Changing the narrative in Indigenous kidney health: undertaking strengths-based analysis using linked data

Amandi Hiyare

Flinders University, South Australia, Australia 27th September, 2024

Kylie-Ann Mallitt, Eleonora Dal Grande, Siah Kim, Victoria Sinka, Michelle Dickson, Armando Teixeria-Pinto, Allison Jaure, Germaine Wong, Natasha Nassar, David Lyle, Stephen I Alexander, Jacqueline H Stephens, Jonathan C Craig.



Acknowledgement of Country



Kaurna Yarta, City of Adelaide, South Australia

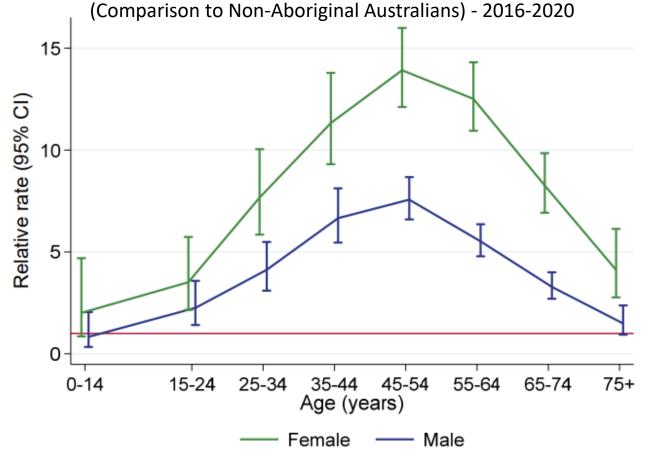


Three Sisters, Gundungurra and Darug land, Blue Mountains National Park, NSW

Background



Figure 1 - Relative Incidence Rate of Treated Kidney Failure for Aboriginal Patients by Gender



Source: ANZDATA Annual Report 2021

Missing longitudinal, population-based studies for Aboriginal and non-Aboriginal children in Australia to understand the trajectory of chronic kidney disease and to inform strategies to prevent its progression.



The ARDAC Story





16-month-old
Aboriginal baby
presented with kidney
failure [received kidney
transplant]

Early 2000s

Aboriginal Health
Education Officer – Ms
Rita Williams initiated
the ARDAC study



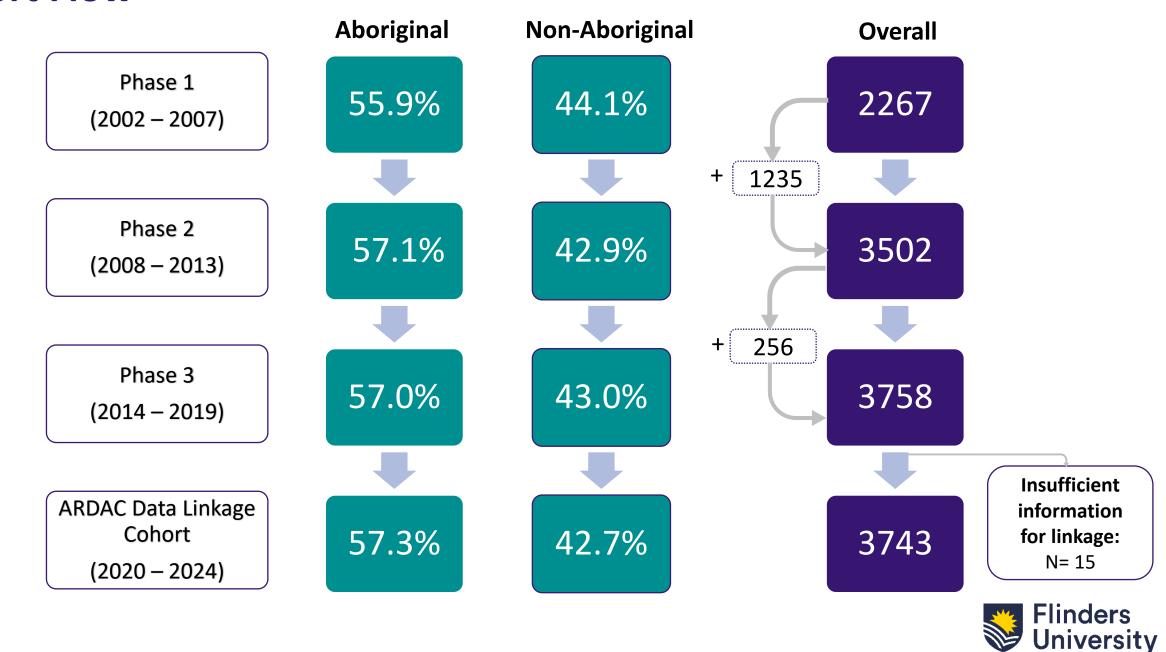
In 2002

Previous findings from ARDAC suggest kidney health may be preventable in childhood

