

Kidney disease in Tasmania

This webinar will start shortly.

Kidney disease in Tasmania

Zoom webinar – Tuesday 22 July 2025

6.30 pm to 8 pm

Acknowledgement of traditional owners


We acknowledge the Tasmanian Aboriginal people as the traditional owners and ongoing custodians of the land on which we are meeting today. We pay our respects to Elders past and present.

We would also like to acknowledge Aboriginal people who are joining us today.



Learning outcomes

After this session, I will be able to:

- Describe the epidemiology of kidney disease for Tasmania overall, as well as your own practice location
 - Identify risk factors for kidney disease and implement early detection strategies applicable to your clinical practice
 - Summarise the pharmacological interventions indicated for kidney disease once detected
 - Review optimal kidney disease management in your clinical practice
- 
- A decorative footer consisting of various shades of blue geometric shapes, including triangles and polygons, arranged in a modern, abstract pattern.

Some housekeeping

- Tonight's webinar is being recorded
- Please use the Zoom Q&A feature to ask questions
- At the end of the webinar your browser will automatically open an evaluation survey. We appreciate you taking the time to complete this to help us improve our events programme
- Please don't forget to register for your next webinar at:
<https://www.primaryhealthtas.com.au/for-health-professionals/events/>

Presenters



Matthew D. Jose



Jan Radford



Lisa Shelverton

Kidney disease in Tasmania

Matthew D. Jose

Head of Renal Unit, Royal Hobart Hospital
Professor of Medicine, University of Tasmania

Lisa Shelverton

Renal Nurse Practitioner, Royal Hobart Hospital

Jan Radford

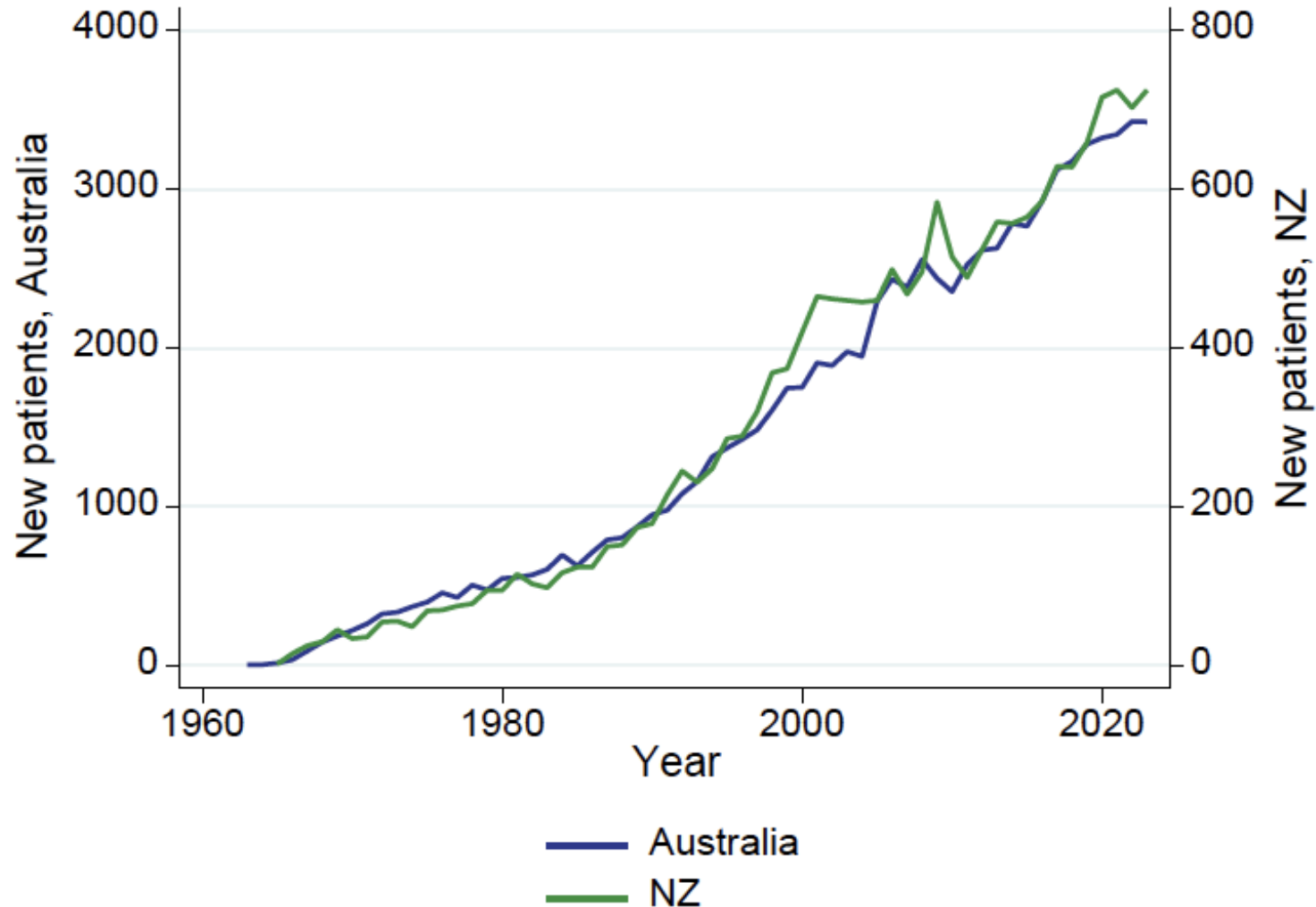
Professor in General Practice, University of Tasmania

Acknowledgement of country

This work was undertaken in *Iutruwita* (Tasmania) Aboriginal land, sea and waterways. I acknowledge, with deep respect the traditional owners of this land, the muwinina and palawa peoples, their elders past, present and emerging.



Incident (treated) End-Stage kidney disease in ANZ



Cost

Dialysis A\$60,000 per year

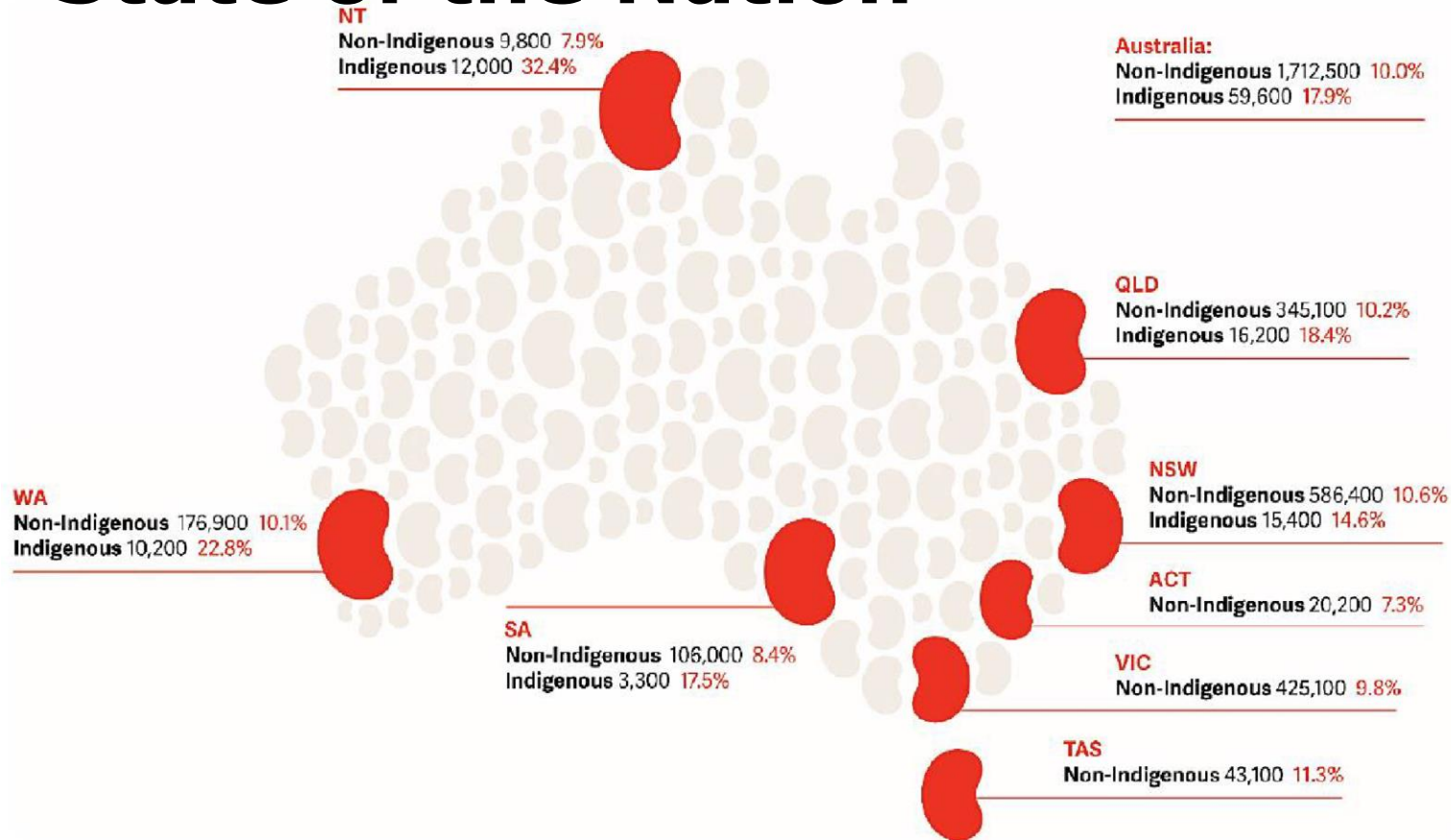
Transplant:

1st year A\$70,000

Beyond A\$10,000

Tasmania: highest (non-Indigenous) state prevalence of CKD

State of the Nation



Non-Indigenous CKD:

Australia 10.0%

Tasmania 11.3%

SA 8.4%

Vic 9.8%

Figure 5. Living with signs of CKD – Number of people (proportion of population)⁸

Tasmania: lowest state prevalence of dialysis

Prevalence of dialysis

~25% lower. ANZDATA 2023

Tasmania 410 per million pop

SA 581 pmp

Vic 537 pmp

NSW 566 pmp

Australia 586 per million
pop.

Overall Aim:

To understand how
high CKD in the community
translates into
low use of dialysis.

Figure 1.4.7
New Patients by Age Group - TAS

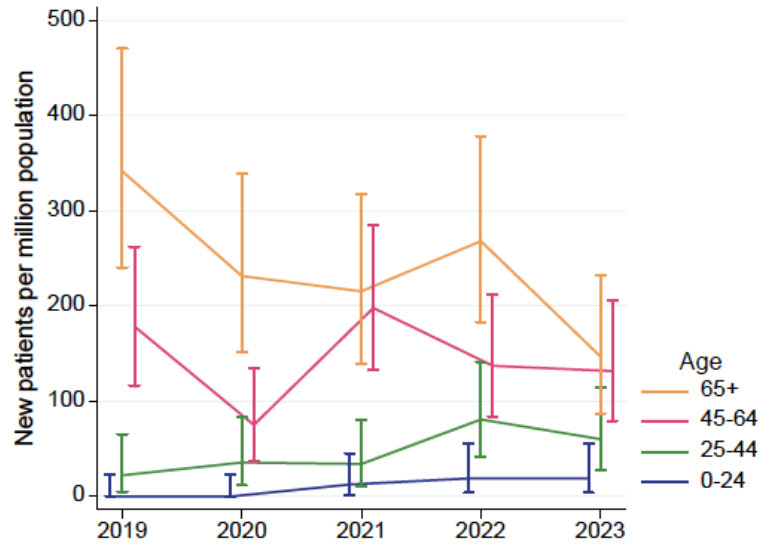
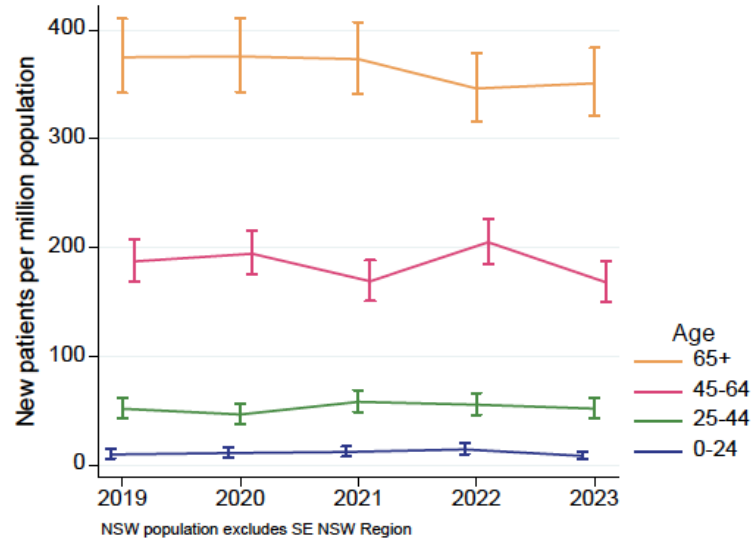


Figure 1.4.2
New Patients by Age Group - NSW



**Tasmania has lower
rates of treated kidney
failure for age ≥ 45 years
old**

Figure 1.4.3
New Patients by Age Group - VIC

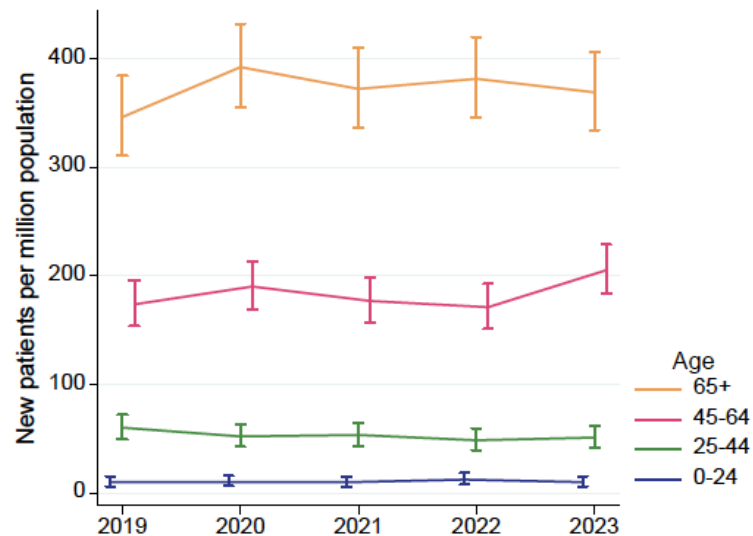
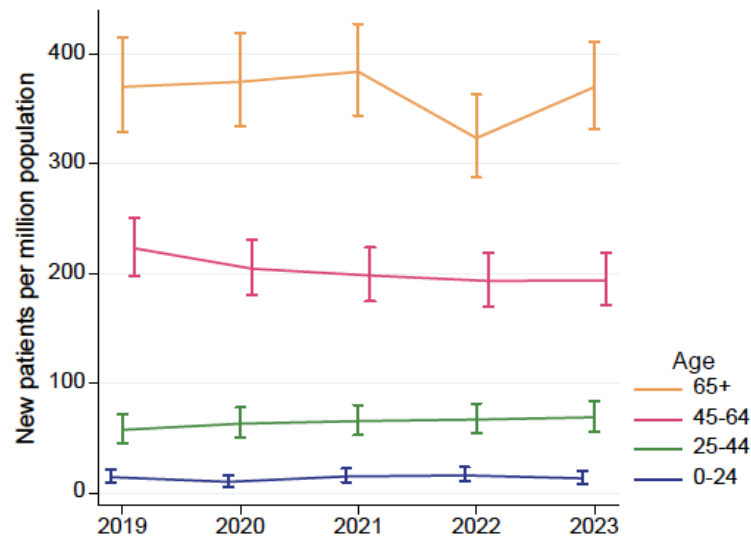


