

The hidden threat: Primary care-driven strategies for earlier chronic kidney disease detection in Australia

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Background

Chronic kidney disease (CKD) is a growing and underdiagnosed public health challenge. General practitioners are central to early detection, yet missed opportunities persist. Evidence suggests up to 90% of stage 3 CKD remains unrecorded, despite simple, cost-effective screening and treatment that slows progression. With a projected 39% rise in the number of Australians with stage 3–5 CKD by 2034, immediate action is required.

Objective

Three roundtable meetings were conducted across Australia with 35 primary care providers and kidney experts to identify barriers and supporting solutions to increase CKD detection and diagnosis.

Discussion

Six themes were identified to improve early CKD detection: (1) education, audits and supportive technologies; (2) guideline harmonisation and a unified risk calculator; (3) automated pathology integrated into practice software, linked to the *CKD management in primary care* handbook and decision tools; (4) defined CKD codes reflecting staging and cause; (5) multidisciplinary communication and models of care; and (6) government policy and advocacy to raise public awareness.

CHRONIC KIDNEY DISEASE (CKD) represents a significant yet largely hidden health burden. Approximately three in four Australian adults are at risk,^{1,2} and 1.7–2 million have reduced kidney function,^{1,3} although fewer than 10% are aware of their diagnosis.^{1,3} The insidious nature of the disease allows up to 90% of kidney function to be lost before symptoms manifest.^{1,3} Although Australia's prevalence 1 in 10) aligns with other high-income countries,^{3,4} Aboriginal and Torres Strait Islander peoples experience double this rate (1 in 5)^{1,3} and have among the highest prevalence globally.^{5,6}

The escalating burden of CKD is a public health crisis, costing Australia over \$9.9 billion annually.^{7,8} Each year CKD claims 20,000 lives^{1,2} and leads to 3300 Australians requiring renal replacement therapy² – equivalent to 55 deaths and 14 new people progressing to kidney failure each day.² Diabetes and cardiovascular disease often co-exist, and people with CKD are 20 times more likely to die from cardiovascular causes than to progress to kidney failure.^{1,9–11} By 2032, Australia's CKD burden is expected to surge, with a 85.9% increase in dialysis and a 39% rise in stage 3–5 CKD.¹² These projections underscore the urgency for earlier identification and timely diagnosis to improve patient outcomes.

General practitioners (GPs) play a pivotal part in early detection and intervention, which can slow the rate of kidney deterioration¹ and, if implemented early,

delay the progression to kidney failure by more than 15 years.^{13–15} To support this, Kidney Health Australia (KHA) updated the *CKD management in primary care* handbook (hereafter the CKD Handbook) in 2024, including a revised algorithm for initial detection and diagnosis of CKD in individuals at increased risk of developing CKD (Box 1, Figure 1).¹ Screening is simple and cost effective,¹⁶ involving a urine test for albuminuria, a blood test for estimated glomerular filtration rate (eGFR) – calculated from serum creatinine – and blood pressure (BP) measurement.¹ This targeted strategy could prevent 38,200 premature deaths, generate 164,956 healthy life years and deliver \$45 for every \$1 invested over the next 20 years.⁷

Evidence reveals substantial gaps in identification of CKD, with up to 80–90% of Australians with stage 3 disease lacking a recorded diagnosis.^{17–19} Screening rates among certain high-risk groups – such as those with hypertension, cardiovascular disease (CVD) or obesity – remain significantly lower than in people with diabetes (unpublished data Turner GL, Aggar C, Rafferty R, et al). Notably, up to 90% of patients with CKD lack any urine albumin–creatinine ratio (uACR) assessment,^{20–22} which is essential for CKD screening, diagnosis and staging.¹ Late referral to a kidney service is common, with 15% of people commencing dialysis within 90 days of referral to a kidney service.^{1,23}

Aims

This article examines real-world barriers to CKD screening and detection in Australian primary care and outlines strategies to raise GP and patient awareness, promoting early diagnosis for better kidney health.

This consensus paper is based on discussion from three roundtable meetings held between October 2023 and April 2024 with primary care providers and kidney experts (Table 1). Participants were invited to attend one of the three roundtable meetings

and were briefed on the format of the meeting including the audio recording of the meetings for data analysis. Verbal consent was obtained at the start of the roundtable. Presentations were made drawing on the literature on gaps and barriers to CKD screening. The roundtable meeting identified local and contextualised barriers to CKD detection and screening, and participants explored potential solutions. The notes of the meeting were reviewed by RA and CG using multiple horizontal (line by line) and

vertical (sections of notes) passes to allow the development of barriers and themes emerging from the data.