2002 onwards



Cohort Flow



AIMS

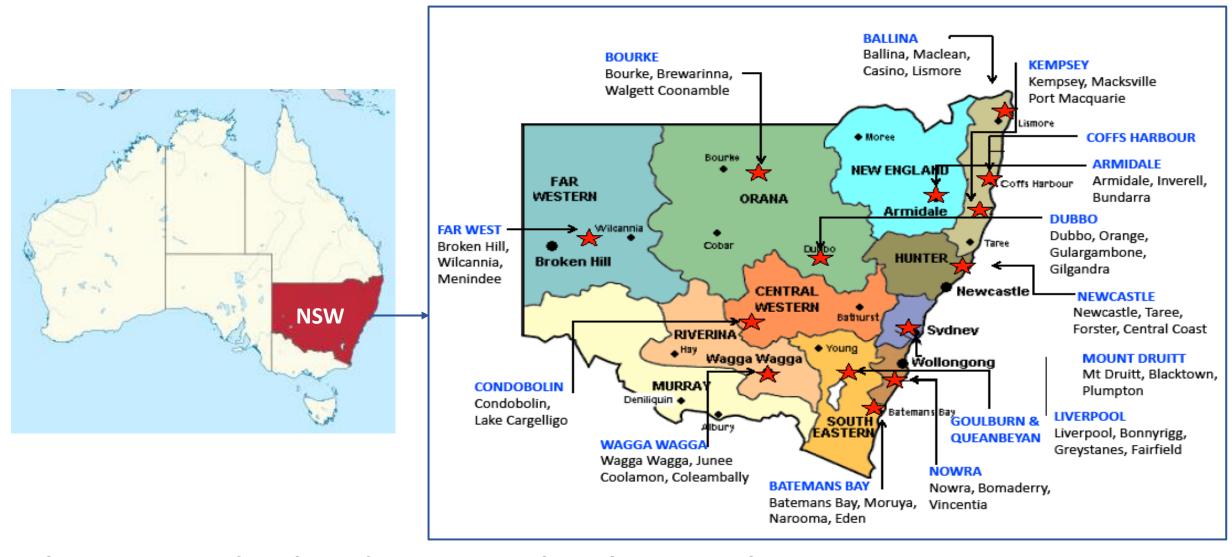
 To estimate the cumulative incidence of chronic kidney disease (CKD) among young Aboriginal people.

 To understand risk factors for development over time.



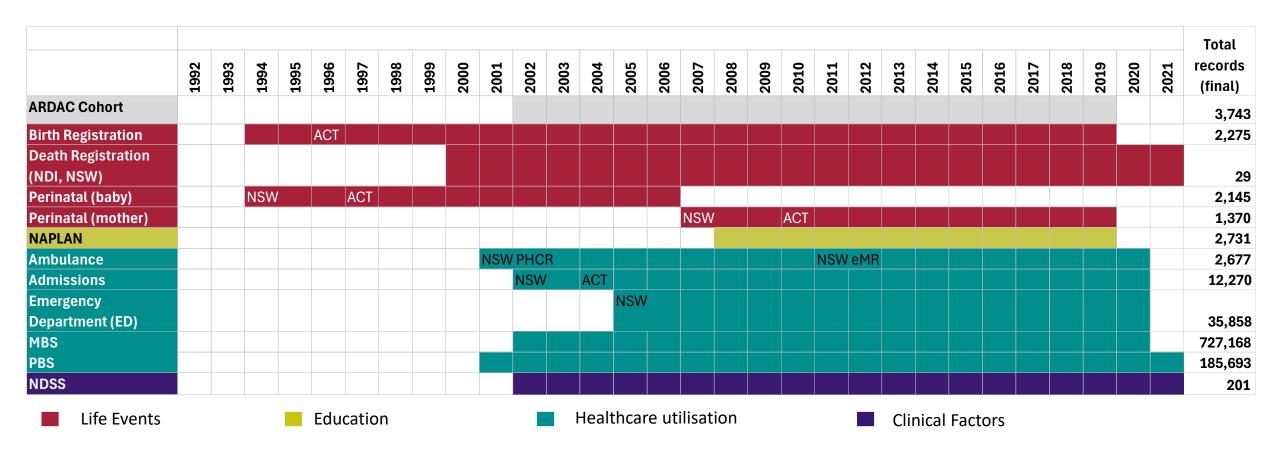


Participating communities



- The ARDAC study is based in New South Wales, Australia
- Over 60 participating communities