Figure 1.4.4
New Patients by Age Group - QLD



**ANZDATA
47th Annual Report 2024**

Tasmania: lowest state prevalence of dialysis

Prevalence of dialysis

~25% lower. ANZDATA 2023

Tasmania 410 per million pop

SA 581 pmp

Vic 537 pmp

NSW 566 pmp

Australia 586 per million

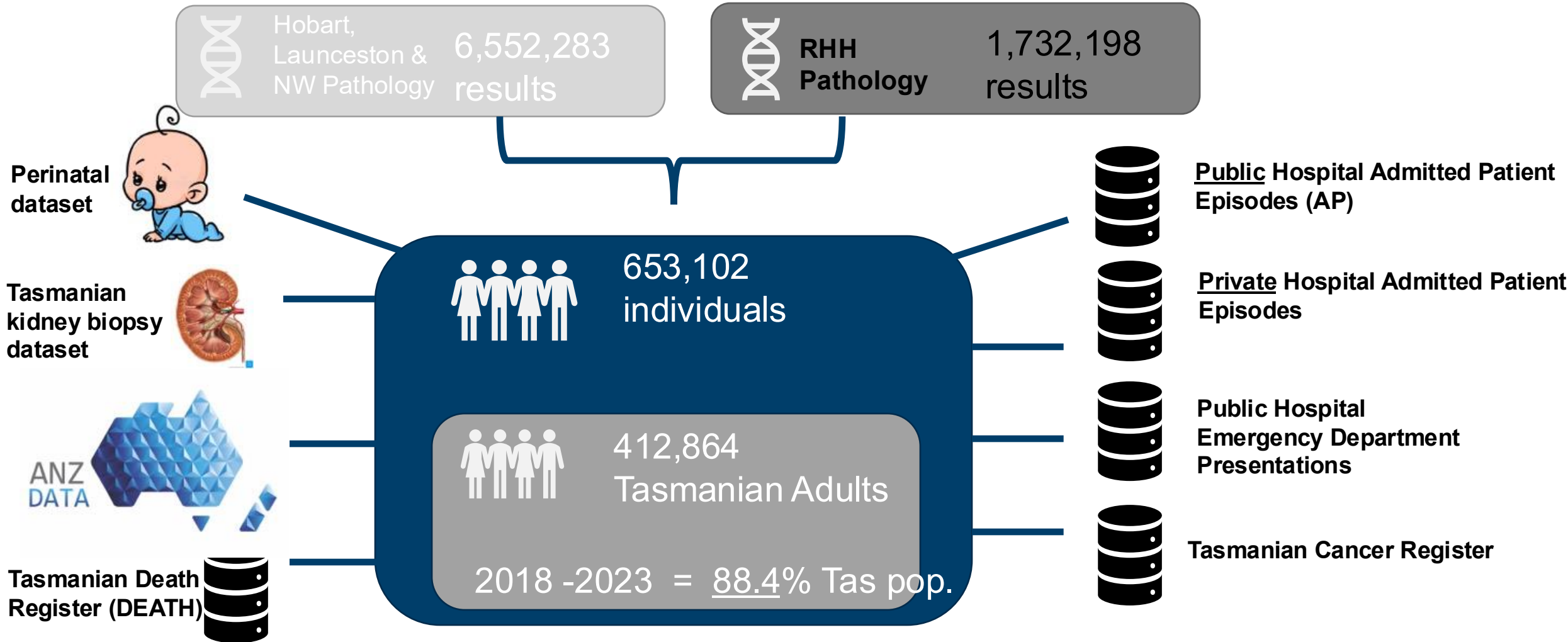
pop.

Overall Aim:

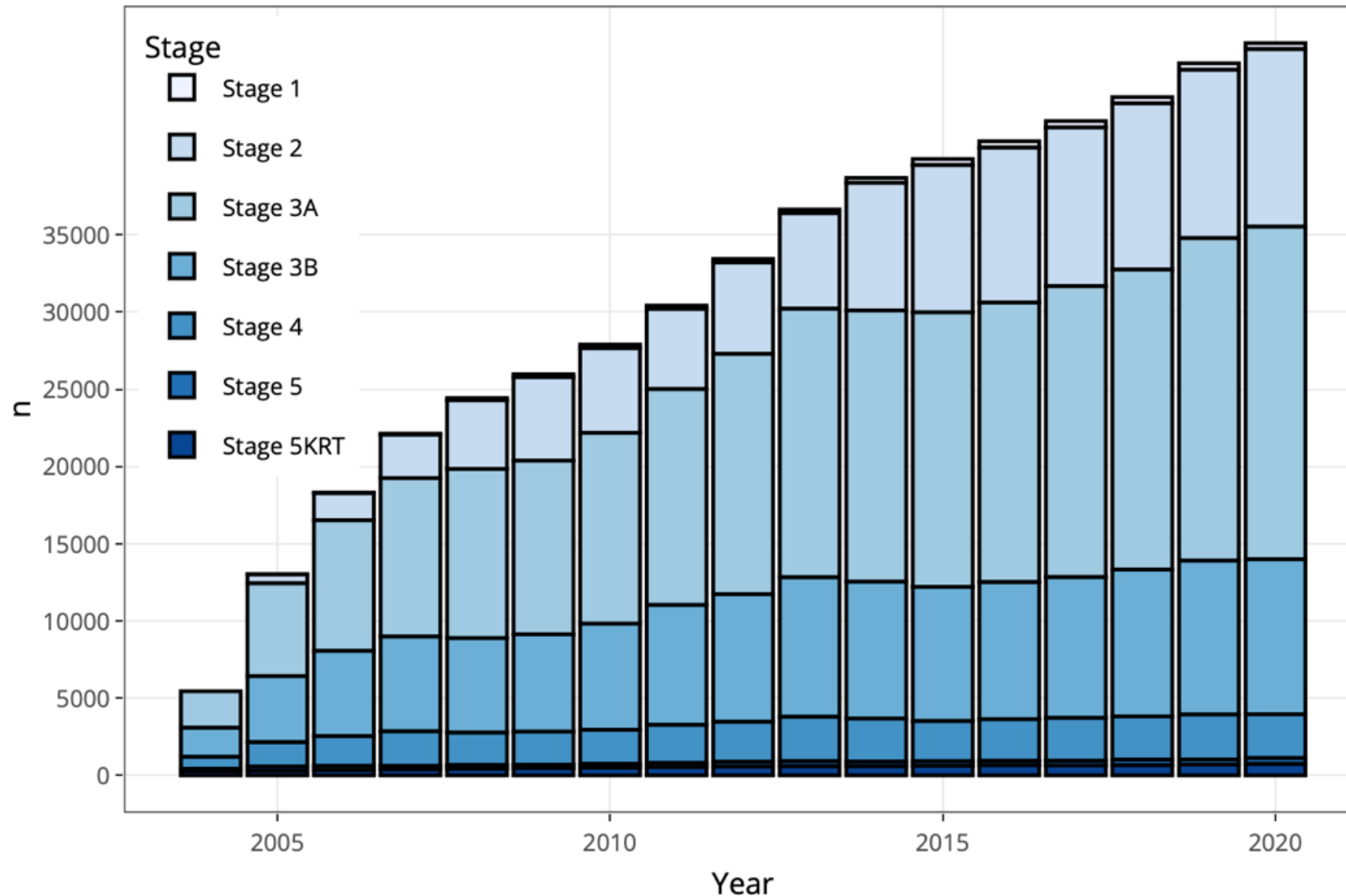
To understand how
**high CKD in the
community translates
into
low use of dialysis.**

- What have we done?
- **Kidney disease in Tasmania?**
 - Epidemiology & growth of chronic disease
 - Competing risk of death or dialysis
 - Co-design of decision aids
- What next?