An editorial committee developed a publication plan and prioritised outcomes. Literature searches were performed in November 2024 to provide evidence-based validation of themes.

Results and discussion

Six key barriers limiting CKD detection in general practice were identified, along with corresponding strategies to address each (Figure 2). Many solutions are interconnected across barriers, highlighting the potential for system-wide approaches such as clinical decision support tools and automated enhanced laboratory reporting to improve diagnosis.^{19,24-27}

Barrier 1: Low awareness and clinical inertia in primary care

Although CKD is twice as prevalent as diabetes,^{1,3} it has historically received less primary care focus, often being overshadowed by other comorbidities.²⁸ Studies indicate gaps in CKD primary care training and education.²⁸ A recent anonymous survey²⁹ of 399 Australian GPs found 33% were unaware of the KHA CKD Handbook.¹

CKD sits in the shadow of other chronic diseases. (Sydney GP)

Strategy: Educational activities, tools and supportive technology

Multidisciplinary education, practice audits and supportive technologies are recommended to help identify and diagnose at-risk patients.^{19,24-27} CKD screening is straightforward and cost effective (approximately \$26.00 for blood and urine test); fasting blood or first-void urine samples are not essential for opportunistic screening (Figure 1). Tools such as PenCS and integrated clinical decision models have demonstrated effectiveness in improving CKD diagnosis and management.^{19,24-27}

Greater promotion and utilisation of KHA resources such as the CKD Handbook,¹ CKD-Go! App and Kidney Health Professional Hub can strengthen primary care knowledge and practice (Figure 2).

Box 1. Case study

Patient profile: Mr RA

History

- Age: 40 years
- Occupation: Accountant
- Family history:
 - Mother: T2DM diagnosed at 51 years
 - Father: Smoker, died of MI at 65 years
- Lifestyle/medication:
 - Works long hours; sedentary lifestyle
 - 8 hours sleep/night; no medication

Examination

- BP: 138/92 mmHg; pulse: 76 bpm, regular
- BMI: 34 kg/m² (central obesity); waist circumference: 102 cm
- General examination: Unremarkable

Investigations/tests

- FBE: normal
- U&Es: eGFR: 61 mL/min/1.73m²; K⁺: 4.1 mmol/L
- Lipids: TC 5.8; HDL 1.0; LDL 3.0; TG 1.9 mmol/L
- HbA1c: 5.7%

Is opportunistic kidney screening warranted?

- Yes, obesity/metabolic syndrome and elevated blood pressure are risk factors for CKD (refer to Figure 1 for people who should be offered a kidney health check)¹
- uACR (in addition to eGFR and BP) is essential to check for kidney disease (Figure 1)
- A uACR was ordered. Test result: 15 mg/mmol

What are the next steps?

- Repeat eGFR and uACR to diagnose and stage CKD (Figure 1)
- Repeated test results show eGFR of 63 mL/min/1.73m² and uACR of 20 mg/mmol indicating stage 2 renal disease with microalbuminuria (Figure 1)
- Refer to KHA *CKD management in primary care* handbook to identify CKD cause and follow the 'yellow' management plan for stage 2 CKD (refer to <https://kidney.org.au/health-professionals/ckd-management-handbook>)¹

BMI, body mass index; BP, blood pressure; bpm, beats per minute; CKD, chronic kidney disease; eGFR, estimated glomerular filtration rate; FBE, full blood examination; HbA1c, haemoglobin A1c; HDL, high-density lipoprotein; K⁺, potassium; KHA, Kidney Health Australia; LDL, low-density lipoprotein; MI, myocardial infarction; T2DM, type 2 diabetes mellitus; TC, total cholesterol; TG, triglycerides; uACR, urine albumin-creatinine ratio; U&Es, urea and electrolytes.

