-drug combination (order not meant to denote any specific preference – choice dependent on a variety of patient and disease-specific factors.)

TRIPLE THERAPY: METFORMIN + LIFESTYLE MANAGEMENT

| Sulfonylurea + | | Thiazolidine-dione + | | DPP-4 Inhibitor + | | SGLT2 Inhibitor + | | GLP-1 Receptor Agonist + | | Insulin (Basal) + | |
|----------------|----------|----------------------|----------|-------------------|---------|-------------------|----------|--------------------------|---------|-------------------|----------|
| or | TZD | or | SU | or | SU | or | SU | or | SU | or | TZD |
| or | DPP-4-i | or | DPP-4-i | or | TZD | or | TZD | or | TZD | or | DPP-4-i |
| or | SGLT2-i | or | SGLT2-i | or | SGLT2-i | or | DPP-4-i | or | SGLT2-i | or | SGLT2-i |
| or | GLP-1-RA | or | GLP-1-RA | or | Insulin | or | GLP-1-RA | or | Insulin | or | GLP-1-RA |
| or | Insulin | or | Insulin | | | | | | | | |

If A1C target not achieved after approximately 3 months of triple therapy and patient (1) on oral combination, move to basal insulin or GLP-2-RA, (2) on GLP-1-RA, add basal insulin, or (3) on optimally titrated basal insulin, add GLP-1-RA or mealtime insulin. Metformin therapy should be maintained, while other oral agents may be discontinued on an individual basis to avoid unnecessarily complex or costly regimen (i.e., adding a fourth antihyperglycemic agent)

COMBINATION INJECTABLE THERAPY LIFESTYLE MANAGEMENT

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SELECTED ORAL DIABETIC AGENTS

| Drug Class/Name | MOA | A1C reduction | Clinical Pearls |
|--|---|---------------|--|
| Biguanine (<i>Metformin</i>) | Decreases Glucose production | Up to 2% | <ul style="list-style-type: none"> • Max dose 2,550 • OK to use Crcl 30-50ml/min • GI SE minimized with slow titration; Weight loss |
| Sulfonylureas (<i>Glipizide, Glimepiride, Glyburide</i>) | Stimulate insulin release | Up to 2% | <ul style="list-style-type: none"> • Low durability, Don't use Glyburide. Hypoglycemia, weight gain. Stop when insulin is started |
| DPP4I's | Prolong Action of endogenous Incretins, Enhancing 1st phase of insulin response | 0.6-1% | <ul style="list-style-type: none"> • Trajenta: No renal adjustments, Weight neutral; No hypoglycemia; SE: GI, joint pain • Pancreatitis, Onglyza increases HF hospitalizations |
| Thiazolidinediones (<i>Pioglitazone, rosiglitazone</i>) | Insulin resistance reducers | Up to 1.5% | <ul style="list-style-type: none"> • Risk of heart failure, weight gain, edema, arthralgia, LFT increase. Inc HDL and Dec TG Actos may reduce risk of stroke/MI with pre-existing disease |

SELECTED DIABETIC MEDICATIONS

| Drug Class/Name | MOA | A1C reduction | Clinical Pearls |
|---|---|---------------|---|
| Meglitinides (<i>Nateglinide, repaglinide</i>) | Increase insulin secretion | 0.5-1% | <ul style="list-style-type: none"> • Short duration of action. PRN use, weight gain, hypoglycemia |
| SGLT2 Inhibitors (<i>Canagliflozin, dapagliflozin, empagliflozin</i>) | Decrease glucose reabsorption from kidney And increase urinary glucose excretion | 0.5-1.5% | <ul style="list-style-type: none"> • Weight loss, lower bp, no hypoglycemia, hyperkalemia, UTI's, mycotic genital infections, ketoacidosis. Renal dosing Possible CV benefit |
| α-Glucosidase Inhibitors (<i>Acarbose, miglitol</i>) | Decrease glucose absorption | Less than 0.5 | <ul style="list-style-type: none"> • GI disturbances. Difficult to take |
| GLP-1 receptor agonists (<i>Exenatide, liraglutide, albiglutide, dulaglutide</i>) | Incretin mimetics | 0.7-1% | <ul style="list-style-type: none"> • Less hypoglycemia, weight loss; Pancreatitis, thyroid cancer, injectable |

SELECTED ORAL DIABETIC AGENTS

| Preparation | Onset | Peak | Duration | Doses per day |
|--|---------|-----------|----------|---------------|
| Rapid-acting Insulin glulisine (Apidra) | 20 min | 0.5-1.5 h | 3-4 h | 3 |
| Insulin lispro (Humalog) | 15 min | 0.5-1.5 h | 3-4 h | 3 |
| Insulin aspart (NovoLog) | 30 min | 1-3 h | 3-5 h | 3 |
| Short acting insulin (e.g., Humulin, Novolin) | 0.5-1 h | 2-3 h | 5-8 h | 3 |
| Intermediate-acting NPH (e.g., Humulin, Novolin) | 1-1.5 h | 4-12 h | 24 h | 1-2 |

*Insulin preparations are 100U/ml and come 10cc vials and 3cc pens
70/30 Formulations with NPH & rapid or short acting insulins

LONG-ACTING INSULIN PRODUCTS

| Agent | Onset | Peak | Duration |
|-----------------------|---|--|-------------------------------------|
| Onset | 1.1 hour | 1.1 to 2 hours | Develops over 6 hours |
| Peak | No significant peak | | |
| Duration | 10.8 to > 24 hours | Mean 7.6 to > 24 hours | > 24 hours |
| Administration | SC once daily | SC once or twice daily (12hr) | SC once daily |
| Formulation | 100 units/mL 10 ml vial, 3 ml SoloStar pen | 100 units/ml 10 ml vial, 3 ml FlexTouch | 300 units/ml 1.5 ml SoloStar pen |
| Compatibility | Do not mix with other insulins or dilute | | |