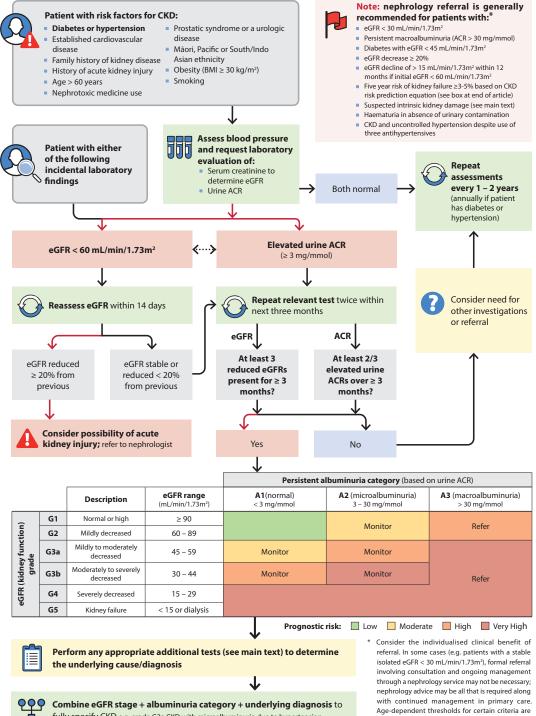
bpac^{nz} Quick Clinical Knowledge



B-QuiCK: Chronic kidney disease (CKD)

Summary: detection and diagnosis of patients with CKD in primary care



fully specify CKD e.g. grade G3a CKD with microalbuminuria due to hypertension

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currently being debated in the literature, however,

no consensus has been reached.

Summary: management of patients with CKD in primary care

Weight loss	• Reduce BMI to at least \leq 30 kg/m ² with an ideal target of \leq 25 kg/m ²
Increase exercise participation	 Tailor recommendation to the patient; general targets include 150–300 min/week moderate intensity exercise or 75–150 min/week vigorous exercise Advise strength/resistance training on at least two days/week, if possible
Nutrition	 Recommend a balanced diet emphasising fruits, vegetables, nuts, low-fat dairy products, whole grains and fish (e.g. <u>DASH diet</u>); reduce sodium intake and avoid trans fats, processed meats, refined carbohydrates and sweetened beverages (N.B. water should be the main fluid but avoid over-consumption) Avoid high levels of protein intake but do not advise a protein-restricted diet unless under dietician or nephrologist supervision
Reducing alcohol intake	 Advise a maximum of one or two standard drinks daily, with at least two alcohol free days per week, if applicable
Smoking cessation	Encourage cessation, if applicable

Discuss healthy lifestyle changes with consideration of individual or cultural barriers to achieving these.

Initiate and optimise blood pressure management

- Aim for a target systolic blood pressure of < 130 mmHg for most patients
- An ACE inhibitor or ARB is first line. Titrate to the maximum (approved) tolerated dose while monitoring for acute reductions in eGFR and hyperkalaemia (within one to two weeks initially, then less frequently once targets have been achieved). If hyperkalaemia becomes an issue, consider other options of controlling serum potassium before withholding or reducing dose of the ACE inhibitor/ARB, e.g. reducing dietary potassium intake, stopping/switching beta-blocker (if applicable).
- If treatment escalation is required, add another antihypertensive, e.g. calcium channel blocker or loop diuretic. Thiazide diuretics are less effective in patients with worsening eGFR.

Initiate and optimise management of hyperglycaemia if the patient has diabetes

- Aim for a target HbA₁ of < 53 mmol/mol for most patients
- Start with metformin; the maximum tolerated dose is dependent on kidney function (see NZF)
- Add a SGLT-2 inhibitor (empagliflozin) if treatment escalation is required (Special Authority criteria applies); also consider a SGLT-2 inhibitor in patients with CKD without diabetes (but this is **not** funded)
- If further glucose-lowering treatment is need, consider adding a GLP-1 receptor agonist (preferred, but dual SGLT-2 inhibitor/GLP-1 receptor agonist treatment is not funded), or other options, e.g. vildagliptin, pioglitazone (unless the patient has heart failure), a sulfonylurea or insulin

Consider other pharmacological treatments

- Assess whether the patient is taking any other potentially nephrotoxic medicines and consider if dose adjustments, switching or discontinuation is required
- Discuss the benefits of statin treatment with all patients, particularly those with high triglyceride and low HDL-C levels
- Consider long-term aspirin treatment for patients with CKD and established CVD or in those with a high risk of atherosclerotic events (unless they have an increased bleeding risk)

Provide advice about managing acute illness

 Advise patients with CKD to temporarily stop the following medicines if they experience acute illness (using the Mnemonic: SAD MANS): sulfonylureas, ACE-inhibitors, diuretics, metformin, ARBs, NSAIDs and SGLT-2 inhibitors

Establish a monitoring plan

- Patients with established CKD should have their eGFR, ACR and blood pressure reviewed at least annually, in addition to other relevant laboratory investigations, e.g. serum electrolytes, lipids, HbA_{1c} (if diabetes present). More frequent review is appropriate for patients with greater eGFR impairment or proteinuric CKD, e.g. three to six monthly (see full text for specific criteria).
- Patients should be referred to nephrology if progressive CKD or other changes indicating significant kidney function deterioration are identified

It is strongly recommended to review the original resource at your convenience for full details of recommendations and evidence. See full article here: <u>bpac.org.nz/2022/ckd.aspx</u>