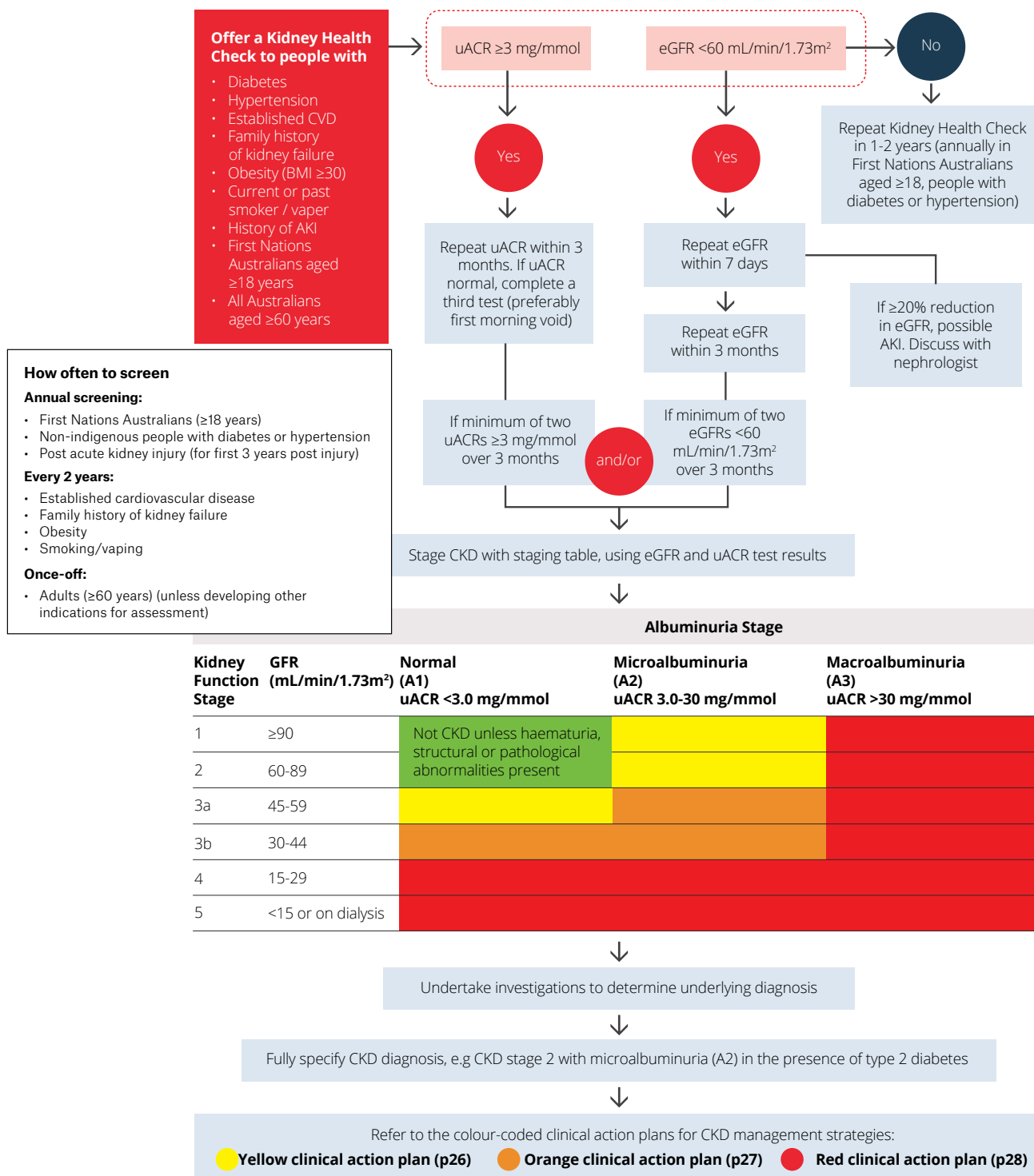


Figure 1. Kidney Health Australia algorithm for detection and diagnosis of chronic kidney disease.

Reproduced from Kidney Health Australia. Chronic kidney disease (CKD) management in primary care (5th edn). Kidney Health Australia, 2024, with permission. Please refer to the handbook for further information including management strategies and action plans.

Barrier 2: Multiple and disparate clinical guidelines

Kidney disease, heart disease and metabolic disorders, collectively known as cardiovascular-kidney-metabolic (CKM) syndrome,³⁰ are closely interconnected and should trigger assessment of all related conditions.³¹ Separate guidelines for CKD, diabetes and heart disease can confuse GPs and hinder effective care, especially if there are multiple comorbidities.^{24,28}

Strategy: Guideline harmonisation

Recognising metabolic syndrome as a shared precursor of CKM syndrome and adopting a multisystem approach³⁰ may enhance GP awareness and facilitate guideline adherence. CKD information in primary care resources, such as The Royal Australian College of General Practitioners' (RACGP's) *Guidelines for preventive activities in general practice* (Red Book) and HealthPathways, should align with the KHA CKD Handbook¹ to ensure consistency. Additionally, a unified risk calculator could help identify individuals at risk for both CVD and CKD, enabling more comprehensive care.