Serum Creatinine between Jan 1 2004 to Dec 31 2023



Results Chronic disease on the rise



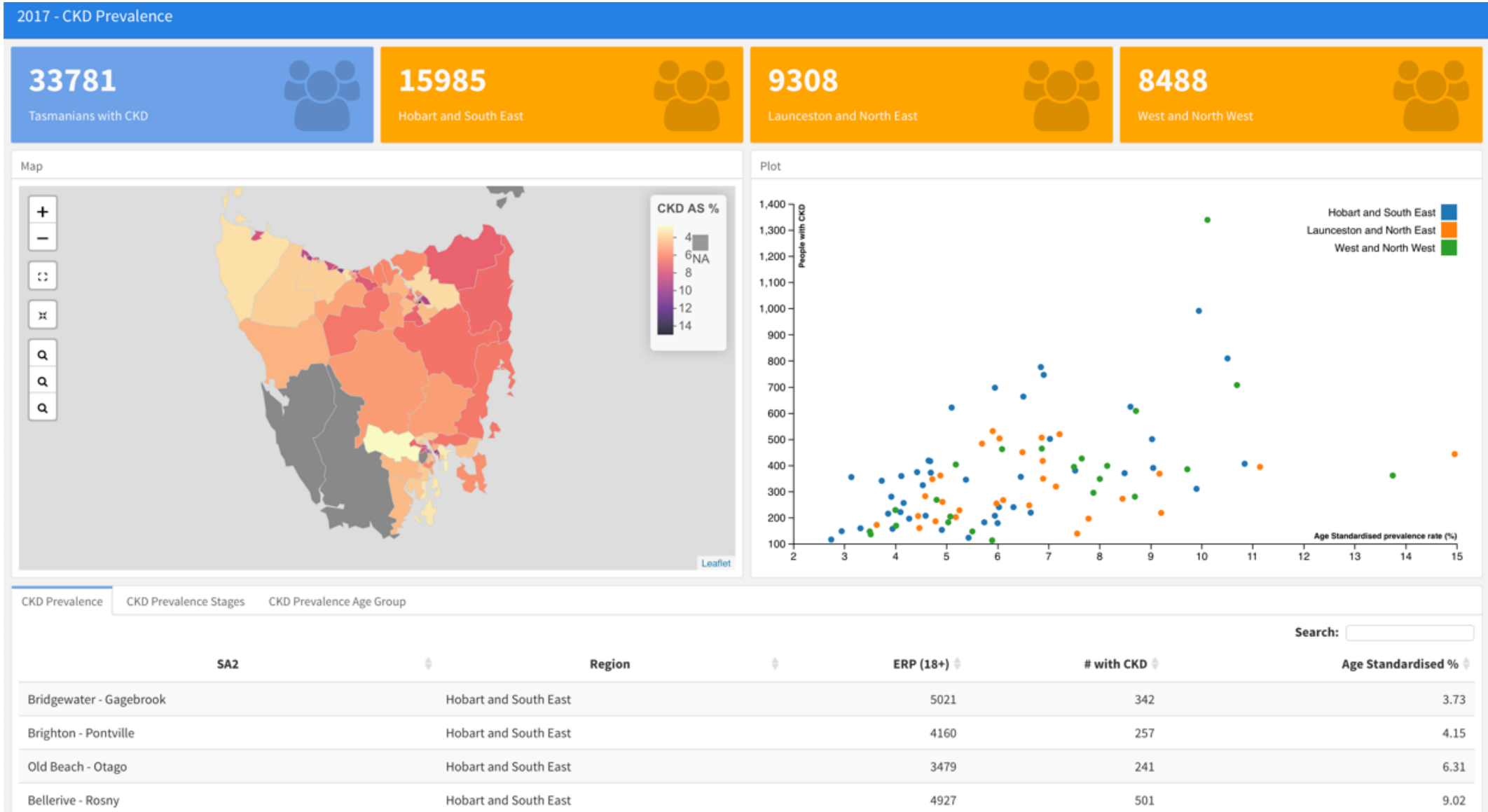
Total Stage 3-5

2010 = 22,544

2020 = 37,747

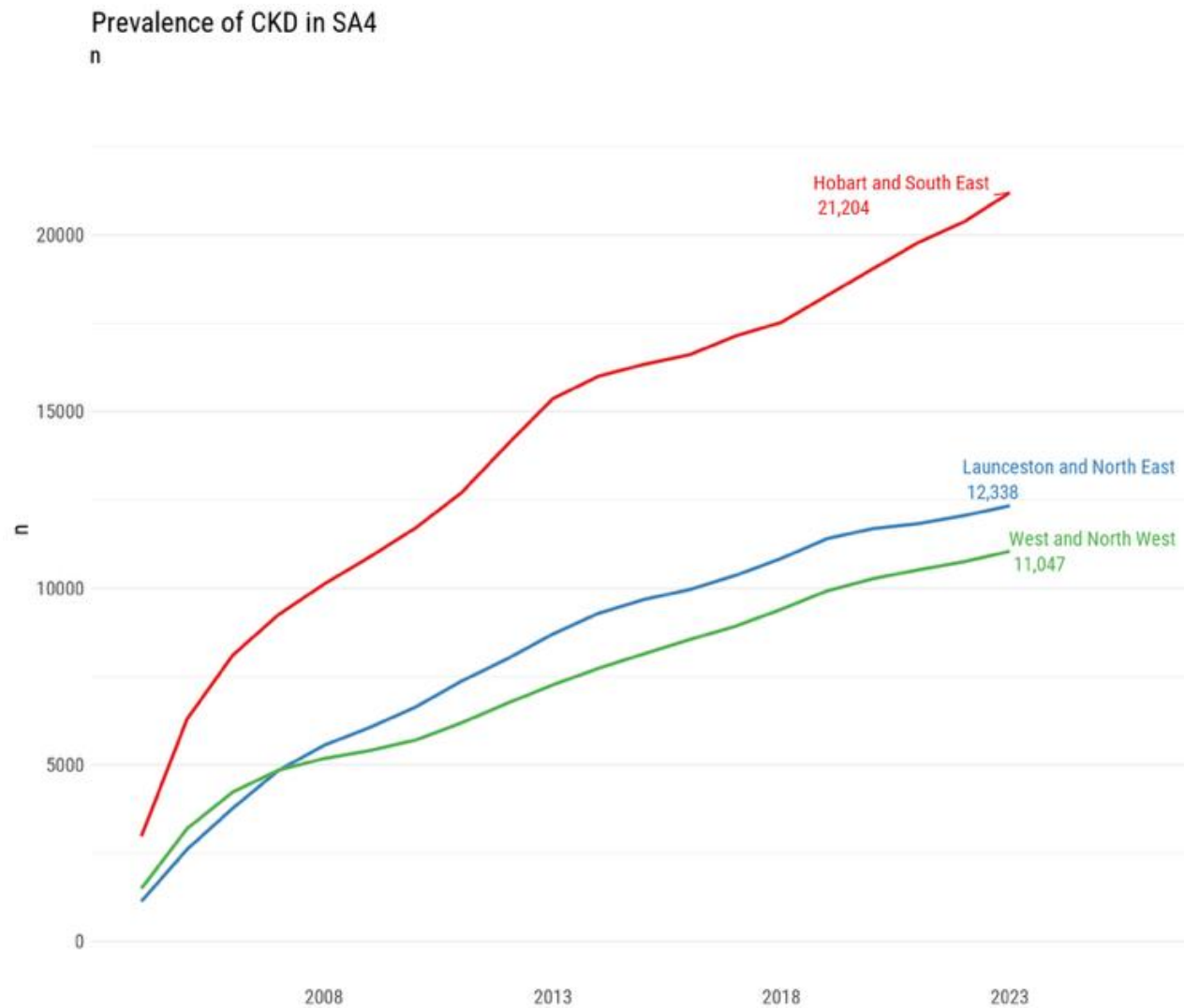
67% increase
in number of
people

CKD prevalence & treatment locations

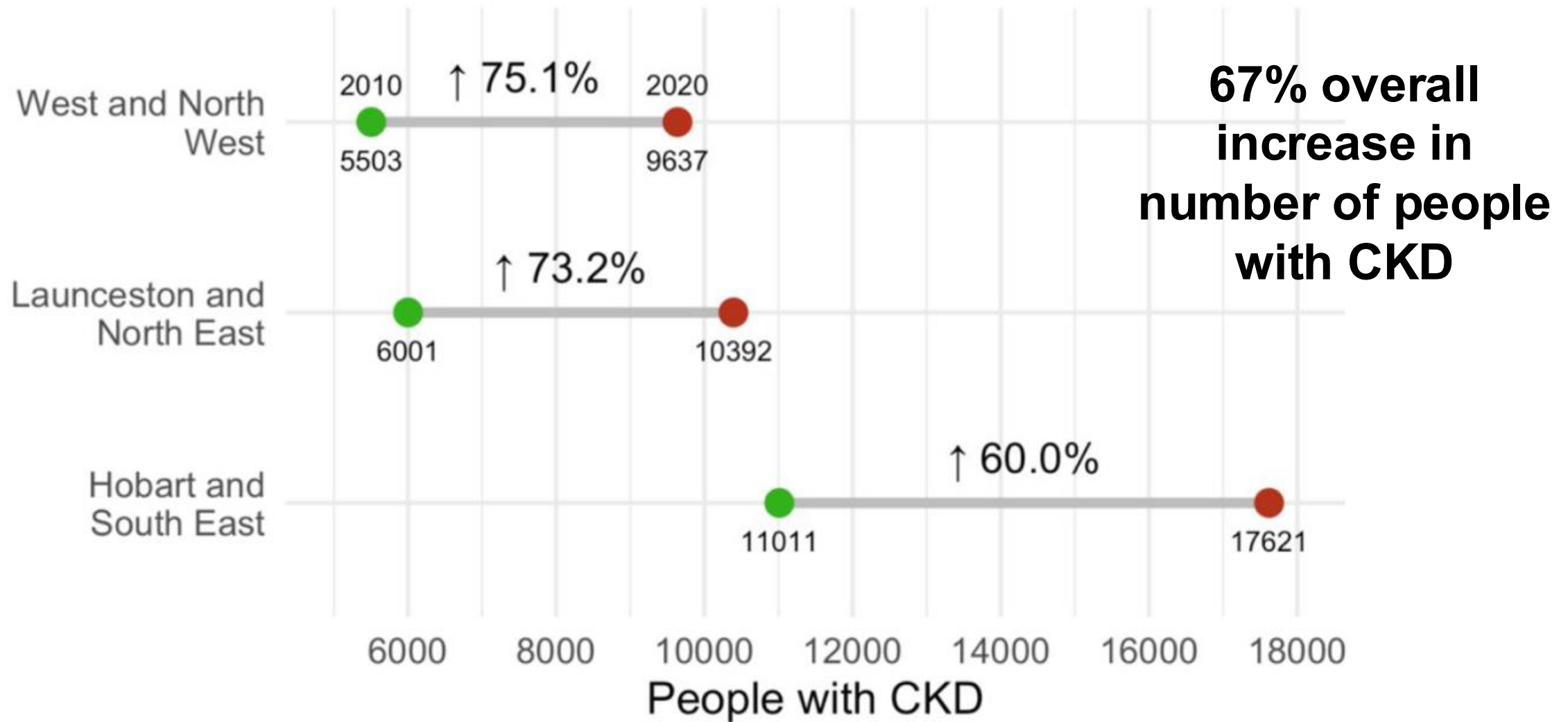


Results

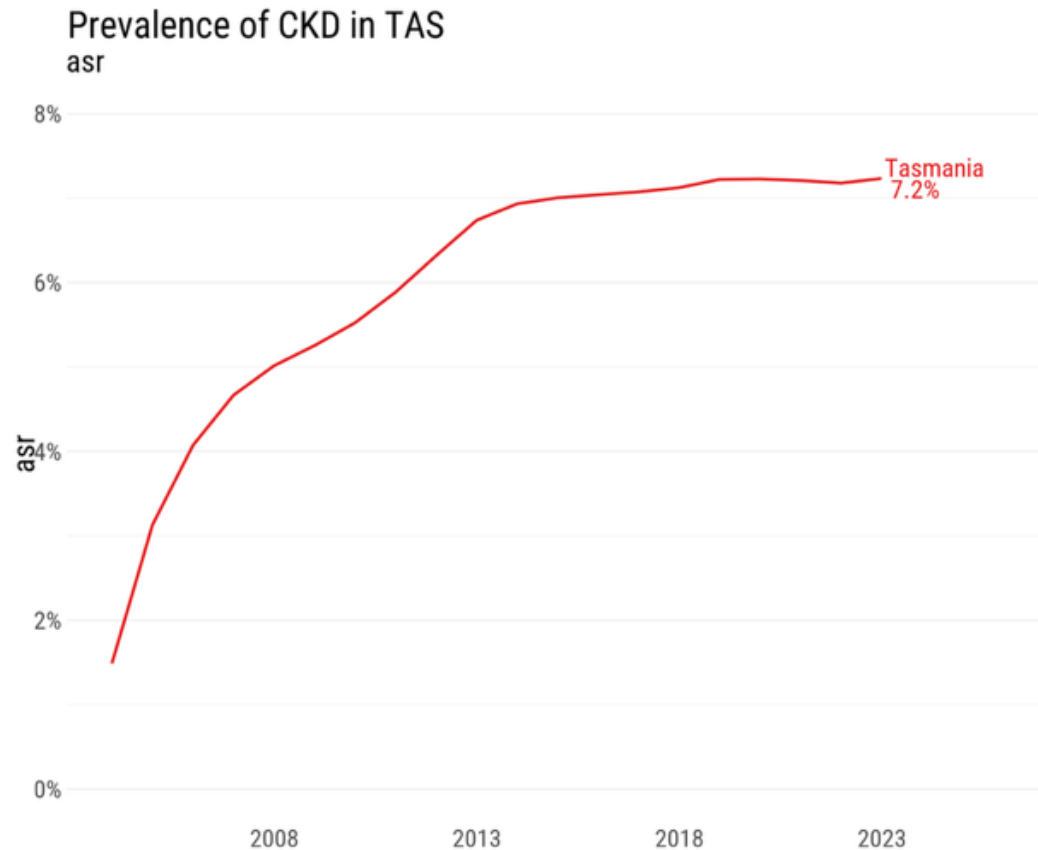
Geographic variation by SA4



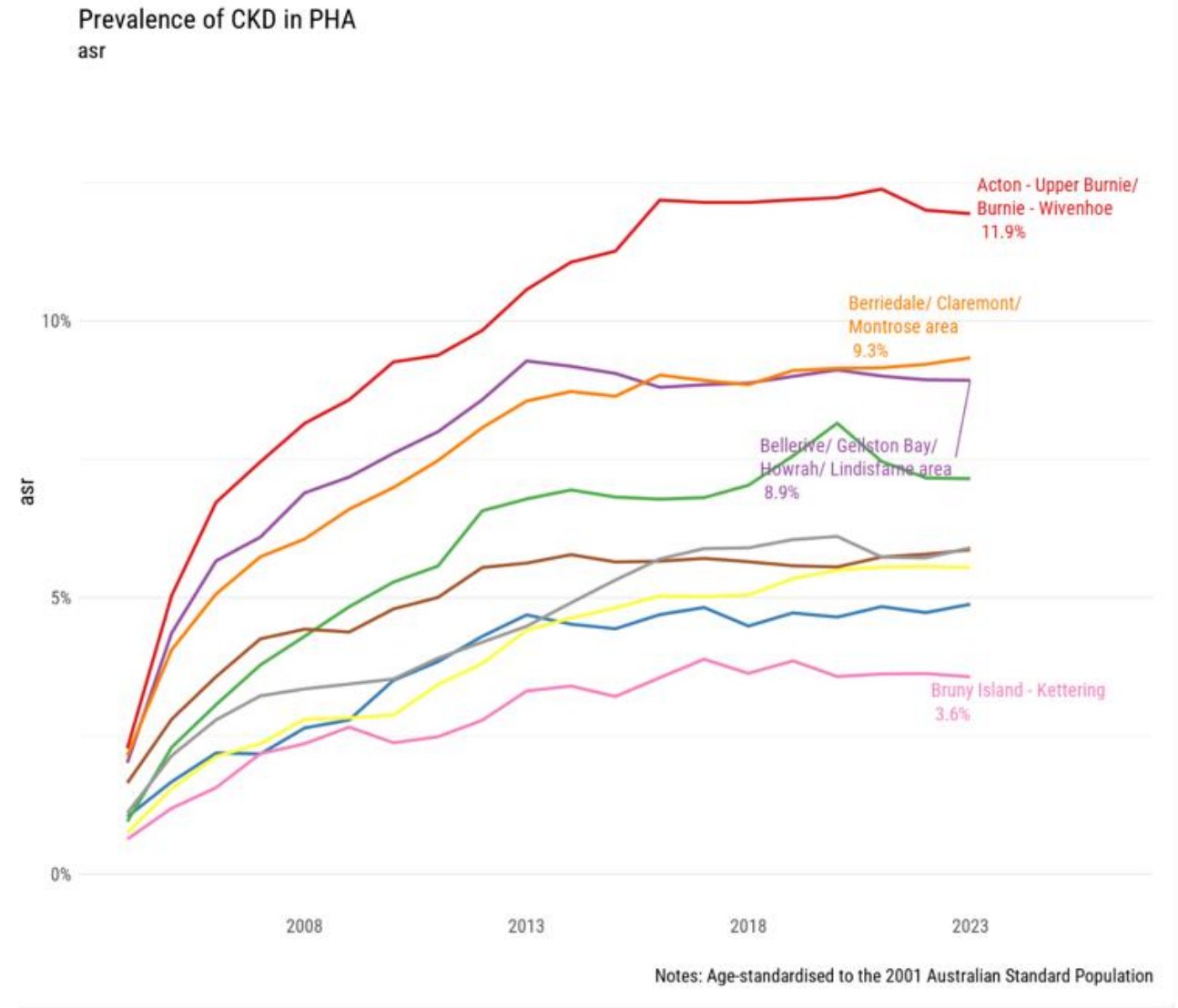
Results Geographic variation by SA4



Results Geographic variation by PHA

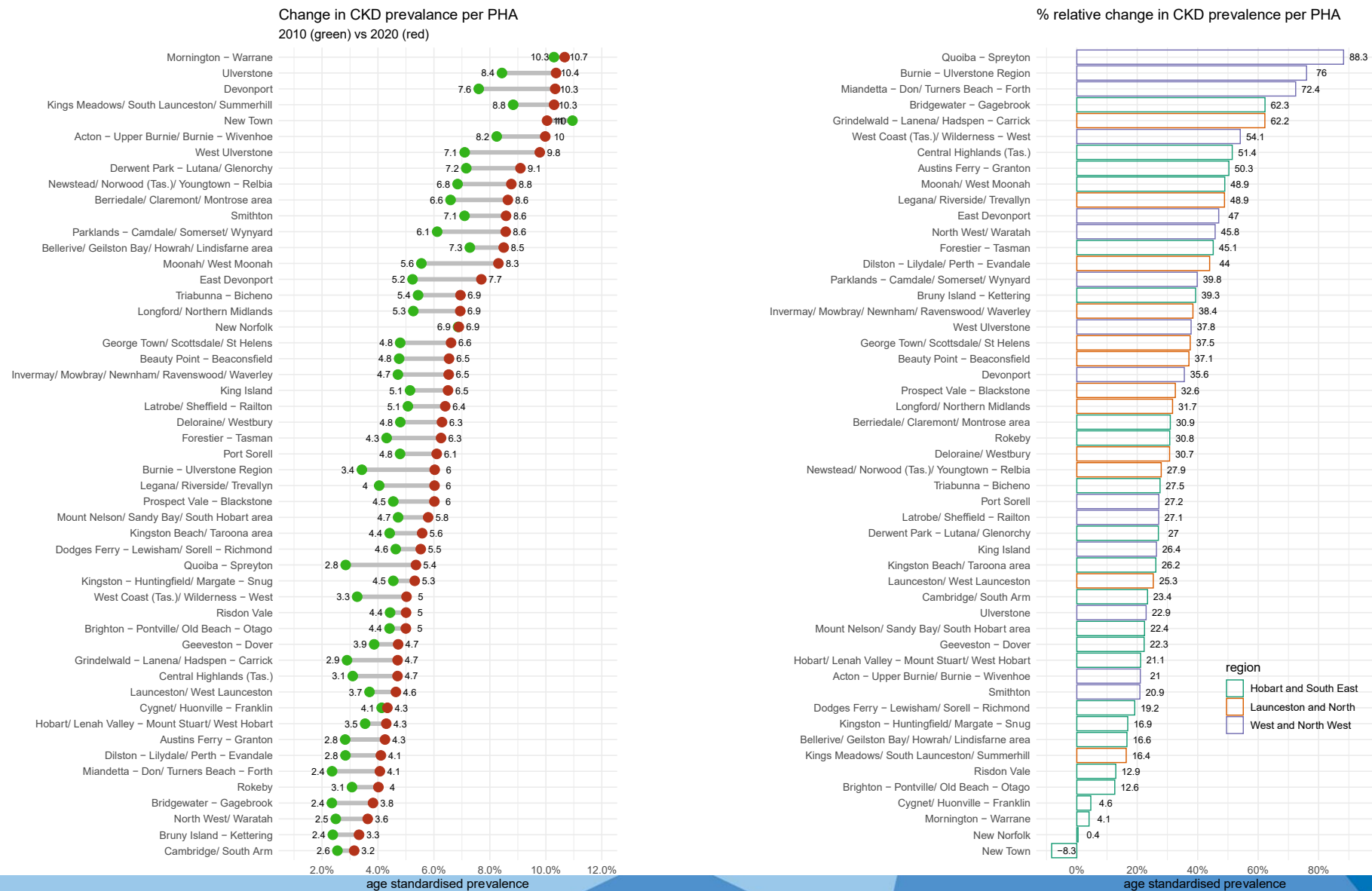


Notes: Age-standardised to the 2001 Australian Standard Population



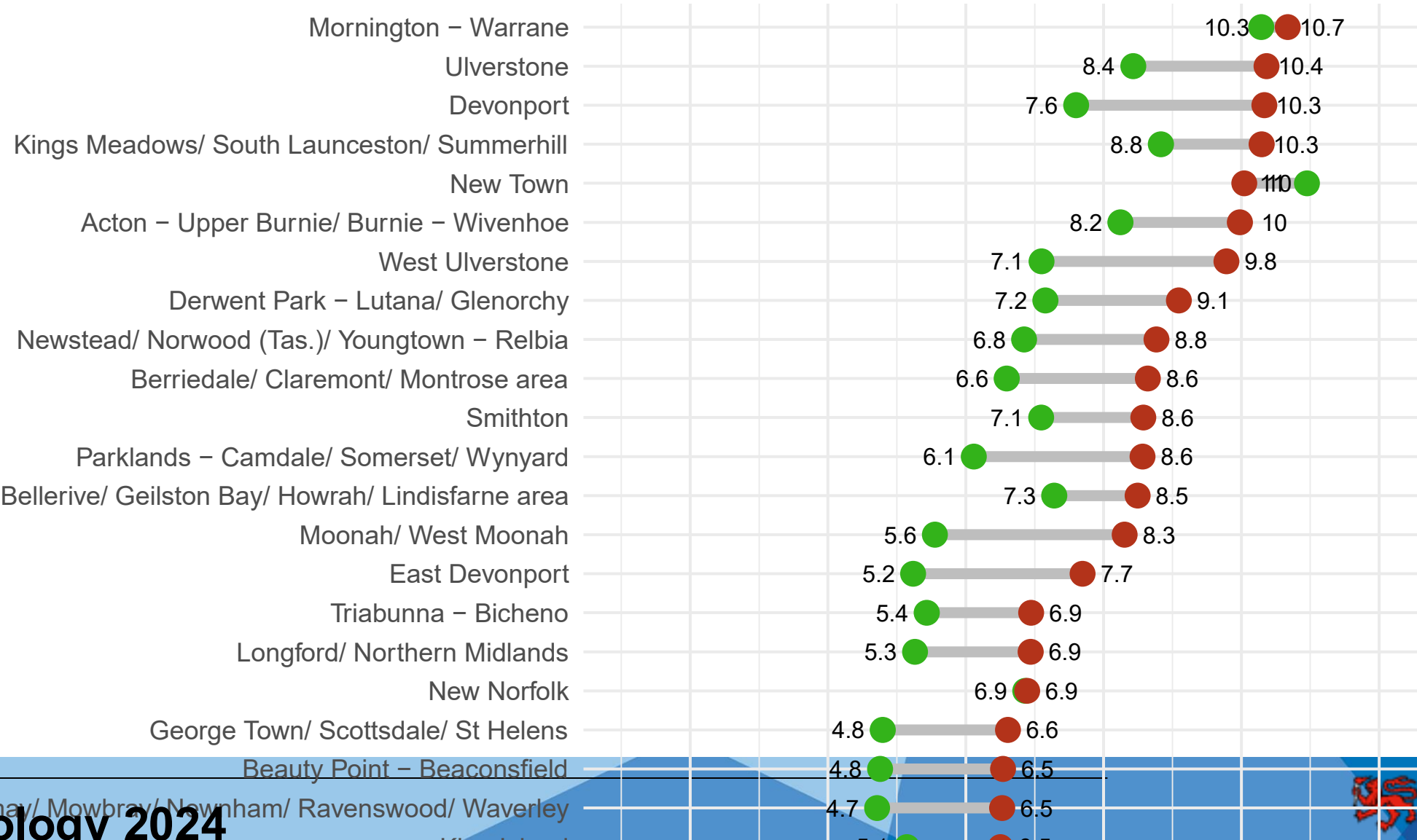
Notes: Age-standardised to the 2001 Australian Standard Population