Data Linkage



- ARDAC biomedical cohort follow up time: 14.5 years
- Data Linkage follow up time: 27 years



CKD Case Definition

At least **ONE** of the following criteria:

- An uACR Measurement of > 3.4mg/mmol PLUS prescribed a CKD related medication
- Primary or secondary diagnosis for CKD within NSW & ACT hospitals
- Accessed a CKD-related Medicare service
- Self-report in the ARDAC sociodemographic survey
- Main or other cause of death as 'CKD'





Baseline characteristics

Characteristic	Non-Aboriginal (n = 1593)	Aboriginal (n = 2150)	
Female	781 (49%)	1067 (50%)	
Age, median (IQR)	10.8 (8.5 – 13.3)	11.0 (8.2 – 13.6)	
SES [IRSAD]*, mean (SD)	933 (55)	921 (59)	
Major cities	339 (21%)	343 (16%)	
Remote/very remote living*	82 (5.1%)	337 (16%)	
Obese*	186 (12%)	357 (17%)	
Birthweight [grams], mean (SD)	3349 (578)	3212 (627)	
Gestational age [weeks], mean (SD)	39.2 (1.70)	39.0 (2.09)	

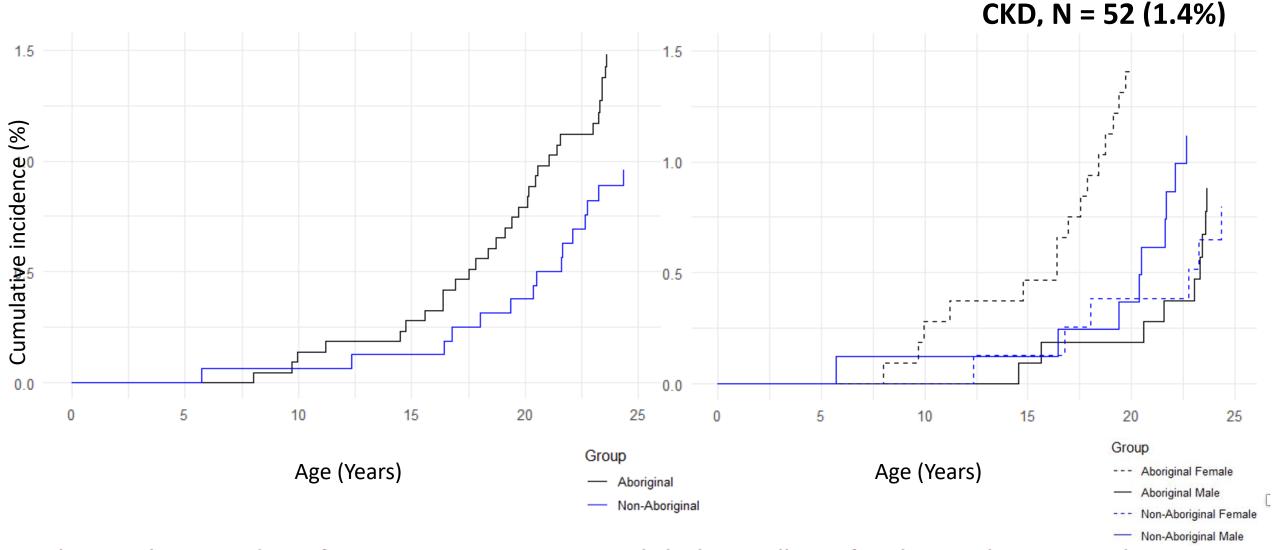
^{*} **Obesity** – Based on CDC cut-offs and percentiles



^{*} Remoteness – Based on 2011 Rural, Remote and Metropolitan Area

^{*} IRSAD – Index of Relative Socio-economic Advantage and Disadvantage

Cumulative Incidence of CKD for Aboriginal and non-Aboriginal young people



- The cumulative incidence from age 10 years is consistently higher at all ages for Aboriginal young peoples
- Young Aboriginal females have a higher cumulative incidence across all groups

