## Clinician Decision Aid

# **Guideline Directed Medical Therapy for T2D**



SGLT2 inhibitors and incretin mimetics are first-line treatments for T2D in patients with cardiorenal disease. *This aid is meant to support the use of GLP-1/GIP receptor agonists and SGLT2 inhibitors, alongside your own clinical judgement, to guide patient-centered diabetes treatment.*<sup>1</sup>

ALL PATIENTS WITH T2D

Off Label

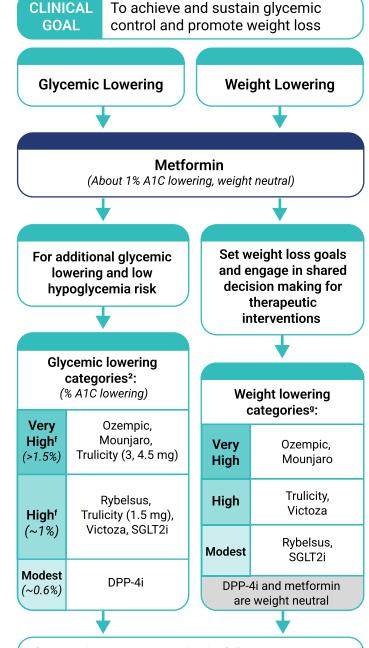
Recommend lifestyle change with reduced carb intake and weight loss if indicated. Promote diabetes self management education and support (DSMES) upon diagnosis.

## **CLINICAL** To reduce cardiorenal risk independent **GOAL** of A1C in high risk patients Heart Established ASCVD<sup>a</sup> **CKD<sup>c</sup>** Failure<sup>b</sup> On maximum tolerated dose of **Shared Decision** ACEI/ARB Making (GLP-1 RA or Farxiga SGLT2i): Jardiance Ozempic/Rybelsus, Trulicity, Victoza Farxiga<sup>e</sup> Invokana Invokana<sup>d</sup> Jardiance<sup>e</sup> Steglatro<sup>d</sup> Invokana, Jardiance or Brenzavvy Ozempic If A1C is above target, consider the following: · Additional agents9 (based on glycemic, weight lowering, and comorbidity needs) Initiation of a CGM Referral or re-referral to DSMES FDA Labeled (preferred) No Evidence - low cost

a. Consider use in HIGH risk patients: ADA Standards gives a weaker recommendation for use given cardiovascular outcomes trial (CVOT) data were not powered for primary analysis in this subgroup. This subgroup includes: age ≥ 55 with 2 or more risk factors (e.g, obesity, hypertension, smoking, hyperlipidemia, albuminuria).

(see footnote f)

- **b.** Heart failure includes: HFpEF (preserved EF) and HFrEF (reduced EF, ≤40%).
- c. Diagnosis of CKD (eGFR<60) and/or presence of albuminuria: Determine after repeating measures of GFR and albuminuria twice, 3 months apart.
- d. Invokana in HF: Data for use from secondary outcomes of CVOT T2D trials. Steglatro in HF: Data for use from a non-inferiority trial, no proven benefit.
- e. When using for HF and/or CKD protection: Irrespective of T2D diagnosis, dose is 10mg daily, and can be used at renal functions below renal cutoffs for glycemic lowering (See chart on next page).
- **f. Brenzavvy:** Low cost option (\$50-60/month) if guideline directed SGLT2i is not affordable. No evidence to support added ASCVD/CKD benefits.



#### If A1C is above target, consider the following:

- Additional agents<sup>g</sup> (based on glycemic, weight lowering, and comorbidity needs)
- Initiation of a CGM
- · Referral or re-referral to DSMES
- g. Additional Glycemic Agents: Insulin (very high), sulfonylureas and pioglitazone (high) are effective glycemic lowering agents, but are not included because they are associated with weight gain. These agents also cause hypoglycemia.

## **Prescribing Reminders and Recommendations**

	Incretin Mimetics	SGLT2 Inhibitors	
Contraindications	Personal or family history of medullary thyroid cancer or MEN-2-syndrome, pregnancy, lactation, or allergy to medication.	T1D, very low carbohydrate diet (less than 50g/day), pregnancy, lactation, or allergy to medication.	
Precautions	History of:     Pancreatitis     Severe GI disease (gastroparesis)     Gallbladder disease	On dialysis: Contraindicated.h  High risk for amputation (e.g. active foot ulcers, prior amputations, or severe peripheral artery disease). See MCT2D Precautions Guide.  If A1c greater than 10%: Consider alternative medication to lower glucose to avoid excess glucosuria.	
Considerations for impaired renal function	Monitor for volume depletion: Use caution in patients with GI side effects or acute illness (e.g. vomiting, diarrhea, dehydration) when initiating or escalating doses.	For glycemic lowering benefit - Avoid if: • eGFR < 45: Farxiga and Steglatro • eGFR < 30: Jardiance, Invokana, and Brenzavvy	
Considerations for medication adjustments	If A1C is less than 9% and on:  • Basal insulin: lower by 10%²  • Prandial insulin: lower by 30-40%²	If A1C less than 8.5% and patient is on:  • Insulin: lower total daily dose by 10-20% - Avoid insulin discontinuation to minimize risk of euglycemic DKA  • Sulfonylurea: Discontinue or lower dose by 50%	
Key counseling points	<ul> <li>GI mitigating strategies and expected time for improvement</li> <li>Sensitizing doses have negligible impact on glucose</li> </ul>	<ul><li>Hydration</li><li>Sick day management</li><li>Perineal hygiene</li><li>Blood pressure monitoring</li></ul>	
Drug interactions	DPP-4 Inhibitors (all incretin mimetics): No benefit from combination  Oral contraceptives (Tirzepatide only): Add backup method for 4 weeks after initiation and for 4 weeks after each dose escalation.	Diuretic therapy - Monitor for:  Volume depletion  Hypotension with other antihypertensive therapy Adjustment upon initiation of SGLT2i is not required.	
Prior to surgery	Daily agents: Hold on the day of surgery³ Weekly agents: Hold at least 7 days prior³ Bridge with other glucose lowering medications if necessary. Use shared decision-making with surgery and anesthesia to hold incretin mimetic based on patient risk.	<b>Hold for 4 days prior.</b> Resume after full oral intake is established.	

**h.** Per KDIGO guidelines, contraindicated given lack of safety studies in those receiving dialysis. Studies are ongoing in the ESRD population.

## SGLT2i Renal Cutoffs:

For Patients with CKD/HF and T2D

	eGFR 25 - 45	eGFR 20 - 30	
	Farxiga	Jardiance	
CKD	10 mg daily	10 mg daily	
Heart Failure	(can continue if eGFR < 25)	(can continue if eGFR < 20)	
On Dialysis: <b>Contraindicated</b> <sup>h</sup>			

## **Medication Dosing Guide**



michmed.org/Zxn7B

## **Patient Medication Handouts**



michmed.org/bmx5B

## References

- 1. ADA 2025 doi: 10.2337/dc25-S009
- 2. Van Dril 2022 doi: 10.1016/j.ahjo.2022.100163
- 3. Kindel 2024 doi: 10.1016/j.soard.2024.08.033