CKD in Tasmania increased by 28% in past decade (2010 to 2020)

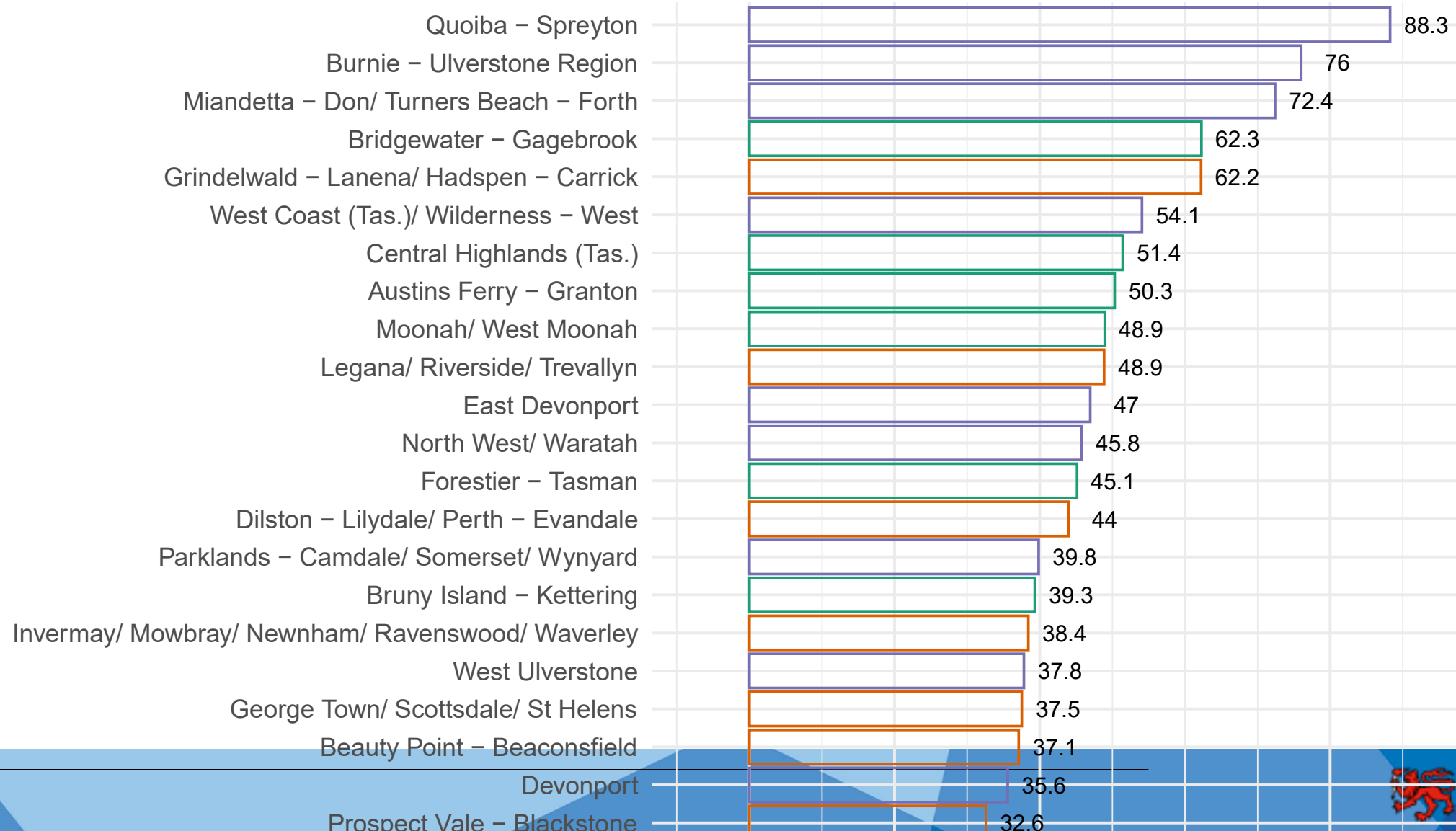


CKD in Tasmania increased by 28% in past decade (2010 to 2020)

Change in CKD prevalence per PHA
2010 (green) vs 2020 (red)



% relative change in CKD prevalence per PHA



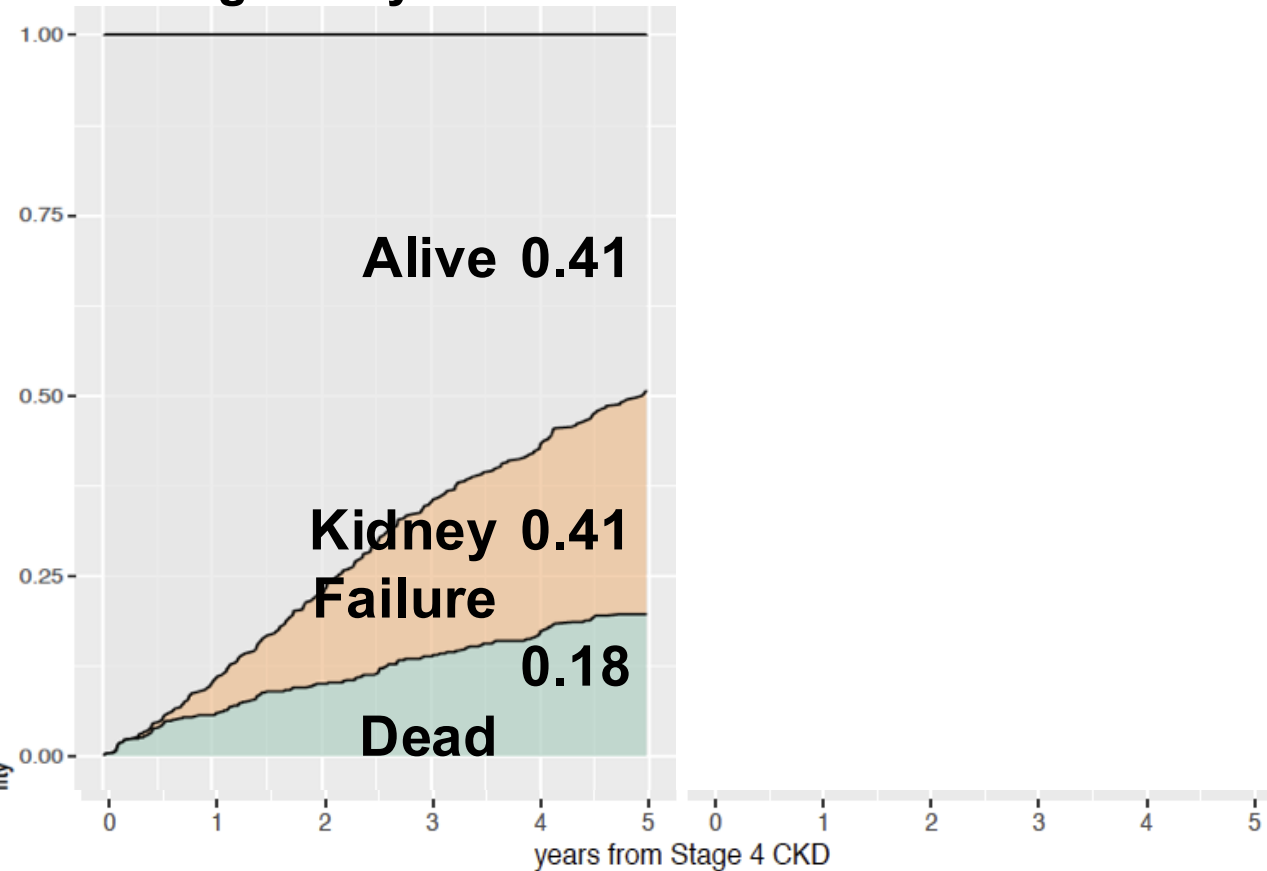
- What have we done?
- **Kidney disease in Tasmania?**
 - Epidemiology & growth of chronic disease
 - **Competing risk of death or dialysis**
 - Co-design of decision aids
 - Mother and baby outcomes with early kidney disease
- What next?

Competing risks analysis:

Stage 4 CKD to death or kidney failure* ($<10\text{mL/min/1.73m}^2$) at 5 yrs?

*Ravani et al, JAMA 2020

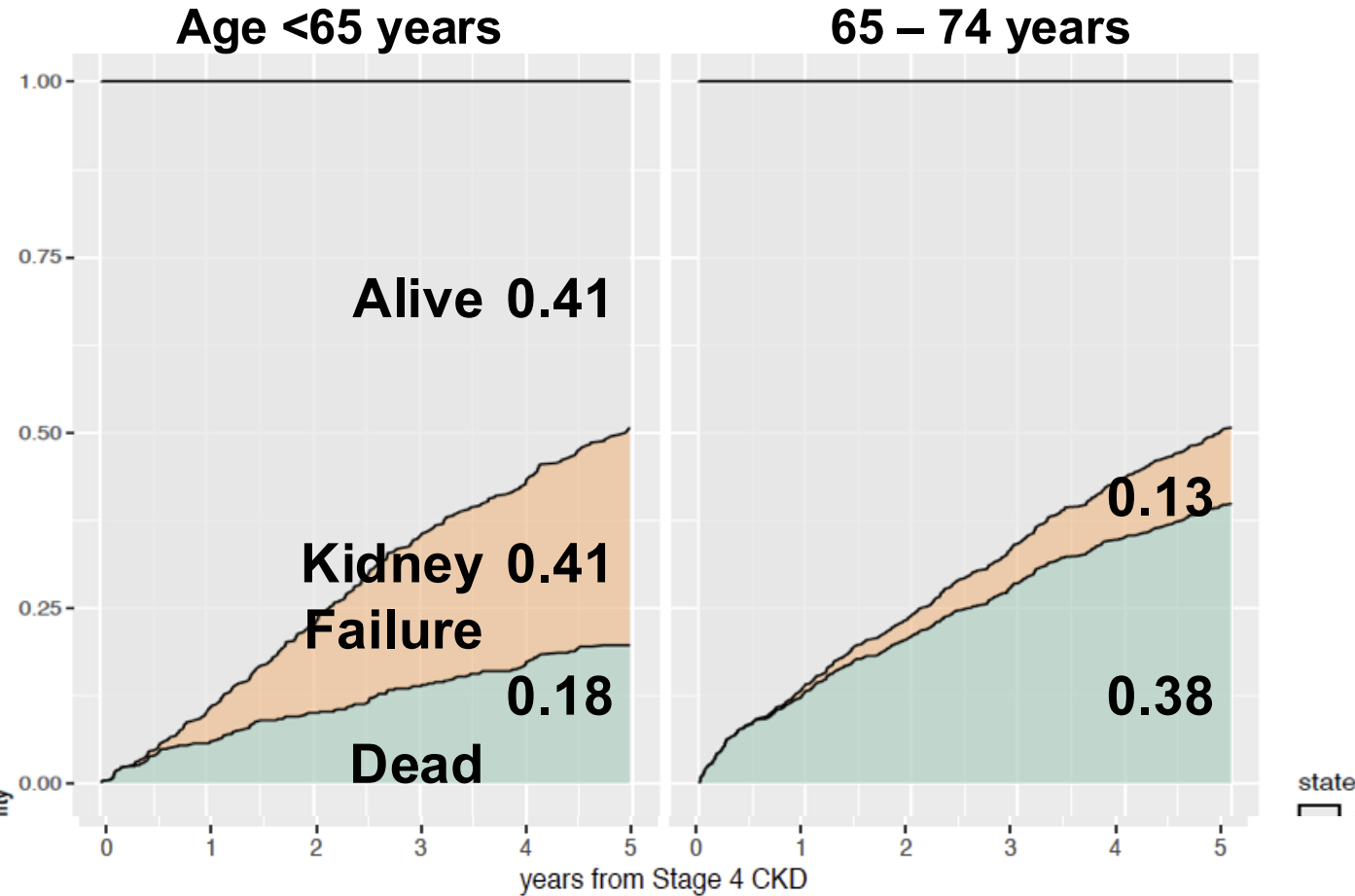
Age <65 years



Competing risks analysis:

Stage 4 CKD to death or kidney failure* ($<10\text{mL/min/1.73m}^2$) at 5 yrs?

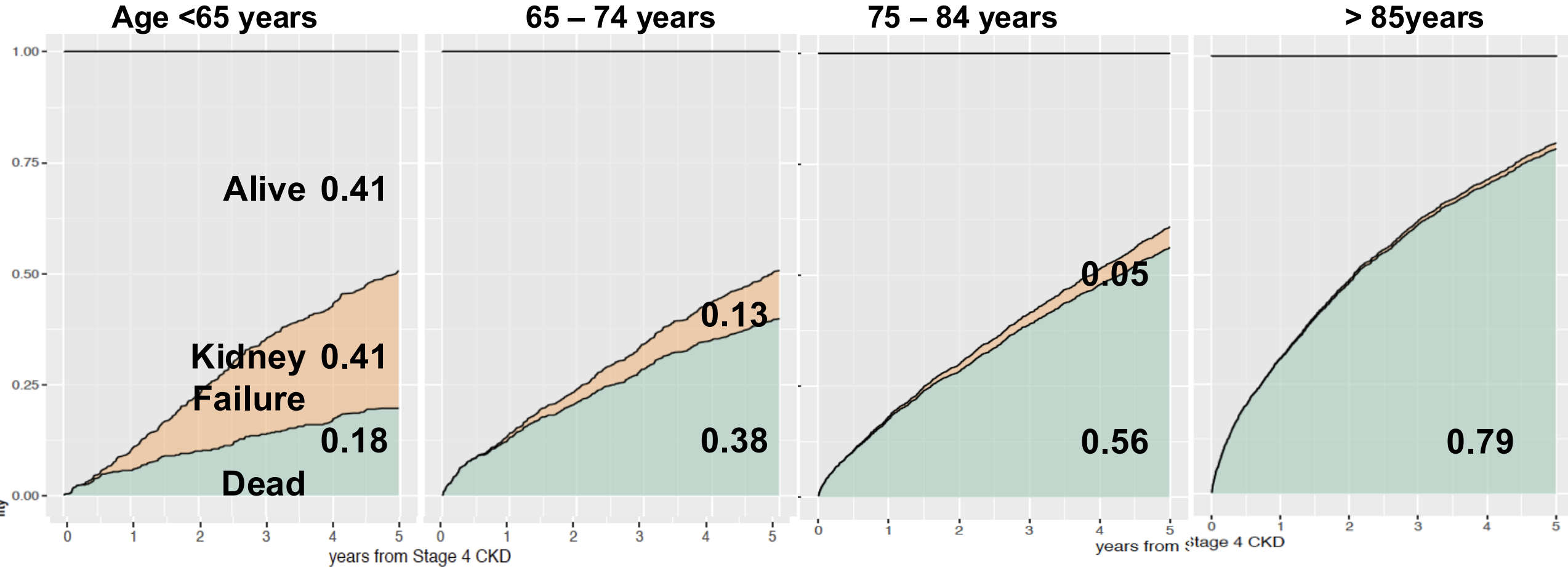
*Ravani et al, JAMA 2020



Competing risks analysis:

Stage 4 CKD to death or kidney failure* ($<10\text{mL/min/1.73m}^2$) at 5 yrs?