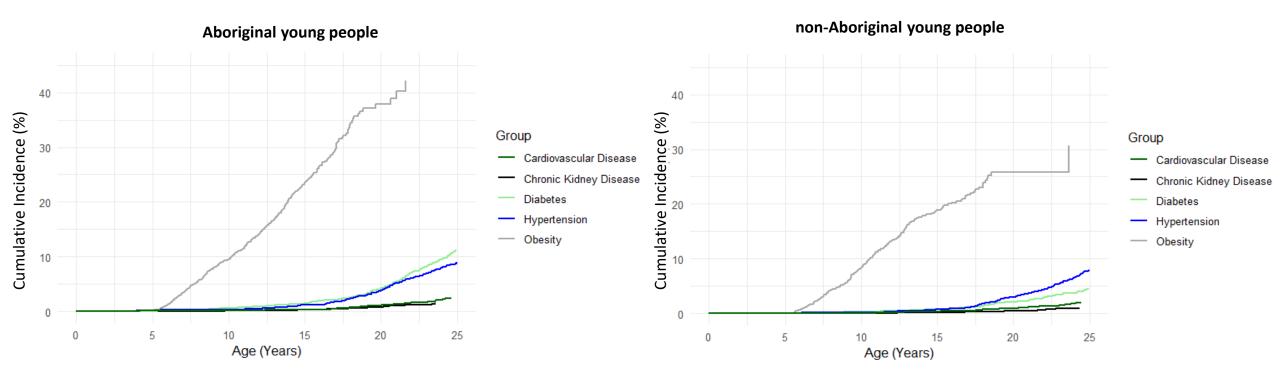


Sociodemographic Risk Factors for Chronic Kidney Disease

		95% CI		
	Adjusted Hazard Ratio	Lower	Upper	P-value
Non-Aboriginal [ref]	1.00			
Aboriginal	1.47	0.81	2.69	0.21
Male [ref]	1.00			
Female	1.51	0.87	2.63	0.14
SES (Per 10 units of IRSAD Scale)	0.99	0.99	1.01	0.75
Major cities [ref]	1.00			
Inner regional	2.22	0.76	6.53	0.14
Outer Regional	3.67	1.18	11.4	0.02
Remote/Very Remote	3.57	1.04	12.3	0.04
No obesity prior to CKD [Ref]	1.00			
Obesity prior to CKD	1.92	1.08	3.41	0.02



Cumulative Incidence of Associated Risk Factors and CKD



- The cumulative incidence of associated risk factors Cardiovascular disease, diabetes, hypertension and obesity are all higher than CKD for both Aboriginal and non-Aboriginal young people
- The cumulative incidence of these associated risk factors is slightly elevated in Aboriginal young people



^{*} Obesity – Based on CDC cut-offs and percentiles

^{*} Diabetes and CVD - Created using various data sources

^{*} Hypertension – Creating using SBP/DBP values measured at biomedical screenings

Conclusions

- ARDAC study fills the missing biomedical data for CKD Young Aboriginal peoples have an increased risk of CKD
- Importance of tailored interventions The cumulative incidence is higher in young Aboriginal females
- Modifiable risk factors Obesity increases the risk of CKD by nearly 2 times
- Risk factors for CKD such as diabetes, obesity, hypertension and CVD are elevated in the early years in young Aboriginal peoples.
- Importance of earlier screening and increased funding for communities.



Acknowledgements

ARDAC Investigators

- Prof Jonathan Craig*
- **Prof Allison Jaure**
- A/Prof Michelle Dickson
- Prof Armando Teixeira-Pinto*
- Prof David Lyle
- **Prof Germaine Wong**
- Prof Natasha Nassar
- A/Prof Jacqueline Stephens*
- Dr Kylie Ann-Mallitt *

ARDAC Research Team

- Dr Eleonora Dal Grande
- Ms Victoria Sinka
- Dr Siah Kim

ARDAC Advisory Committee ARDAC Study Participants

Visit our website:

https://www.ardac.org.au/



Ms. Rita Williams

Aboriginal Health Education Officer, Children's Hospital at Westmead



















Curtin University











- Flinders University Research Student **Travel Grant**
- PHAA SA Award
- **HDA Travel Grant**

















WESTERN SYDNEY UNIVERSITY









CKD Specific ICD-10 Codes (Used for Deaths and Hospital Admissions)