He found in his education with GPs that despite seeing the KHA book 20 times over, they have never engaged with it. (Sydney consultant pharmacist)

Barrier 3: CKD pathology and interpretation

Australian pathology reports define an eGFR >60 mL/min/1.73m² as 'normal' without considering any change over time or clinical context, such as patient age, history or concomitant illness. Moreover, uACR is often missed as part of CKD screening,²⁰⁻²² although it is vital for staging CKD to determine the best management approach.¹

Strategy: Automated pathology tests integrated with clinical pathways

Integrating a predefined kidney health request into general practice software is recommended to automate eGFR and uACR testing, accompanied by digital reminders to check blood pressure. Standardised pathology reporting, tailored to the clinical context, should seamlessly integrate with patients' electronic medical records (EMRs). Decision support tools within prescribing software, linked to the KHA CKD Handbook,¹ can further assist GPs.^{19,24-27} Additionally, any abnormal pathology should automate a prompt in clinical software to initiate further assessment.

PSA [prostate specific antigen] historical graph is included in pathology reports. eGFR longitudinal graph would be useful with a low eGFR flag, that shows historical data. (Brisbane GP)

Barrier 4: Inconsistent CKD documentation and coding

Currently, GPs do not consistently record a complete diagnosis of CKD,^{24,32} which requires eGFR, uACR and underlying aetiology.¹ Furthermore, EMR systems do not clearly distinguish CKD staging and have limited functionality to link comorbid conditions, impairing accurate diagnosis and effective management.

Strategy: Comprehensive CKD documentation

General practice software should integrate standardised CKD codes³¹ to record aetiology and automate longitudinal eGFR results and uACR tests to monitor disease staging and progression. Other recommendations include digital prompts to remind clinicians to screen at-risk patients

and repeat eGFR/uACR testing as needed for monitoring or rescreening. Moreover, incorporating CKD on death certificates would enable more precise measurement of disease burden.

Coding in the system is a big barrier and easily solved. (Brisbane GP)

Barrier 5: Insufficient time and funding

Limited time and funding are major barriers to CKD screening in primary care.^{28,33} Time pressures are exacerbated by poor communication among healthcare providers and fragmented care.^{28,34} Collaborative multidisciplinary approaches can improve diagnosis and optimise outcomes, particularly for patients with multiple comorbidities.^{28,31,35}

Strategy: Multidisciplinary communication and enhanced care models

A CKD RACGP Specific Interest group, or broader 'metabolic disease' subspecialty, is recommended to advocate for increased CKD investment. Opportunistic 'spot testing' in high-risk groups should be encouraged wherever possible³³ and effective recall mechanisms incorporated into clinical practice.³¹ Utilisation of home medication reviews and alternative workforce, such as practice nurses, chronic care or CKD nurses and diabetes educators, can enhance CKD screening efforts.^{31,36} New models of care that enhance interdisciplinary communication are recommended to improve patient outcomes.^{35,37}

Biggest barrier with CKD is limited time where GPs are faced with so many other factors and are unlikely to prioritise CKD. (Brisbane GP)

Table 1. Chronic kidney disease roundtable attendees, according to state and profession (n=35)

State	Renal physician	General practitioner	Nurse or nurse practitioner	Pharmacist	Patient or professional organisation	Number of participants
Victoria	1	4	1	2	2	10
New South Wales	0	8	2	3	0	13
Queensland	0	12	0	0	0	12
Total	1	24	3	5	2	35