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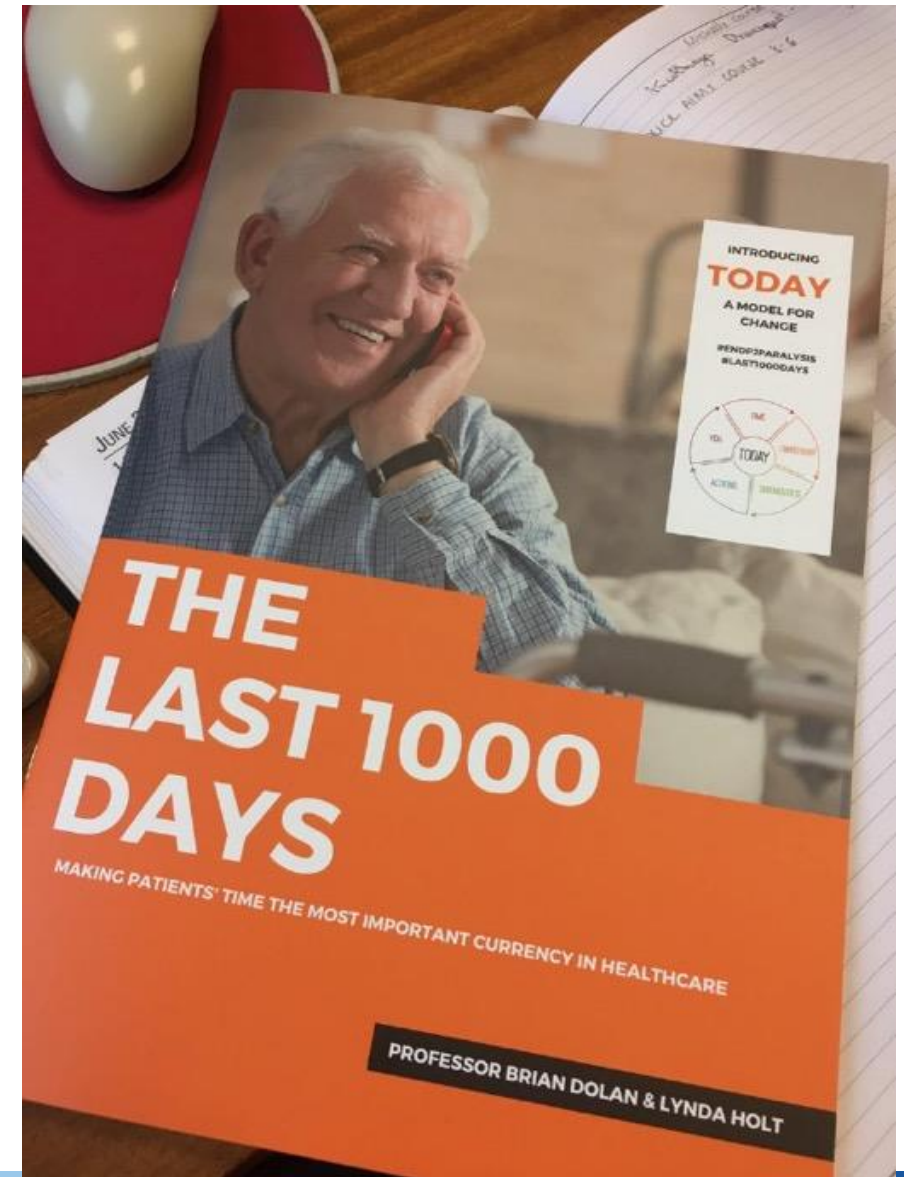


- What have we done?
- Kidney disease in Tasmania?
 - Epidemiology & growth of chronic disease
 - Competing risk of death or dialysis
 - **Co-design of decision aids**
 - Mother and baby outcomes with early kidney disease
- What next?

First 1000 days of Life



Last 1000 days



***If I don't have dialysis,
I'll die.***

comment heard about once a week

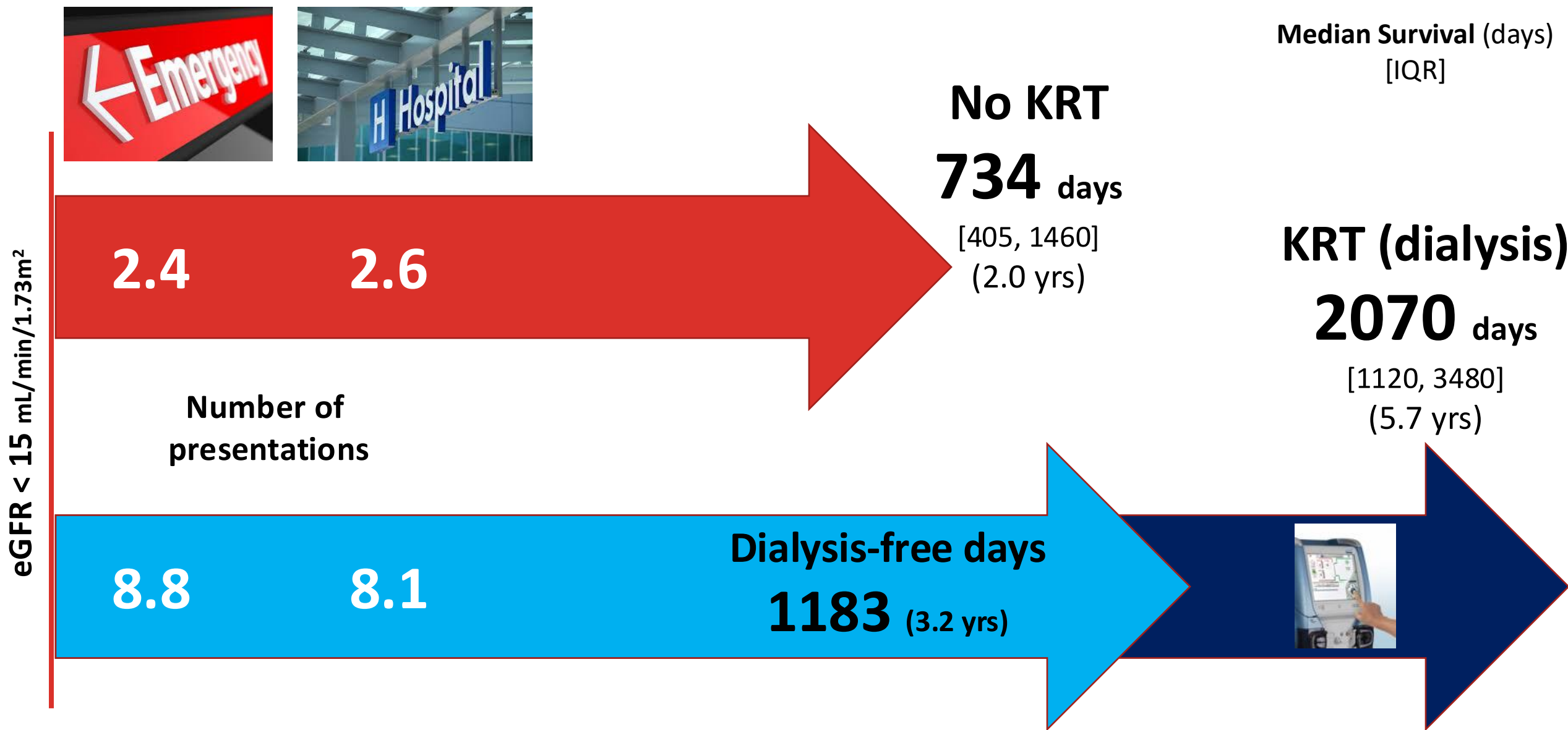


Treatment pathways with CKD – it's about time



Age <65 years

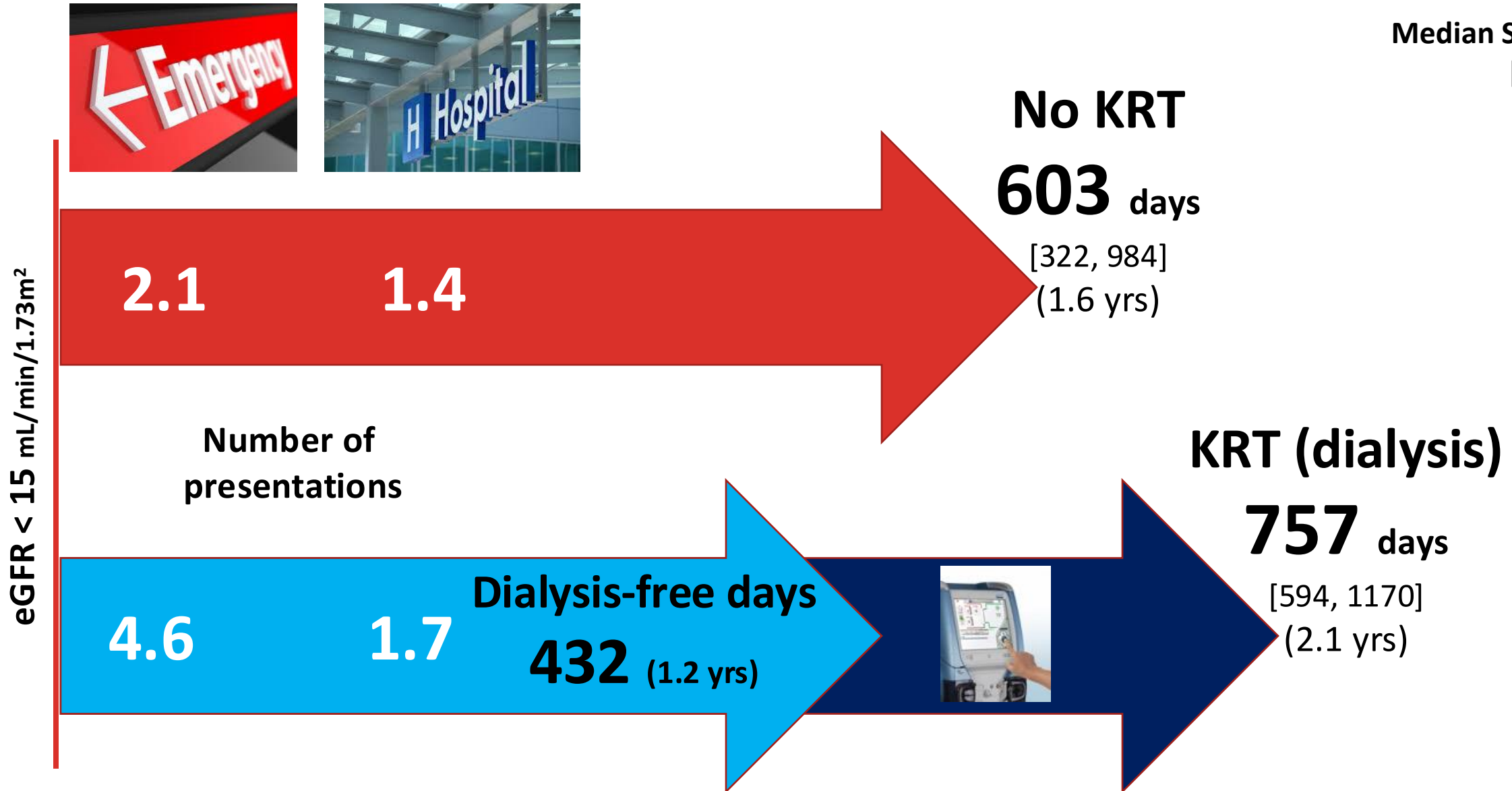
(from incident eGFR<15 mL/min/1.73m²)



Age 85+ years

(from incident eGFR < 15 mL/min/1.73m²)

Median Survival (days)
[IQR]



The *last 1000 days* - living with kidney failure

**Making sense of new
(Tasmanian) information**



**Co-design of
new Australian
decision aids**



- What have we done?
- Kidney disease in Tasmania?
 - Epidemiology & growth of chronic disease
 - Competing risk of death or dialysis
 - Co-design of decision aids
- **What next?**

Managing CKD in primary care



CKD Management in Primary Care Handbook

5TH EDITION

The #1 guide for to help detect, manage and refer patients
in your practice with CKD.

CKD-Go! App
New app available
now.

The best bits of the CKD
handbook in an App.



Get your copy!

kidney.org.au/ckdhandbook

Kidney Health[®]
Australia

5th Edition

Chronic Kidney Disease (CKD) Management in Primary Care

Guidance and clinical tips to help identify, manage, and refer
patients in your practice with CKD.



CKD in Australia - Common

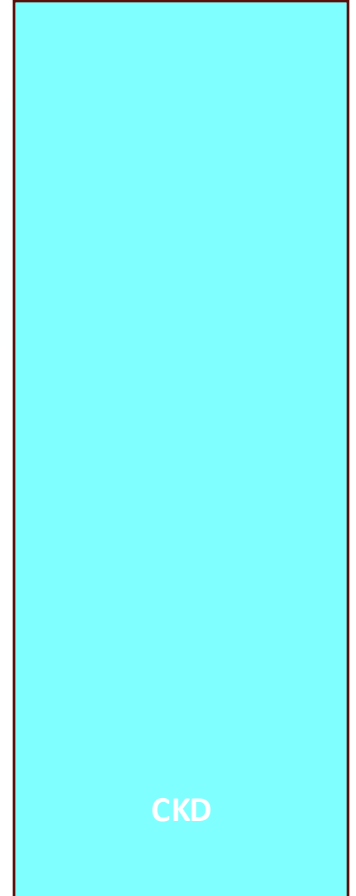


Australian adults (1 in 7) aged 18 years and over have indicators of CKD such as reduced kidney function and/or albumin in the urine.¹



Australian adults have at least one factor increasing their risk of CKD.³

CKD is **twice as common** as diabetes.²



1. Deloitte Access Economics. *Changing the chronic kidney disease landscape: the economic benefits of early detection and treatment*. 2023:62. February 2023. Accessed January 23, 2024. <https://kidney.org.au/get-involved/advocacy/deloittereport>
2. Australian Bureau of Statistics. *Australian Health Survey: Biomedical Results for Chronic Diseases*, 2025.
3. Australian Institute of Health Welfare. *Chronic kidney disease: Australian facts*. 2023.

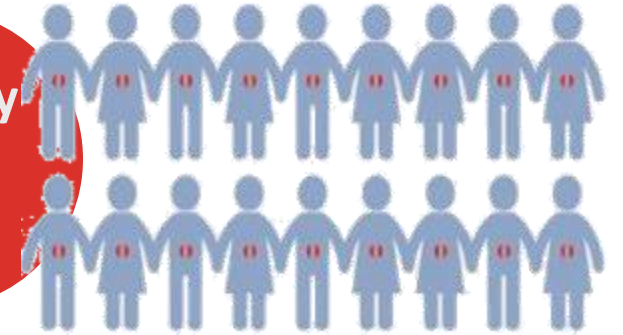
CKD in Australia - Harmful



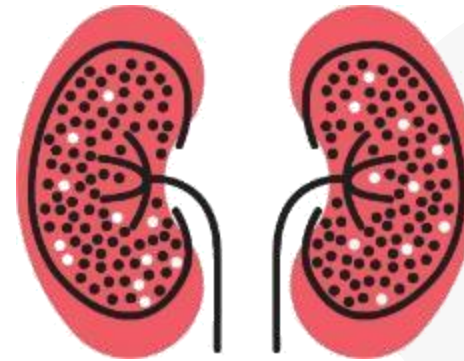
People with CKD are up to **20 times** more likely to **die from a heart attack** or stroke than they are to progress to kidney failure. ²

~20,000

Australians **die every year** with kidney disease. ¹



The number of people needing treatment for kidney failure has **doubled** in the last 20 years. ¹



The **burden of CKD** is greatest in people experiencing socioeconomic disadvantage, living rurally and in First Nations Australians. ¹

1. Australian Institute of Health Welfare. Chronic kidney disease: *Australian facts*. 2023.

2. Tonelli M, Wiebe N, Culleton B, et al. Chronic kidney disease and mortality risk: systematic review. *J AM Soc Neph*. 2006;17:2034-2047.

CKD in Australia - Treatable



If CKD is **detected early** and managed appropriately, deterioration in kidney function can be **reduced by as much as 50%**.¹



New treatments can slow the progression of CKD by up to **15 years**, or potentially longer if started early.²

1. Johnson DW. Evidence-based guide to slowing the progression of early renal insufficiency. *Intern Med J.* 2004;34(1-2):50-57.