CKD Stage	ICD-10 Description	ICD-10 Codes
	Chronic kidney failure	N18
Stages 1-4	Chronic kidney disease, stage 1	N18.1
	Chronic kidney disease, stage 2 (mild)	N18.2
	Chronic kidney disease, stage 3	N18.3
	Chronic kidney disease, stage 4 (severe)	N18.4
	Hypertensive chronic kidney disease with Stage 1 through 4 chronic kidney disease, or unspecific chronic kidney disease	I12.9
	Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease	I13.0
	Hypertensive heart and chronic kidney disease without heart failure	l13.1
	Hypertensive heart and chronic kidney disease with heart failure and with stage 4 chronic kidney disease, or end stage kidney disease	I13.2
Stage 5	Chronic kidney disease, stage 5	N18.5
	End-stage renal disease	N18.6
	Hypertensive chronic kidney disease with stage 5 chronic kidney disease or end stage kidney disease	I12.0
Stage 5 Regular dialysis	Preparatory care for dialysis	Z49.0
	Haemodialysis	Z49.1
	Peritoneal dialysis	Z49.2
	Kidney transplant and dialysis status	Z94.0, Z99.2
	Complications related to dialysis and kidney transplant	T82.4, T86.1

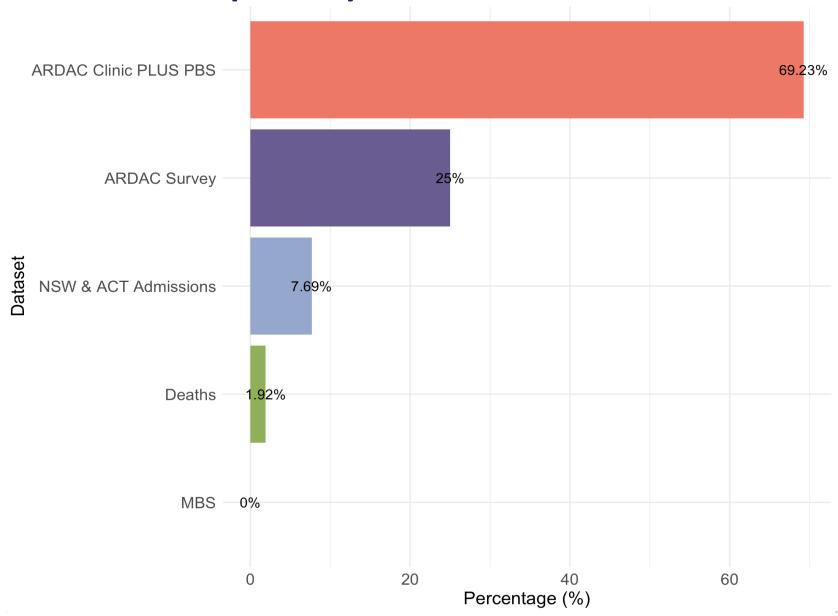
CKD Specific ICD-10 Codes

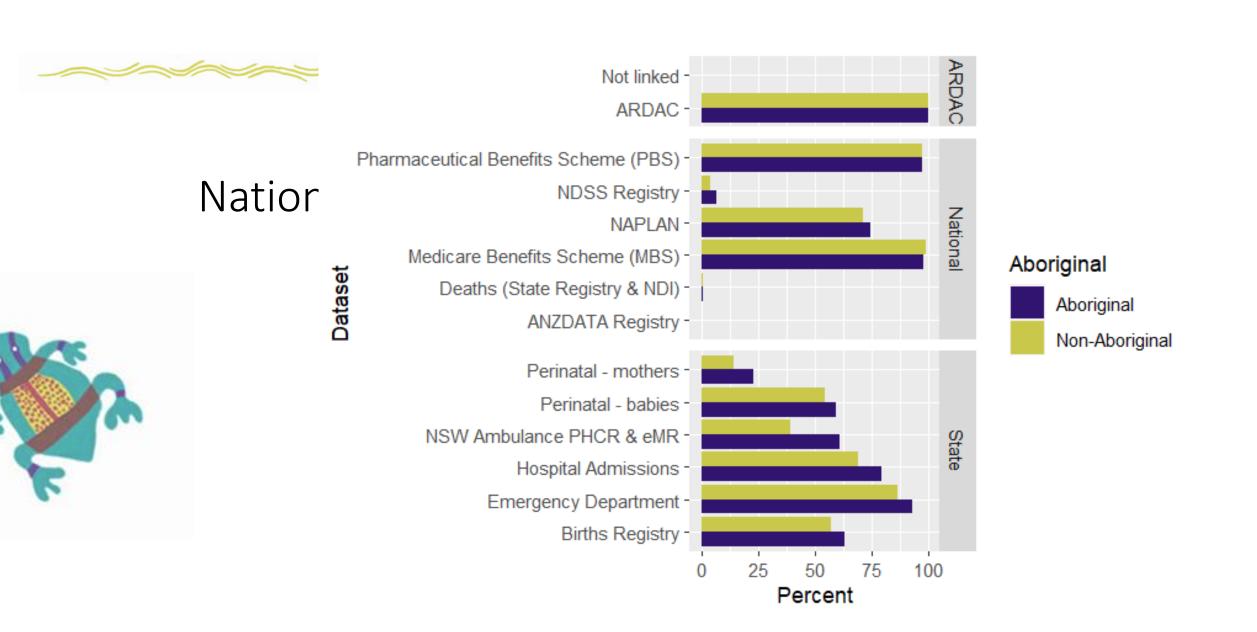
MBS Number	Category	Group	Subgroup	Description
13100-13110	3	T1 – Miscellaneous therapeutic produces	2 – Dialysis	SUPERVISION IN HOSPITAL by a medical specialist of haemodialysis, haemofiltration, haemoperfusion or peritoneal dialysis, including all professional attendances, where the total attendance time on the patient by the supervising medical specialist exceeds 45 minutes in 1 day
36503	3	T08 – Surgical Operations	5-UROLOGICAL	RENAL TRANSPLANT (not being a service to which item 36506 or 36509 applies)
36543	3	T08 – Surgical Operations	5-UROLOGICAL	Nephrolithotomy or pyelolithotomy, or both, extended, for one or more renal stones, including one or more of nephrostomy, pyelostomy, pedicle control with or without freezing, calyorrhaphy or pyeloplasty
36506	3	T08 – Surgical Operations	5-UROLOGICAL	RENAL TRANSPLANT, performed by vascular surgeon and urologist operating together vascular anastomosis including aftercare