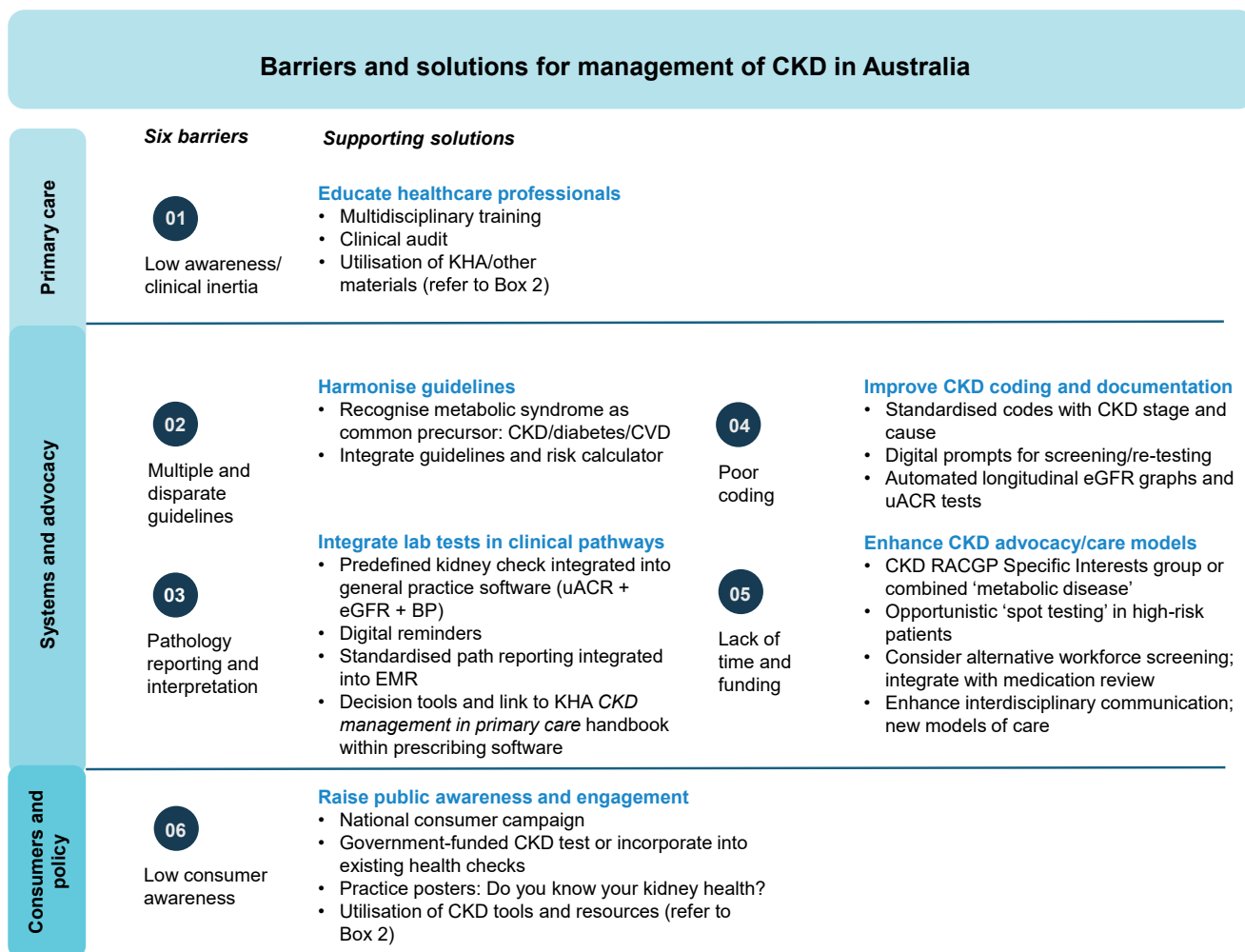


Figure 2. Key barriers to CKD screening and detection in Australian primary care and supporting strategies.

BP, blood pressure; CKD, chronic kidney disease; CVD, cardiovascular disease; eGFR, estimated glomerular filtration rate; EMR, electronic medical record; KHA, Kidney Health Australia; RACGP, The Royal Australian College of General Practitioners; uACR, urine albumin-creatinine ratio.

Barrier 6: Low consumer awareness

Limited consumer knowledge and understanding of CKD and its risk factors remains a significant barrier to early detection.^{31,34,38,39}

Strategy: Government policy and tools to raise public awareness and engagement

Government-funded programs are essential to improve CKD screening in at-risk populations. Recommendations include a national consumer campaign and either a Medicare Benefits Schedule-funded kidney health check or expansion of the Heart Health Check to

integrate heart/kidney/metabolic screening. Patient-focused surveys and posters in general practice can prompt earlier detection. Greater promotion and utilisation of the KHA Helpline (<https://kidney.org.au/ways-we-help/kidney-helpline>) and patient resources (<https://kidney.org.au/your-kidneys>), including Aboriginal and Torres Strait Islander-specific information, can empower shared decision making to improve patient outcomes.^{31,40} Engagement with Aboriginal and Torres Strait Islander communities and elders is essential to formulate targeted strategies for this at-risk population.^{5,41}

If there was more patient awareness and advocacy that helps keep these things front of mind. Consumer demand can improve doctor performance. (Brisbane GP)

Conclusion

The projected 39% rise in the number of Australians with stage 3–5 CKD from 2022 to 2034¹² demands immediate action. Earlier detection through targeted screening can reduce the healthcare burden and improve patient outcomes. With effective treatments now available to slow disease progression

Box 2. Useful resources

Primary provider materials

- Kidney Health Australia CKD Management in Primary Care handbook, <https://kidney.org.au/health-professionals/ckd-management-handbook>
- CKD-Go! App, <https://kidney.org.au/new-ckd-go-app>
- Kidney Health Professional Hub, <https://kidney.org.au/health-professionals/hp-hub-sign-up>
- Tools to identify at-risk patients (eg PenCAT, POLAR, Cubiko)

Patient resources

- Kidney helpline, <https://kidney.org.au/ways-we-help/kidney-helpline>
- Patient support materials, <https://kidney.org.au/your-kidneys>

and prevent both kidney failure and cardiovascular complications, the investment in early detection is compelling.^{1,7} Although targeted CKD screening is cost effective¹⁶ and straightforward, shifting from late-stage treatment to early intervention requires innovative approaches. A coordinated, multipronged strategy is essential to expand screening to all at-risk groups. Success will require systematic strategies, government support and improved coding and documentation to monitor disease impact and intervention effectiveness.