2. Fernandez-Fernandez B, Sarafidis P, Soler MJ, Ortiz A. EMPA-KIDNEY: expanding the range of kidney protection by SGLT2 inhibitors. *Clin Kidney J.* Aug 2023;16(8):1187-1198. doi:10.1093/ckj/sfad082

CKD in Australia - Overlooked

< 7%

Fewer than 7% of people with CKD are aware they have this condition. ¹



90% of kidney function can be lost before people experience symptoms. ¹

17%

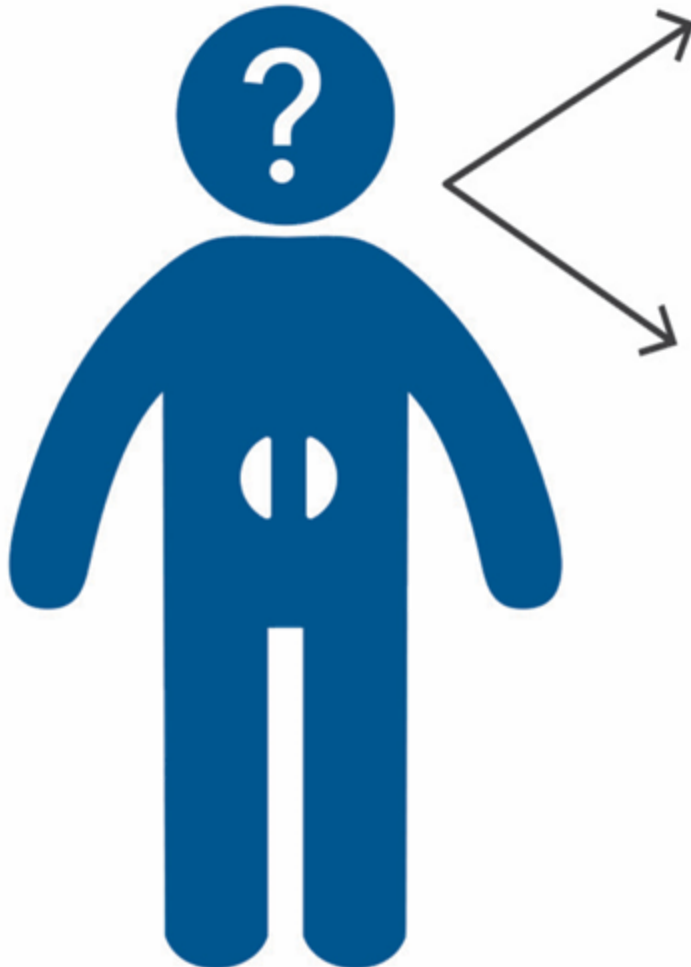
Late referral is common. 17% of people commence dialysis within 90 days of being referred to a kidney service. ²

1. Australian Bureau of Statistics. *Australian Health Survey: Biomedical Results for Chronic Diseases*, 2011-12. 2013.

2. ANZDATA Registry. 46th Report, Chapter 1: Incidence of Kidney Failure with Replacement Therapy. 2023. Accessed December 04, 2023. <https://www.anzdata.org.au/report/anzdata-46th-annual-report-2023-data-to-2022/>

What is CKD?

CKD is defined as...



An estimated or measured glomerular filtration rate (GFR) $<60 \text{ mL/min/1.73m}^2$ that is present for ≥ 3 months with or without evidence of kidney damage.

Or

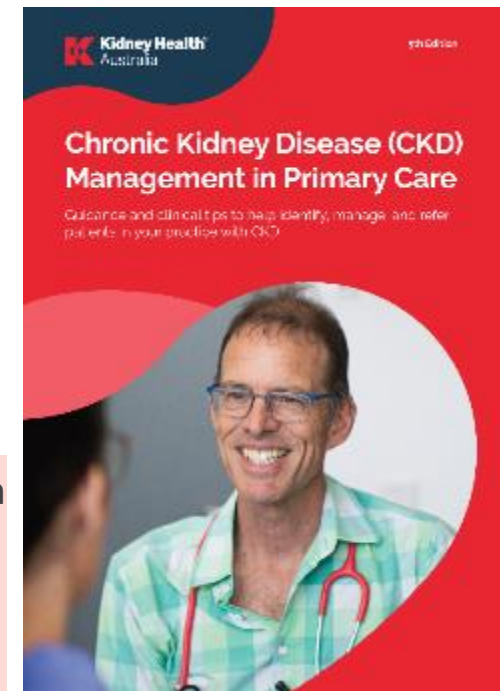
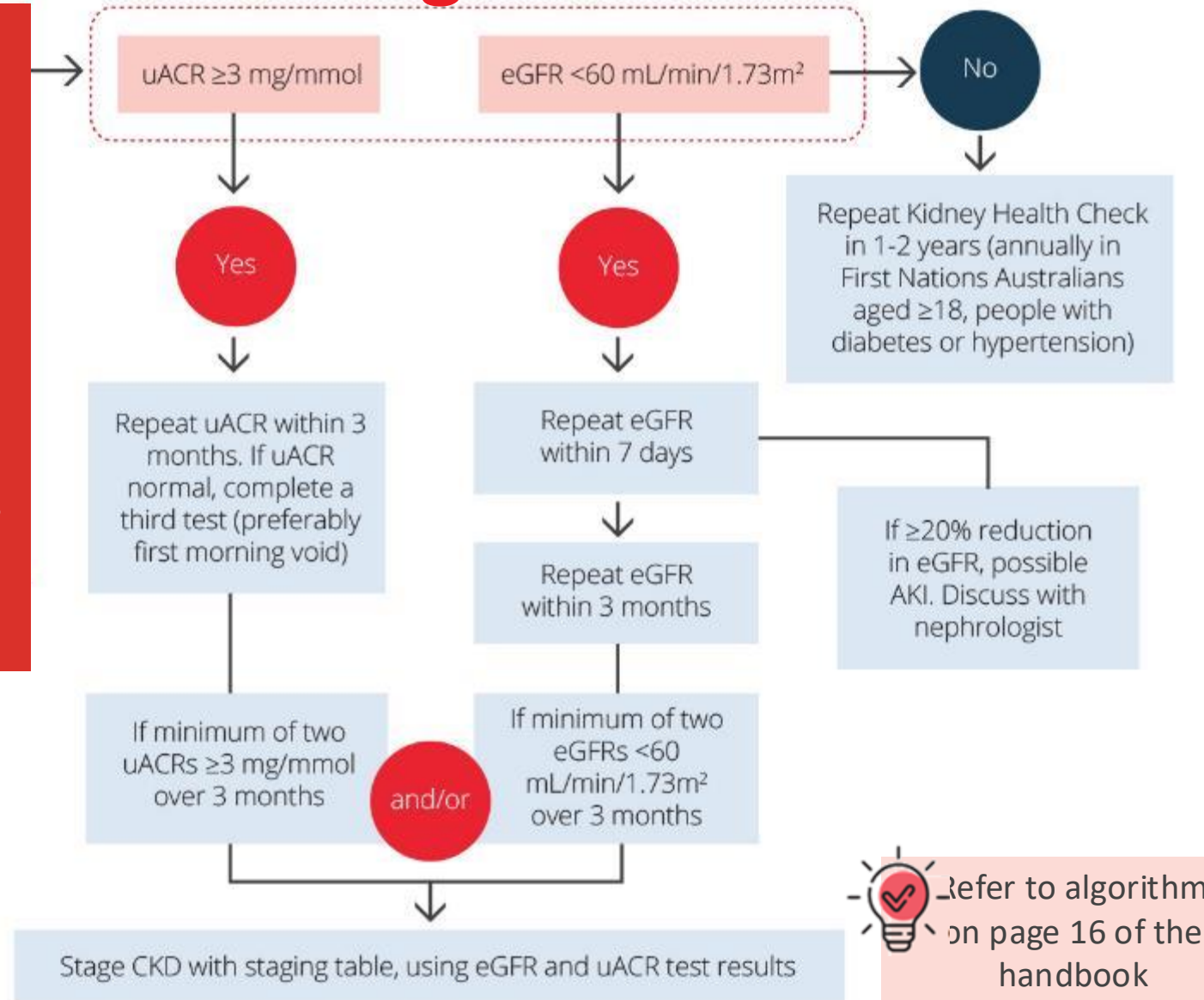
Evidence of kidney damage with or without decreased GFR that is present for ≥ 3 months as evidenced by the following, irrespective of the underlying cause:

- Albuminuria
- Haematuria after exclusion of urological causes
- Structural abnormalities (e.g. on kidney imaging tests)
- Pathological abnormalities (e.g. renal biopsy)

Initial detection and diagnosis of CKD

Offer a Kidney Health Check to people with

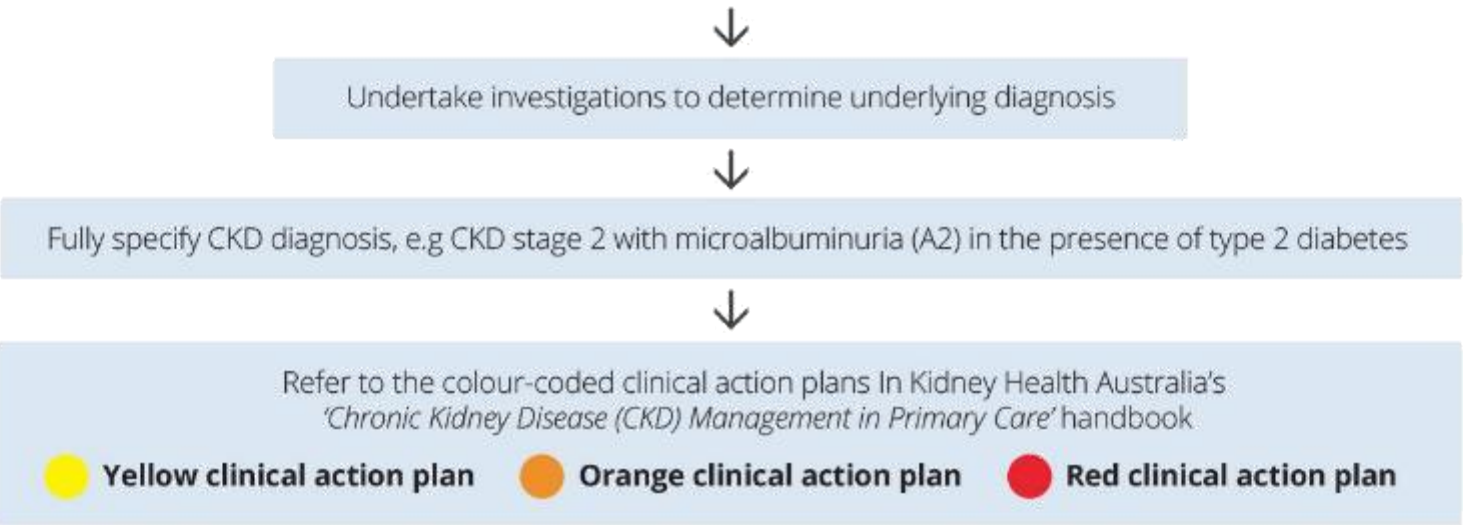
- Diabetes
- Hypertension
- Established CVD
- Family history of kidney failure
- Obesity (BMI ≥ 30)
- Current or former smoker or vaper
- History of AKI
- First Nations Australians aged ≥ 18 years
- All Australians aged ≥ 60 years



Algorithm for initial detection and diagnosis of CKD cont...

↓

Albuminuria Stage				
Kidney Function Stage	GFR (mL/min/1.73m ²)	Normal (A1) uACR <3.0 mg/mmol	Microalbuminuria (A2) uACR 3.0-30 mg/mmol	Macroalbuminuria (A3) uACR >30 mg/mmol
1	≥90	Not CKD unless haematuria, structural or pathological abnormalities present		
2	60-89			
3a	45-59			
3b	30-44			
4	15-29			
5	<15 or on dialysis			



Refer to algorithm on page 16 of the handbook

Enter diagnosis into the practice software as a coded diagnosis

What key management strategies can be implemented in primary care to manage CKD?

- a) Slow decline in eGFR**
- b) Reduce albuminuria**
- c) Maintain blood pressure below 130/80 mmHg**
- d) Lower cardiovascular risk**
- e) Avoid further damage to kidneys**
- f) Nothing. CKD cannot be effectively managed in primary care**

Question

Answer:

Key management strategies for CKD

Slow decline
in eGFR

Reduce
albuminuria

Maintain blood
pressure below
130/80 mmHg

Lower CVD
risk

Avoid further
damage to
kidneys



Clinical tip

Use the corresponding colour-coded action plan for key management strategies and goals on pages 26-28 of the handbook.

Colour-coded action plan

Yellow clinical action plan

eGFR ≥ 60 mL/min/1.73m² with microalbuminuria (A2) or
eGFR 45–59 mL/min/1.73m² with normoalbuminuria (A1)

Management goals

- Slow progression of CKD.
 - Slow decline in eGFR.
 - Reduce albuminuria by at least 30%.
- Assess and lower cardiovascular risk.
- Avoid nephrotoxic medications or volume depletion.
- Encourage positive lifestyle changes and self-management practices.



Management strategies

Frequency of review

- Every 12 months

Clinical assessment

- Blood pressure
- Weight and waist circumference
- Smoking/vaping history

Laboratory assessment

Recommended:

- uACR
- eGFR
- Urea, creatinine, and electrolytes
- Full blood count

Also consider:

- Screening for diabetes (fasting blood glucose or HbA1c)
- HbA1c (for people with diabetes)
- Dipstick urinalysis for haematuria detection
- Lipid studies (Trig, Chol, HDLC, LDLC)

Treatment checklist

- Complete investigations to determine underlying cause of CKD.
- Provide advice on positive lifestyle changes (addressing smoking/vaping, nutrition, alcohol use, physical activity, sleep, stress).
- Maintain blood pressure consistently below target.
- Complete cardiovascular risk assessment.
- Prescribe medications to slow CKD progression, e.g., ACE inhibitor or ARB, SGLT2 inhibitor, non-steroidal MRA.
- Consider lipid lowering treatment where appropriate.
- Optimise glycaemic control.
- Avoid nephrotoxic medications or volume depletion.
- Discuss contraception with individuals of child-bearing age.
- Recommend vaccinations.



Refer to action plans on pages 26–28 of the handbook

Colour-coded action plans

Orange clinical action plan

eGFR 30-59mL/min/1.73m² with microalbuminuria (A2) or
eGFR 30-44 mL/min/1.73m² with normoalbuminuria (A1)



Enter review
reminders and
into practice
software

Management goals

- Slow progression of CKD.
 - Slow decline in eGFR.
 - Reduce albuminuria by at least 30%.
- Assess and lower cardiovascular risk.
- Avoid nephrotoxic medications or volume depletion.
- Encourage positive lifestyle changes and self-management practices.



- Early detection and management of complications.
- Adjust medication doses to levels appropriate for kidney function.
- Appropriate referral to a nephrologist when indicated.



Management strategies

Frequency of review

- Every 3-6 months

Clinical assessment

- Blood pressure
- Weight and waist circumference
- Smoking/vaping history

Laboratory assessment

Recommended:

- uACR
- eGFR
- Urea, creatinine, and electrolytes
- Full blood count

Also consider:

- Screening for diabetes (fasting blood glucose or HbA1c)
- HbA1c (for people with diabetes)
- Dipstick urinalysis for haematuria detection
- Lipid studies (Trig, Chol, HDLC, LDLC)
- Iron studies
- Calcium and phosphate
- Parathyroid hormone (6-12 monthly if eGFR <45mL/min/1.73m²)

Treatment checklist

- Complete investigations to determine underlying cause of CKD.
- Provide advice on positive lifestyle changes (addressing smoking/vaping, nutrition, alcohol use, physical activity, sleep, stress).
- Maintain blood pressure consistently below target.
- Complete cardiovascular risk assessment.
- Prescribe medications to slow CKD progression, e.g., ACE inhibitor or ARB, SGLT2 inhibitor, non-steroidal MRA.
- Consider lipid lowering treatment where appropriate.
- Optimise glycaemic control.
- Avoid nephrotoxic medication or volume depletion and adjust doses to levels appropriate for kidney function.
- Assess for common issues presenting in CKD.
- Appropriate referral to nephrologist when indicated.
- Discuss contraception with individuals of child-bearing age.
- Recommend vaccinations.



Refer to action
plans on page 27
of the handbook

Colour-coded action plan

Red clinical action plan

Macroalbuminuria irrespective of eGFR or
eGFR <30 mL/min/1.73m² irrespective of albuminuria

Management goals

- Slow progression of CKD.
 - Slow decline in eGFR.
 - Reduce albuminuria by at least 30%.
- Assess and lower cardiovascular risk.
- Avoid nephrotoxic medications or volume depletion.
- Encourage positive lifestyle changes and self-management practices.



- Early detection and management of complications.
- Adjust medication doses to levels appropriate for kidney function.
- Appropriate referral to a nephrologist when indicated.