36509	3	T08 – Surgical Operations	5-UROLOGICAL	RENAL TRANSPLANT, performed by vascular surgeon and urologist operating together ureterovesical anastomosis including aftercare
66671	6	P2 – Chemical		Quantitation of serum aluminium in a patient in a renal dialysis program - each test

CKD Specific Medications

Code	Code & Prescriber	Description		
Angiotensin converting enzyme (ACE) Inhibitors or Angiotensin Receptor Blockers (ARBs)				
CO9A		ACE Inhibitors, plain		
C09B		ACE Inhibitors, combinations		
		ACE inhibitors and diuretics:		
		Enalapril + Hydrochlorothiazide		
C09BA		Fosinopril + Hydrochlorothiazide		
		Perindopril + Indapamide		
		Quinapril + Hydrochlorothiazide		
C09C		Angiotensin II Receptor Blockers (ARBs), plain		
C09D		Angiotensin Ii Receptor Blockers (ARBS), combinations		
Non-loop diure	etics and loop diuretics			
C03AA	1484D	Thiazides, plain		
C03CA		Sulfonamides, plain		
CU3CA		Furosemide (Frusemide)		
Beta blockers				
C07AB	1081X, 2243C	Atenolol		
	8732N, 8733P, 8734Q, 8735R	Metoprolol succinate		
	1324Q, 1325R	Metoprolol tartrate		
Calcium Channel Blockers				
C08C		Selective calcium channel blockers with mainly vascular effects		
C08D		Selective calcium channel blockers with direct cardiac effects		

CKD % in each dataset (n = 52)





Risk Factors for Chronic Kidney Disease [Univariate]

Chronic Kidney Disease, N = 52		95% CI		
	Adjusted Hazard Ratio	Lower	Upper	P-value
Non-Aboriginal [ref]	1.00			
Aboriginal	1.72	0.96	3.11	0.07
Male [ref]	1.00			
Female	1.51	0.87	2.63	0.14
SES (Per 10 units of IRSAD Scale)	0.99	0.99	1.00	0.17
Major cities [ref]	1.00			
Inner regional	2.18	0.76	6.21	0.15
Outer Regional	3.77	1.23	11.6	0.02
Remote/Very Remote	4.17	1.28	13.6	0.02
No obesity prior to CKD [Ref]	1.00			
Obesity prior to CKD	1.93	1.09	3.41	0.02



Strengths and limitations

• Strengths

- Investigator-advisory committee nexus
- Large proportion of young Aboriginal peoples in our cohort
- Follow-up time increased from 14.5 to 27 years due to data linkage

Limitations

- Lack of eGFR data (most recent papers use eGFR) as younger cohort
- Lack of data linkage studies on CKD in young people to validate the definition