Key points

- The asymptomatic nature of CKD demands proactive, repeated screening for early identification.
- Early detection and intervention delay disease progression.
- Screening is simple, effective and cheap: non-fasting blood + urine sample + BP (Figure 1).
- It is important to make the link between the kidneys, diabetes and the heart, with detection of disease at one site prompting investigation at the others.
- Resources available for GPs include the KHA CKD Handbook, CKD-Go! App, Professional Hub, patient materials and helpline.

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Competing interests: RA has been an advisor and paid lecturer for AstraZeneca, Aspen Pharmacare, Seqirus, Boehringer Ingelheim, Novartis, Novo Nordisk, Sevier, Eli Lilly, Bristol Myers Squibb and Menarini; KD has delivered education and received honorarium from AstraZeneca, Boehringer Ingelheim and Bayer, and has been on advisory committees for AstraZeneca and CSL. She is currently receiving funding from Western Alliance, Epworth Medical Foundation. She has received educational grants from Abbott and funding from Deakin University, National Health and Medical Research Council, Medical Research Future Fund and Servier; AM has received honoraria for consultancy, advisory boards and presentations from GSK, CSL, MSD, Moderna, Pfizer, Amgen, Abbott, Pharmaceutical Society of Australia (PSA), Immunisation Coalition, United Clinical, Praxhub and The Royal Australian College of General Practitioners. She has received travel grants from PSA, CSL and MSD; RR has provided medical education and advisory board consultancy for AstraZeneca, Boehringer Ingelheim, Eli Lilly Australia, GSK, MSD, Novo Nordisk and Sanofi-Aventis Australia; GT has received honoraria for education and presentations from AstraZeneca, Amgen and Fresenius; CMH received honoraria for education consultancy, advisory boards, speaker presentations from Amgen, AstraZeneca, CSIRO, CSL, Seqirus, GSK, Lundbeck, Menarini, Moderna, MSD, Novartis, Novo Nordisk and Pfizer, and she has received research support from Amgen and Sanofi.

Funding: The logistics of the roundtable meetings and medical writing support were funded by AstraZeneca, Australia. The authors did not receive any funding for preparation of the manuscript and were solely

responsible for its content and all related decisions. The authors had full access to all relevant data in this study, and the sponsor (AstraZeneca) had no involvement in data analysis and interpretation, or in the writing of the article.

Provenance and peer review: Not commissioned, externally peer reviewed.

AI declaration: The authors confirm that there was no use of artificial intelligence (AI)-assisted technology for assisting in the writing or editing of the manuscript and no images were manipulated using AI.

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Acknowledgements

The authors wish to thank the roundtable participants for their participation and extend gratitude to Celia Green, Director at Bioscript, Melbourne, Victoria, for her medical writing assistance.

References

1. Kidney Health Australia. Chronic kidney disease (CKD) management in primary care. 5th edn. Kidney Health Australia, 2024.
2. Australian Institute of Health and Welfare (AIHW). Chronic kidney disease: Australian facts. AIHW, 2023. Available at www.aihw.gov.au/reports/chronic-kidney-disease/chronic-kidney-disease/contents/impact-of-chronic-kidney-disease/burden-of-chronic-kidney-disease# [Accessed 2 December 2024].
3. Australian Bureau of Statistics (ABS). Australian health survey: Biomedical results for chronic diseases, 2011–12. ABS, 2013.
4. Bello AK, Levin A, Lunney M, et al. Status of care for end stage kidney disease in countries and regions worldwide: International cross sectional survey. *BMJ* 2019;367:l5873. doi: 10.1136/bmj.l5873.
5. Tunnicliffe DJ, Bateman S, Arnold-Chamney M, et al. Recommendations for culturally safe clinical kidney care for First Nations Australians: A guideline summary. *Med J Aust* 2023;219(8):374–85. doi: 10.5694/mja2.52114.
6. Lakhan P, Cooney A, Palamuthusingam D, et al. Challenges of conducting kidney health checks among patients at risk of chronic kidney disease and attending an urban Aboriginal and Torres Strait Islander primary healthcare service. *Aust J Prim Health* 2022;28(5):371–79. doi: 10.1071/PY21248.
7. Deloitte Access Economics. Changing the chronic kidney disease landscape: The economic benefits of early detection and treatment. Kidney Health Australia, 2023.
8. Chadban S, Arıcı M, Power A, et al. Projecting the economic burden of chronic kidney disease at the patient level (Inside CKD): A microsimulation modelling study. *EClinicalMedicine* 2024;72:102615. doi: 10.1016/j.eclinm.2024.102615.
9. Tonelli M, Wiebe N, Culleton B, et al. Chronic kidney disease and mortality risk: A systematic review. *J Am Soc Nephrol* 2006;17(7):2034–47. doi: 10.1681/ASN.2005101085.
10. Keith DS, Nichols GA, Gullion CM, Brown JB, Smith DH. Longitudinal follow-up and outcomes among a population with chronic kidney disease in a large managed care organization. *Arch Intern Med* 2004;164(6):659–63. doi: 10.1001/archinte.164.6.659.
11. Thompson S, James M, Wiebe N, et al; Alberta Kidney Disease Network. Cause of death in patients with reduced kidney function. *J Am Soc Nephrol* 2015;26(10):2504–11. doi: 10.1681/ASN.2014070714.