- Prepare for kidney replacement therapy if appropriate.
- Prepare for comprehensive conservative care if appropriate.

Management strategies – as for Orange action plan, plus...

Frequency of review

- Every 1-3 months

Clinical assessment

- Oedema

Treatment checklist

- Address high cardiovascular risk (handbook page 48)
- Discuss potential progression to kidney failure with patient and treatments
- Initiate advance care planning (handbook page 75).

Frequently asked question

What dietary changes could help preserve kidney function?

Please enter your answers in the Chat

Question

Nutrition and diet – lifestyle modification

Target	Detail
Healthy dietary pattern	<ul style="list-style-type: none">• Vegetables, fruit, wholegrains, nuts and legumes, dairy foods, lean meat poultry, fish and plant protein.• Associated with reduced risk of mortality, kidney failure, developing CKD, and progression of CKD• Can reduce rate of kidney function decline, decrease body weight and blood pressure, and metabolic acidosis.
Fluid	<ul style="list-style-type: none">• Make water the drink of choice.• No recommended number of glasses to consume daily.• Drink to thirst.• Avoid sugar sweetened beverages – they have shown to elevate risk of and progression of CKD.
Salt	<ul style="list-style-type: none">• Reduce intake to <5g per day
Ultra-processed foods	<ul style="list-style-type: none">• Avoid foods high in fat, sugar and salt e.g. biscuits, cakes, packaged snack foods, takeaway foods, energy drinks, fruit juices and cordials.



Refer to action plans on page 30 of the handbook



Alcohol – lifestyle modification

Australian guidelines recommend **healthy** men and women should drink no more than 10 standard drinks a week and no more than 4 standard drinks on any one day to reduce the risk of harm from alcohol-related disease or injury.

There are no specific recommendations about safe levels of alcohol consumption people with CKD, however... the less you drink, the lower your risk of harm from alcohol.



Physical activity – lifestyle modification

Target

Be active on most days

Aim for 2.5-5 hours of moderate activity per week

Include muscle strengthening activities at least 2x per week

- Increased physical exercise and reduced sedentary time is associated with improved outcomes for people living with CKD.
- Light to moderate physical activity is a safe starting point of most individuals.



Weight management – lifestyle modification

Overweight (BMI 25.1-30) & obese (BMI >30) people are **40% & 80%** more likely to develop CKD compared to normal weight individuals*

Although not as powerful as diabetes or hypertension, obese people are more likely to develop albuminuria and proteinuria

Central obesity more important than generalised obesity



*Wang Y et al. Association between obesity and kidney disease: a systematic review and meta-analysis. *Kidney Int.* 2008;73:19-33.

Self-management programs

**ARE YOU
LIVING WITH
KIDNEY DISEASE
AND WANT MORE
SUPPORT?**

**EXPRESS YOUR
INTEREST IN THE
KIDNEY HEALTH 4 LIFE
PROGRAM!**

KIDNEY HEALTH 4 LIFE provides support for people at all ages and stages of kidney disease to manage their kidney health and wellbeing.

KIDNEY HEALTH 4 LIFE WILL GIVE:

- Improved disease knowledge.
- Tips and advice to support healthy behaviours and goal setting.
- Practical diet and nutrition support.
- Tailored support for all stages of kidney disease.

REGISTER YOUR INTEREST

Please scan the QR code or call our Kidney Helpline on **1800 454 363**



Kidney Health[®]
Australia

Get the Most Out of Life

Information for health and community workers

TASMANIAN
HEALTH
SERVICE

Get the Most Out of Life is a 6-week group program to help Tasmanians self-manage their chronic conditions and symptoms.

Two trained leaders deliver programs for groups of 10-15 participants.

Who can benefit from the program?

The program is suitable for people with chronic conditions, including, but not limited to:

- heart disease,
- arthritis,
- diabetes,
- asthma,
- bronchitis / emphysema / COPD
- chronic kidney disease
- multiple sclerosis and other neurological conditions.

The program is especially helpful for people with multiple conditions, as it teaches skills to coordinate many of the things needed to manage their health.

Partners / family members / friends / carers are also welcome to participate.

What does the program cover?

Get the Most Out of Life encourages participants to become more active self-managers. They share their experiences of learning new tools and support each other in the process.

Get the Most Out of Life helps participants learn how to:

- manage their symptoms
- get started with healthy eating and physical activity
- communicate effectively with their doctor and healthcare team
- manage difficult emotions
- develop an action plan - a key self-management tool
- be systematic with problem solving and decision making
- self-manage medication responsibilities
- practice relaxation techniques and self-help activities to improve sleep and reduce stress, pain and depression.

What does the research show?

This program was originally developed at Stanford University and is now licensed by the [Self Management Resource Center, USA](#), as the [Chronic Disease Self Management Program](#).

More than 50 research studies have found that people who participate in this program generally:

- have fewer symptoms such as depression and shortness of breath,
- better quality of life,
- they exercise more, and
- use health care less.

Primary Health

TASMANIAN
HEALTH
SERVICE



Research indicates that trained peers (non-professionals) with chronic conditions, can facilitate the program as effectively, if not more effectively, than health professionals. In Tasmania we have a mix of health workers and volunteers as leaders.

How do I refer people?

It's best if people self-refer via the contact details listed below (it is a self-management program after all).

They can speak with a coordinator who can answer questions and register them for the next group or add them to a waiting list.

You can help by giving them a brochure about the program and / or the contact details.

Where are programs run?

Get the Most Out of Life programs are held in venues across NW Tasmania.

People may also be able to access a program online.

Is there a cost?

A gold coin contribution for refreshments is the only cost to attend.

What is the length of the program?

The program runs for 2½ hours each week for six weeks with a group of about 10 - 15 people.

Who is licensed in Tasmania?

In Tasmania programs are delivered by [Tasmanian Health Service](#), and [Aural Health Tasmania](#).

Contact:

Michelle Towle (NW):
Email michelle.towle@ths.tas.gov.au
Phone (03) 6477 7347

May 2021

Dale Anderson's story



Dale Anderson, Food Services Manager at the Launceston General Hospital attended the six-week Get the Most Out of Life program and has not looked back since.

Dale has type II diabetes and says doing the program really made him take stock and look at things differently.

"I'm more comfortable since doing the program - I was in denial for a few years after my initial diagnosis. I'd even go to different doctors just to get the scripts I needed so I avoided having the conversation about changes I needed to make."

"The program made me realise I needed to be more responsible about my medication, I needed to look after myself better, and seeing other people in worse situations made me realise others do experience difficulties. I sit back and listen more and am definitely more aware and not as judgemental of others."

I really enjoyed every aspect of the program and have already recommended the program to others."

Pillars of medication management

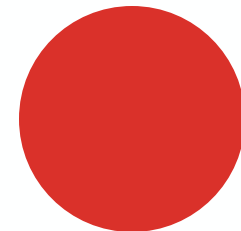


Angiotensin-converting enzyme inhibitors (ACEi) are better than angiotensin receptor blockers (ARB) in reducing albuminuria and kidney function decline.

- a) True
- b) False**
- c) Don't know

ACE inhibitor **or** ARB is recommended first line therapy in primary care.

Combined ACE **and** ARB should **only** be used **under specialist supervision**



Hypertension target



For all people with CKD...
maintain BP below

130/80 mmHg

Treatment should always be individualised and in some patients, it may be appropriate to aim for a lower BP target

Treatment targets should take into account the risk / benefit scenario along with clinical practicalities

Blood pressure reduction



ACE inhibitor
or ARB is
recommended
first line
therapy.

Reducing blood
pressure to
below target
levels is one of
the most
important goals
of CKD
management

Lifestyle
changes
should always
be advocated
and can have
significant
effect on BP

Hypertension
may be difficult
to control, and
multiple (3 or
more)
medications
are frequently
required



Refer to pages 52-56 in
the CKD Handbook

RAS blockade and kidney function



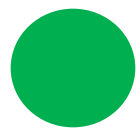
Clinical tip

ACE inhibitors and ARBs cause a reversible reduction in GFR when treatment initiated.

Check eGFR within 2 weeks following initiation.

Provided reduction is $< 25\%$ within 2 weeks of starting therapy, the ACE inhibitor or ARB should be continued.

If reduction is $>$ than 25% below the baseline value, discontinue and consider referral to a nephrologist.



PRESCRIBE: medications to slow CKD progression and reduce CVD risk

ACE inhibitor or ARB	Statin (+/- ezetimibe)	SGLT2 inhibitor*	Non-steroidal MRA*	GLP-1 RA*
<ul style="list-style-type: none">• First-line treatment.• Up-titrate to maximum tolerated dose.	<p>Consider use in:</p> <ul style="list-style-type: none">• People with CKD and a CVD risk $\geq 10\%$ and• First Nations Australians with CKD and a CVD risk $\geq 5\%$.	<ul style="list-style-type: none">• Use in people with CKD and proteinuria, with/without diabetes*.• Do not initiate if eGFR $< 25\text{mL/min/1.73m}^2$.	<ul style="list-style-type: none">• Indicated for use in people with CKD (with albuminuria) associated with type 2 diabetes.• Do not initiate if eGFR $< 25\text{mL/min/1.73m}^2$ or when K$^+$ $> 5.0\text{mmol/L}$.	<ul style="list-style-type: none">• Indicated for use in people with CKD if they also have type 2 diabetes.• Do not use in people with kidney failure.

* Refer to product information for eligibility criteria and dosing

SGLT2 inhibitors e.g. dapagliflozin (e.g. Forxiga), empagliflozin (e.g. Jardiance)

Non-steroidal mineralocorticoid receptor agonist (MRA) = finerenone (Kerendia)

Glucagon-like peptide- 1 receptor agonist (GLP-1 RA) (e.g. semaglutide (e.g. Ozempic)



Refer to pages 34-35 in the CKD Handbook

Statin therapy for CKD

Fasting lipid profile evaluation is recommended for all adults with newly identified CKD. Consider secondary causes and specialist evaluation if LDL-cholesterol >4.9 mmol/L or triglycerides >11.3 mmol/L.

Statin (+/-ezetimibe) for:

- Non-Indigenous people with CKD (eGFR ≥ 15 ml/min/1.73m²) and CVD risk $\geq 10\%$
- First Nations Australians with CKD and CVD risk $\geq 5\%$.

Lifestyle advice if hypertriglyceridaemia is present.

Refer to: CARI Guidelines: Management of cholesterol-lowering therapy in people with chronic kidney disease.



SGLT2 inhibitors treatment for CKD

-
- PBS Criteria:
- Diagnosis of proteinuric CKD (with or without diabetes) present for ≥ 3 months prior to prescribing.
- eGFR 25 - 75 mL/min/1.73m²
- uACR 22.6 - 565 mg/mmol
- Must be stabilised, for at least 4 weeks, on either: (i) an ACE inhibitor or (ii) an angiotensin II receptor antagonist.
- Do not use in combination with another SGLT2 inhibitor.
- Not recommended to initiate if eGFR < 25 mL/min/1.73m².
- May be prescribed by nurse practitioners (continuing therapy only)

SGLT2 inhibitors



Clinical tip

- SGLT2 inhibitors cause a reversible drop in eGFR 4 weeks after initiation, then rebounds.
- Specific testing of eGFR for this purpose is not required.
- SGLT2 inhibitors cause osmotic diuresis, reduce diuretics and/or antihypertensive medications upon initiation of an SGLT2 inhibitor.

SGLT-2 inhibitors slow CKD progression

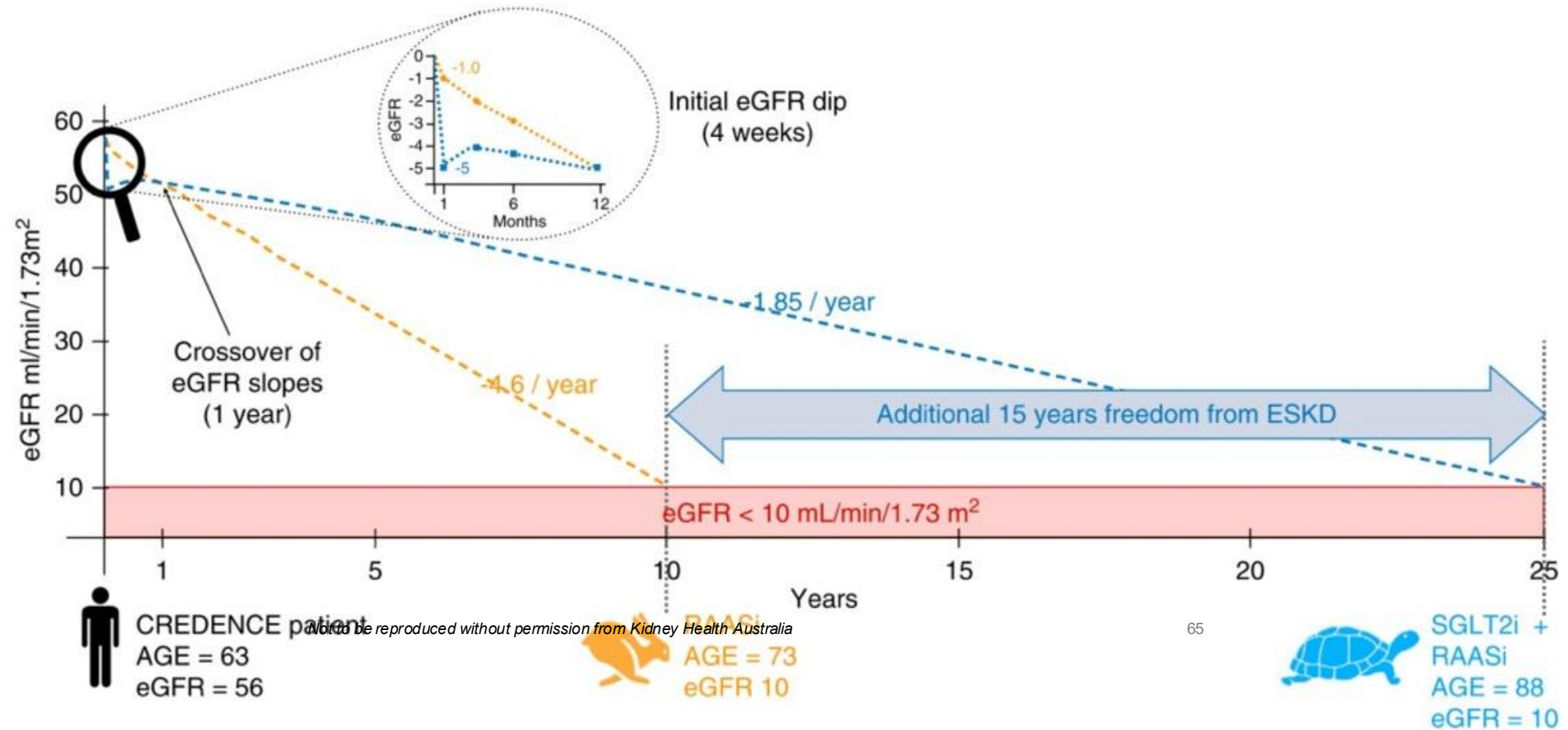


Figure 1. | SGLT2is may delay ESKD by 15 years. A typical patient included in CREDENCE would lose 4.6 ml/min per year of eGFR if treated with RAASi only, reaching ESKD in 10 years. However, if canagliflozin is added to his treatment, he would only lose 1.85 ml/min per year of eGFR, delaying ESKD by 15 years. RAASi, renin-angiotensin-aldosterone system inhibitors; SGLT2i, sodium-glucose cotransporter 2 inhibitor.

Diabetic kidney disease and CKD therapy

Non-steroidal mineralocorticoid antagonist (nsMRA), finerenone, is PBS listed for diabetic kidney disease (DKD) to delay progressive decline of kidney function and reduce risk of CV event in addition to standard care.

