

CHRONIC KIDNEY DISEASE WITH TYPE 2 DIABETES AND CARDIOVASCULAR DISEASE



PRIMARY CARE
MANAGEMENT OF

CKD



CVD is the leading cause of death in patients with CKD, who face an increased risk of adverse CV outcomes



Primary care plays a vital role in preventing CKD progression and reducing adverse CV occurrences



Many patients with T2D and CKD remain at residual risk for CKD progression and CV events despite receiving standard treatment

Monitoring for residual risk

What?



Comorbidity

- CKD monitoring
- CVD and dyslipidemia
- Diabetes



What to measure

- eGFR, uACR, urinalysis
- Blood pressure, cardiovascular risk stratification, lipid status
- Blood glucose, HbA1c

Look out for



- Dose adjustments based on eGFR
- Hyperkalemia if RAS inhibition is included
- CKD/CVD risk factors (e.g. anemia, bone disease, heart failure)

When?



- The KDIGO heatmap can be used as a tool to guide frequency of monitoring

Albuminuria categories Description and range		
A1	A2	A3
Normal to mildly increased	Moderately increased	Severely increased
<30 mg/g <3 mg/mmol	30–299 mg/g 3–29 mg/mmol	≥300 mg/g ≥30 mg/mmol

GFR categories (mL/min/1.73m ²) Description and range	G1	Normal or high	≥90	Screen (1)	Treat (1)	Treat and refer (3)
	G2	Mildly decreased	60–89	Screen (1)	Treat (1)	Treat and refer (3)
	G3a	Mildly to moderately decreased	45–59	Treat (1)	Treat (2)	Treat and refer (3)
	G3b	Mildly to severely decreased	30–44	Treat (2)	Treat and refer (3)	Treat and refer (3)
	G4	Severely decreased	15–29	Treat and refer (3)	Treat and refer (3)	Treat and refer (4+)
	G5	Kidney failure	<15	Treat and refer (4+)	Treat and refer (4+)	Treat and refer (4+)

Numbers: Indicate how often (per year) you should be screening or monitoring. Monitor, treat, or refer: Indicates the recommended course of action. CKD is classified based on Cause (C), GFR (G), Albuminuria (A).

Low risk (if no other markers of kidney disease, no CKD)

Moderately increased risk

High risk

Very high risk

Management of T2D + CKD + CVD

Aim of treatment: Slow CKD progression and prevent adverse CV events



Additional risk factor control

Lipid management
Glycemic control
Blood pressure control

Additional drugs with heart and kidney protection

GLP-1 RA (T2D)
ns-MRA (T2D)
Antiplatelet therapies (ASCVD)

First-line drug therapy

Metformin (T2D)
SGLT2i (T2D)
RAS blockade (HTN)
Statin

Lifestyle and self-management

Diet
Exercise
Weight
Smoking cessation

Management pearls

- SGLT2i should be initiated or continued for those with CKD and T2D when eGFR ≥ 20 mL/min/1.73m² and can be continued after initiation at lower eGFR levels
- Metformin should be initiated/continued when eGFR ≥ 30 mL/min/1.73 m²
- RAS inhibitors (ACEi or ARB) should be initiated or continued as first-line therapy at the maximal tolerated dose
- Intensification of statin therapy is recommended for secondary prevention in patients with established CVD
- Aspirin should be used for lifelong secondary prevention in patients with established CVD
- Long-acting GLP-1 RA are recommended for patients who do not achieve individualized glycemic targets despite use of metformin and SGLT2i
- Additional medications with cardiorenal protective effects, such as non-steroidal MRAs* can be used in patients at residual risk despite standard treatment

When to refer to nephrology

Consult/refer to nephrology if:



KFRE score rises above 3–5%



Developing a treatment plan and primary care practitioner not confident in the recommended first-line treatment



Unexplained decline in eGFR (≥ 5 mL/min/1.73m²) over 12 months or sudden decline over days to weeks



Unexplained significant albuminuria or hematuria



Persistent hyperkalemia



Resistant hypertension



Recurring kidney stones



Hereditary kidney disease

*Finerenone is currently the only nonsteroidal MRA with proven clinical kidney and cardiovascular benefits

References: de Boer IH, et al. *Diabetes Care* 2022; 45: 3075–3090; ISN & KDIGO. ISN-KDIGO early screening booklet. Accessible at: https://www.theisn.org/wp-content/uploads/2023/02/ISN_KDIGO_EarlyScreeningBooklet_PRINT_Updated.pdf (accessed December 2023); Kidney Disease: Improving Global Outcomes (KDIGO) CKD Work Group. KDIGO 2023 Clinical Practice Guideline for the evaluation and management of chronic kidney disease. July 2023 draft. *Kidney Disease: Improving Global Outcomes (KDIGO) Diabetes Work Group. Kidney Int* 2022; 102: S1–S127; Marx N, et al. *Eur Heart J* 2023; 44: 4043–4140.

Acronyms: ACEi, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; CKD, chronic kidney disease; CV, cardiovascular; CVD, cardiovascular disease; eGFR, estimated glomerular filtration rate; GLP-1 RA, glucagon-like peptide-1 receptor agonist; HTN, hypertension; KFRE, kidney failure risk equation; MRA, mineralocorticoid receptor antagonist; RAS, renin-angiotensin system; SGLT2i, sodium-glucose cotransporter-2 inhibitor; T2D, type 2 diabetes; uACR, urine albumin-creatinine ratio.