12. Moura AF, Wheeler D, Brotons-Munto F, et al. WCN24-1204 Multidimensional burden of chronic kidney disease in eight countries: Insights from the IMPACT CKD study. *International Society of Nephrology (ISN) 2024;9(4) Suppl S263*. doi: 10.1016/j.ekir.2024.02.540.
13. Meraz-Muñoz AY, Weinstein J, Wald R. eGFR decline after SGLT2 inhibitor initiation: The tortoise and the hare reimaged. *Kidney360 2021;2(6):1042–47*. doi: 10.34067/KID.0001172021.
14. Fernández-Fernández B, Sarafidis P, Soler MJ, Ortiz A. EMPA-KIDNEY: Expanding the range of kidney protection by SGLT2 inhibitors. *Clin Kidney J 2023;16(8):1187–98*. doi: 10.1093/ckj/sfad082.
15. Heerspink HJL, Stefánsson BV, Correa-Rotter R, et al; DAPA-CKD Trial Committees and Investigators. Dapagliflozin in patients with chronic kidney disease. *N Engl J Med 2020;383(15):1436–46*. doi: 10.1056/NEJMoa2024816.
16. Yeo SC, Wang H, Ang YG, Lim CK, Ooi XY. Cost-effectiveness of screening for chronic kidney disease in the general adult population: A systematic review. *Clin Kidney J 2023;17(1):sfad137*. doi: 10.1093/ckj/sfad137.
17. Pecoits-Filho R, Ribeiro de Castro MCR, Cebrian A, et al. # 3667 REVEAL-CKD: Prevalence of undiagnosed stage 3 chronic kidney disease in Australia, Brazil, Canada and Spain. *Neph Dialysis Transplant 2023;38(Suppl 1)*. doi: 10.1093/ndt/gfad063c_3667.
18. Khanam MA, Kitsos A, Stankovich J, et al. Chronic kidney disease monitoring in Australian general practice. *Aust J Gen Pract 2019;48(3):132–37*. doi: 10.31128/AJGP-07-18-4630.
19. Pefanis A, Botlero R, Langham RG, Nelson CL. eMAP:CKD: Electronic diagnosis and management assistance to primary care in chronic kidney disease. *Nephrol Dial Transplant 2018;33(1):121–28*.
20. Tangri N, Alvarez CS, Arnold M, et al. Suboptimal monitoring and management in patients with unrecorded stage 3 chronic kidney disease in real-world settings: Insights from REVEAL-CKD. *Eur J Clin Invest 2024;54(11):e14282*. doi: 10.1111/eci.14282.
21. Radford J, Kitsos A, Stankovich J, et al. Epidemiology of chronic kidney disease in Australian general practice: National Prescribing Service MedicineWise MedicineInsight dataset. *Nephrology (Carlton) 2019;24(10):1017–25*. doi: 10.1111/nep.13537.
22. Jun M, Wick J, Neuen BL, et al. The prevalence of CKD in Australian primary care: Analysis of a national general practice dataset. *Kidney Int Rep 2023;9(2):312–22*. doi: 10.1016/j.ekir.2023.11.022.
23. Australia and New Zealand Dialysis and Transplant Registry (ANZDATA). Chapter 1: Incidence of kidney failure with replacement therapy. In: 46th Annual Report 2023. ANZDATA, 2023. Available at <https://anzorrg.org.au/reports/anzdata-46th-annual-report-2023-data-to-2022> [Accessed 10 November 2025].
24. Jones JL, Lumsden NG, Simons K, et al. Using electronic medical record data to assess chronic kidney disease, type 2 diabetes and cardiovascular disease testing, recognition and management as documented in Australian general practice: A cross-sectional analysis. *Fam Med Community Health 2022;10(1):e001006*. doi: 10.1136/fmch-2021-001006.
25. Jones JL, Simons K, Manski-Nankervis JA, et al. Chronic disease IMPACT (chronic disease early detection and improved management in primary care project): An Australian stepped wedge cluster randomised trial. *Digit Health 2023;9:20552076231194948*. doi: 10.1177/20552076231194948.
26. Hunter B, Davidson S, Lumsden N, et al. Optimising a clinical decision support tool to improve chronic kidney disease management in general practice. *BMC Prim Care 2024;25(1):220*. doi: 10.1186/s12875-024-02470-w.