Tips for using nsMRA

- PBS approved for use in diabetic kidney disease

eGFR 25 - 75 mL/min/1.73m²

uACR 22.6 - 565 mg/mmol

Add on therapy to both RAS (ACEi or ARB) and SGLT2 inhibitor

Predictable drop in eGFR and rise in serum potassium - monitor carefully

Do not use in combination with steroidal MRAs (e.g. spironolactone)





REDUCE: medications excreted by the kidneys

Medications that may need to be started at a reduced dose or ceased in patients with CKD include but not limited to#:

Anti-infective	Cardiovascular	Diabetes	Pain	Other
<ul style="list-style-type: none">• famciclovir• nirmatrelvir• valaciclovir• antibiotics e.g. ciprofloxacin, trimethoprim, sulfamethoxazole, aminoglycosides, nitrofurantoin	<ul style="list-style-type: none">• apixaban• dabigatran• digoxin• rivaroxaban• sotalol• spironolactone	<ul style="list-style-type: none">• acarbose• all gliptins except linagliptin• insulin• metformin*• sulfonylureas	<ul style="list-style-type: none">• gabapentin• opioid analgesics• pregabalin	<ul style="list-style-type: none">• allopurinol• benzodiazepines• colchicine• baclofen• duloxetine• escitalopram• solifenacin• fenofibrate• denosumab^• lithium

* Metformin reduce dose if eGFR 30-60mL/min/1.73m² and under specialist supervision if eGFR<30mL/min/1.73m²

Medications in **red** have been added to the table in the 5th edition



Refer to pages **34-35** in the CKD Handbook

AVOID: nephrotoxic medications

Commonly prescribed drugs that can adversely affect kidney function in CKD	Commonly prescribed drugs that should be avoided temporarily during a sick day (SADMANS)*
<ul style="list-style-type: none">• Lithium• Aminoglycosides• NSAIDs/COX-2 inhibitors – beware of the ‘triple whammy’	<ul style="list-style-type: none">• Sulfonylureas• ACE inhibitors• Diuretics• Metformin• ARBs• NSAIDs• SGLT2 inhibitors

- *As part of a sick day action plan, it is important that patients are advised to seek guidance from their healthcare professional on temporarily stopping medications during periods of illness.*
- *More on Sick Day action plans later...*



Refer to pages **34-35** in the CKD Handbook



GPs and pharmacists need to discuss appropriate pain relief medication with patients

Use of radiographic contrast agents in the context of CKD

Recent observations studies questions the severity of contrast-induced nephropathy (CIN).

Benefit of the imaging test and risk of CIN needs to be considered individually.

Recommended CIN risk classification for **adults**:

- eGFR ≥ 45 mL/min/1.73m² low risk
- eGFR 30-40 mL/min/1.73m² intermediate risk, diabetes increases risk
- eGFR < 30 mL/min/1.73m² high risk.

Sick day action plan

- Another important element of a CKD management plan is a Sick Day Action Plan

- ✓ People with CKD stage 3-5 are at increased risk of AKI.
- ✓ Avoid NSAIDs and other nephrotoxic medications
- ✓ Identify early people with acute illness (e.g., GI upset or dehydration)
- ✓ Temporarily cease ACE inhibitors, ARBs, diuretics with hypovolaemia / hypotension

Mnemonic for drugs to be avoided on a sick day (SADMANS)

- S** Sulfonylureas
- A** ACE-inhibitors
- D** Diuretics
- M** Metformin
- A** Angiotensin receptor blockers
- N** Non-steroidal anti-inflammatory
- S** SGLT2 inhibitors

How to guides - Sick Day Action Plan

Sick Day Action Plan (template)

NEW

Being prepared for times of illness is an important element in CKD management and care.



4 Sick Day Action Plan

Download this template and complete for people with CKD.

Contacts				
Doctor: Name:		Phone:		
Pharmacy: Name:		Phone:		
Family: Name:		Phone:		
When I am...	Health care	Medications	Self-care	Resources
 Dehydrated (vomiting, diarrhea, extreme heat)	 Contact your doctor. Contact a family member.	 Stop taking medications:	 Rest. Drink water so that you are passing urine every 2-3 hours and that it is straw coloured. Stay calm and contact family/ carer for assistance. Ask your GP to complete a Kidney	 Drink Water Instead factsheet.

'How to guides' available in the Kidney Health Professional Hub

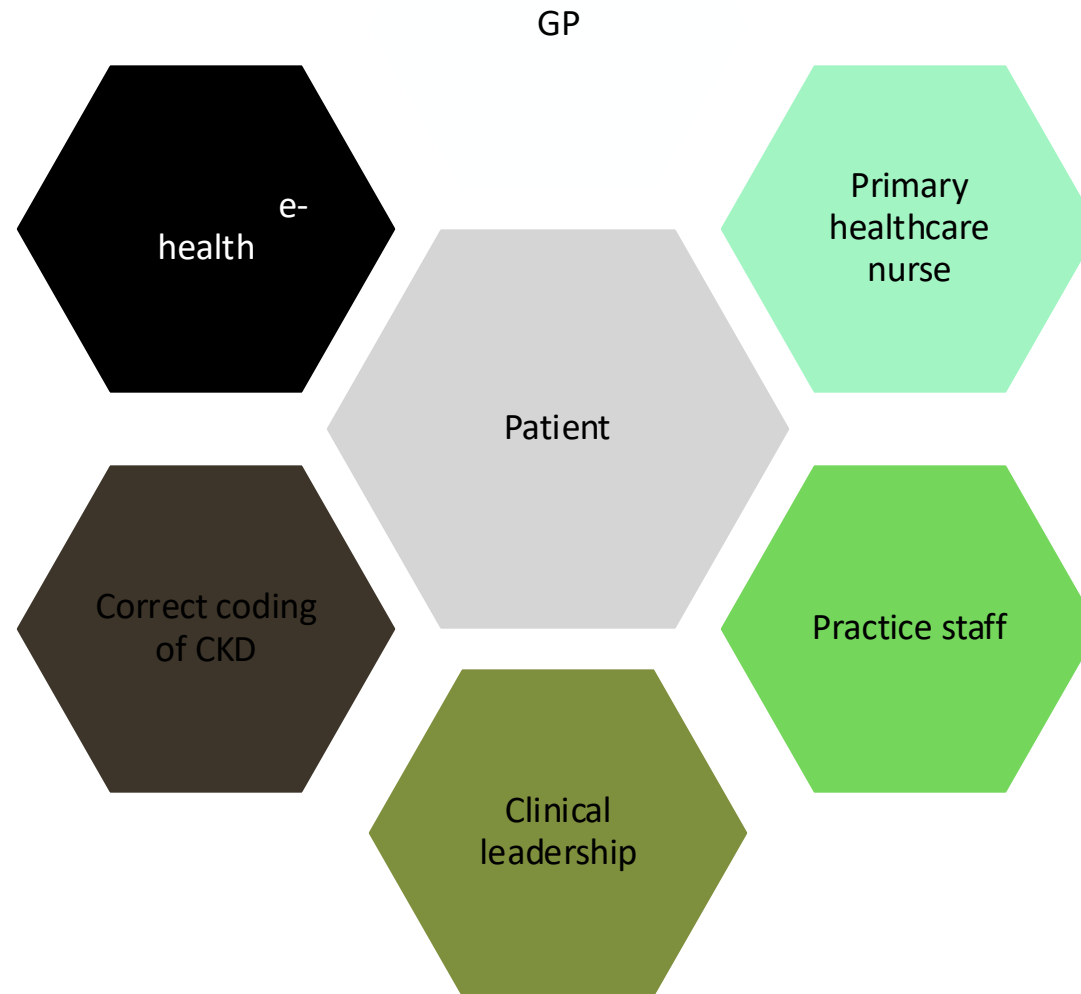


Page 40-42 of the CKD Handbook

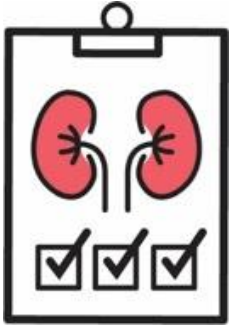
Treatment targets for people with CKD - summary

Parameter	Target	Treatment
Blood pressure	$\leq 130/80$ mmHg	Lifestyle modification ACE inhibitor or ARB
Albuminuria	uACR reduction of at least 30%	ACE inhibitor or ARB
Lipids	No target lipid level is recommended	Dietary advice Statins
Blood glucose (for people with diabetes)	HbA1c $\leq 7.0\%$ / ≤ 53 mmol/mol BGL 6-8 mmol/L (fasting) / 8-10 mmol/L (postprandial)	Lifestyle modification Oral hypoglycaemic Insulin SGLT2 inhibitor and nsMRA

Whole of practice approach to CKD best practice management



Take home messages



Early detection and management is important because CKD is common, harmful, **treatable** and often overlooked.



Actively identify people at risk of CKD in your practice.



Apply to your practice:

- ✓ Algorithm for the initial detection
- ✓ Colour coded clinical action plans to guide CKD management
- ✓ PRESCRIBE, REDUCE AND AVOID classification of medications
- ✓ Sick Day Action Plans



Remember: actively search for CKD indicators and code it in your practice software.



Support

Contact Kidney Health Australia
for non-medical advice,
information, and support.

1800 454 363

Kidney.helpline@kidney.org.au

kidney.org.au

Kidney Health Resources



Treatment options series

Make informed choices about kidney disease treatment options. The series of 'An Introduction to' booklets includes topics on: Treatment Options, Haemodialysis, Peritoneal Dialysis, Comprehensive Conservative Care, Kidney Transplantation, Kidney Donation by Living Donors, and Withdrawing from Dialysis.

First Nations Peoples

Various factsheets, kidney stories toolkit, and flipchart for clinics available to download.



Download
resources



SCAN ME



Eating Out Guide

General advice about good food choices, options, and substitutes when eating out.



Back on the Menu

Easy to follow recipes for a reduced potassium diet.



Dining In

Delicious recipes developed for people with kidney disease.



Living with kidney failure

A practical guide providing a wealth of information about kidney disease, written in Australia, for Australians.



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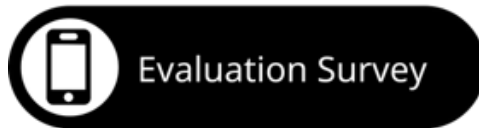
**Kidney Health
Australia**

kidney.org.au

Kidney Helpline: 1800 454 363

Thank you for participating!

1. **Please complete the evaluation** form via the QR code on the screen or on the case study handout.
2. **Follow Kidney Health Australia** on Facebook, LinkedIn and X.



kidney.org.au

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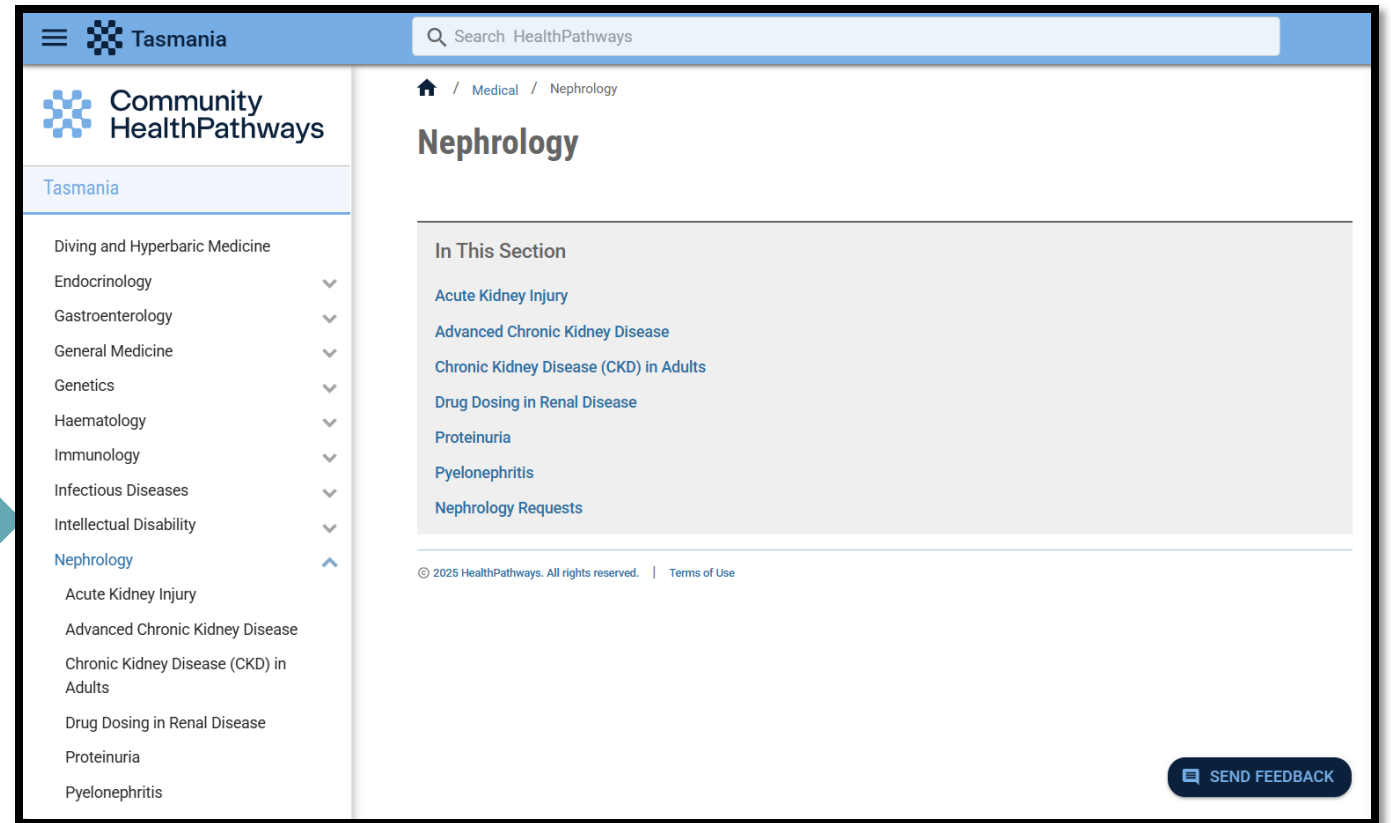
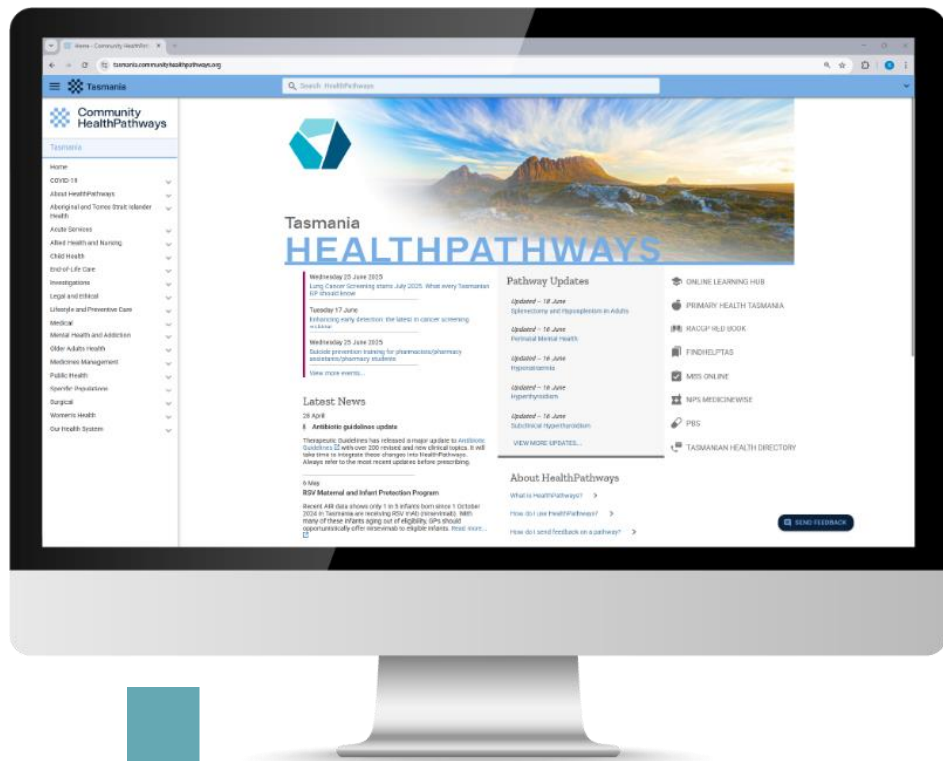


Tasmanian HealthPathways

is a web-based information portal developed by Primary Health Tasmania. It is designed to help primary care clinicians plan local patient care through primary, community and secondary healthcare systems.



tasmania.communityhealthpathways.org



SCAN THE QR CODE OR GO TO
<https://tasmania.communityhealthpathways.org/>

Some final words

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