27. McBride C, Hunter B, Lumsden N, et al. Clinical acceptability of a quality improvement program for reducing cardiovascular disease risk in people with chronic kidney disease in Australian general practice: Qualitative study. *JMIR Hum Factors 2024;11:e55667*. doi: 10.2196/55667.
28. Neale EP, Middleton J, Lambert K. Barriers and enablers to detection and management of chronic kidney disease in primary healthcare: A systematic review. *BMC Nephrol 2020;21(1):83*. doi: 10.1186/s12882-020-01731-x.
29. Guppy M, Bowles EJ, Glasziou P, Doust J. General practitioners' assessment and management of chronic kidney disease in older patients – A mixed methods study. *BMC Prim Care 2024;25(1):312*. doi: 10.1186/s12875-024-02559-2.
30. Ndumele CE, Rangaswami J, Chow SL, et al; American Heart Association. Cardiovascular-kidney-metabolic health: A presidential advisory from the American Heart Association. *Circulation 2023;148(20):1606–35*. doi: 10.1161/CIR.0000000000001184.
31. Robson B, Deed G, Phoon RK. Improving the detection and management of kidney health in primary care. *J Patient Exp 2024;11:23743735241256464*. doi: 10.1177/23743735241256464.
32. Kitsos A, Peterson GM, Jose MD, Khanam MA, Castelino RL, Radford JC. Variation in documenting diagnosable chronic kidney disease in general medical practice: Implications for quality improvement and research. *J Prim Care Community Health 2019;10:2150132719833298*. doi: 10.1177/2150132719833298.
33. Sinclair PM, Day J, Levett-Jones T, Kable A. Barriers and facilitators to opportunistic chronic kidney disease screening by general practice nurses. *Nephrology (Carlton) 2017;22(10):776–82*. doi: 10.1111/nep.12856.
34. Lo C, Teede H, Fulcher G, et al. Gaps and barriers in health-care provision for co-morbid diabetes and chronic kidney disease: A cross-sectional study. *BMC Nephrol 2017;18(1):80*. doi: 10.1186/s12882-017-0493-x.
35. Lo C, Zimbudzi E, Teede H, et al. Models of care for co-morbid diabetes and chronic kidney disease. *Nephrology (Carlton) 2018;23(8):711–17*. doi: 10.1111/nep.13232.
36. Bonner A, Douglas B, Brown L, et al. Understanding the practice patterns of nephrology nurse practitioners in Australia. *J Ren Care 2023;49(4):278–87*. doi: 10.1111/jorc.12444.
37. Manski-Nankervis JA, Alexander K, Biezen R, et al. Towards optimising chronic kidney disease detection and management in primary care: Underlying theory and protocol for technology development using an Integrated Knowledge Translation approach. *Health Informatics J 2021;27(2):14604582211008227*. doi: 10.1177/14604582211008227.
38. Guha C, Lopez-Vargas P, Ju A, et al; PAVE-CKD workshop investigators. Patient needs and priorities for patient navigator programmes in chronic kidney disease: A workshop report. *BMJ Open 2020;10(11):e040617*. doi: 10.1136/bmjopen-2020-040617.
39. Lopez-Vargas PA, Tong A, Howell M, et al. Patient awareness and beliefs about the risk factors and comorbidities associated with chronic kidney disease: A mixed-methods study. *Nephrology (Carlton) 2017;22(5):374–81*. doi: 10.1111/nep.12829.
40. Havas K, Douglas C, Bonner A. Meeting patients where they are: Improving outcomes in early chronic kidney disease with tailored self-management support (the CKD-SMS study). *BMC Nephrol 2018;19(1):279*. doi: 10.1186/s12882-018-1075-2.
41. Tunnicliffe DJ, Bateman S, Arnold-Chamney M, et al. Recommendations for culturally safe and clinical kidney care for First Nations Australians. *CARI Guidelines, 2022*.

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