

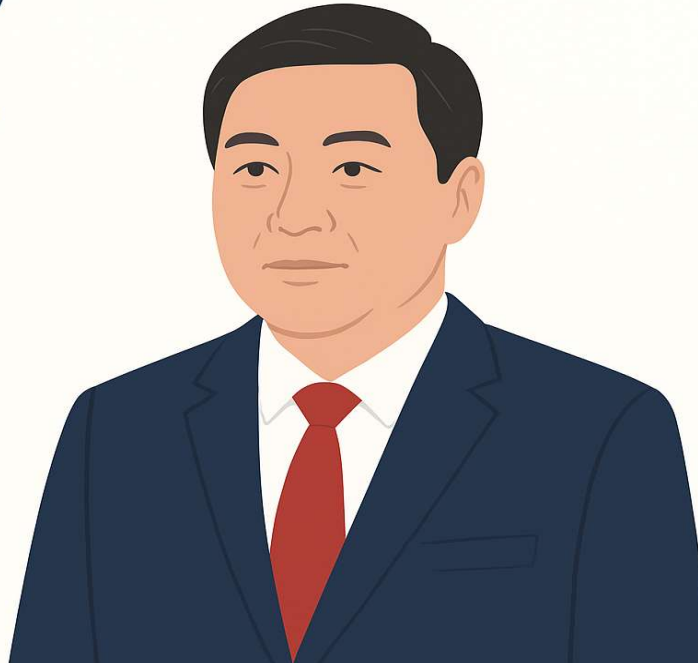
The CKD4/5 Patient. What's going on? Practical Advice

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Renal CME, GLMS
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Mr KL

Case Study

- 52-year-old Chinese New Zealander, living in East Auckland
- Owns a successful family-run business
- Two university-aged children
- Busy schedule, little time for health management



Presents for prescription renewal



- **Hypertension** (10 years)
- **Type 2 Diabetes** (15 years, suboptimal control, HbA1c ~80 mmol/mol)
- -retinopathy and required x2 laser therapy
- **BMI 32,**
- Ex-smoker
- Mild ankle swelling, BP: 160/95mmHg
- **Bloods:** eGFR 28 mL/min/1.73m² Hb 128g/L
- Urine ACR: 250 mg/mmol → CKD Stage G4A2

GFR and ACR categories and risk of adverse outcomes			ACR categories (mg/mmol), description and range		
			<3 Normal to mildly increased	3–30 Moderately increased	>30 Severely increased
			A1	A2	A3
GFR categories (ml/min/1.73 m ²), description and range	>90 Normal and high	G1	No CKD in the absence of markers of kidney damage		
	60–89 Mild reduction related to normal range for a young adult	G2			
	45–59 Mild–moderate reduction	G3a ¹			
	30–44 Moderate–severe reduction	G3b			
	15–29 Severe reduction	G4			
	<15 Kidney failure	G5			

Increasing risk →

↑ Increasing risk

¹ Consider using eGFR_{cystatinC} for people with CKD G3aA1 (see KDIGO recommendations 1.1.14 and 1.1.15)

Abbreviations: ACR, albumin:creatinine ratio; CKD, chronic kidney disease; GFR, glomerular filtration rate

Adapted with permission from Kidney Disease: Improving Global Outcomes (KDIGO) CKD Work Group (2013) KDIGO 2012 clinical practice guideline for the evaluation and management of chronic kidney disease. Kidney International (Suppl. 3): 1–150

Calculating Risk of Progression

- CVD Risk Assessment

www.nzssd.org.nz

CVD and ESRD Risk Assessment
for people with type 2 diabetes in New Zealand

INPUT

Age: 52
Duration of Diabetes: 15 years
Sex: ☒ Male ☐ Female
Smoker: Previously smoked
Systolic BP: 160 mmHg
HbA1c: 82 mmol/mol ☒ mmol/mol ☐ %
Ethnicity: Asian
Previous CVD: ☐ Yes ☒ No

Total Cholesterol: 3.2 mmol/L
HDL: 2.1 mmol/L
Urine ACR: 200 mg/mmol
Serum creatinine: 200 µmol/L
BP lowering medication: ☒ Yes ☐ No ☐ Unknown

Calculate Reset

OUTPUT

5 year CVD Risk: 28.9%
5 year MI Risk: 9.6%
5 year ESKD Risk: 55.3%

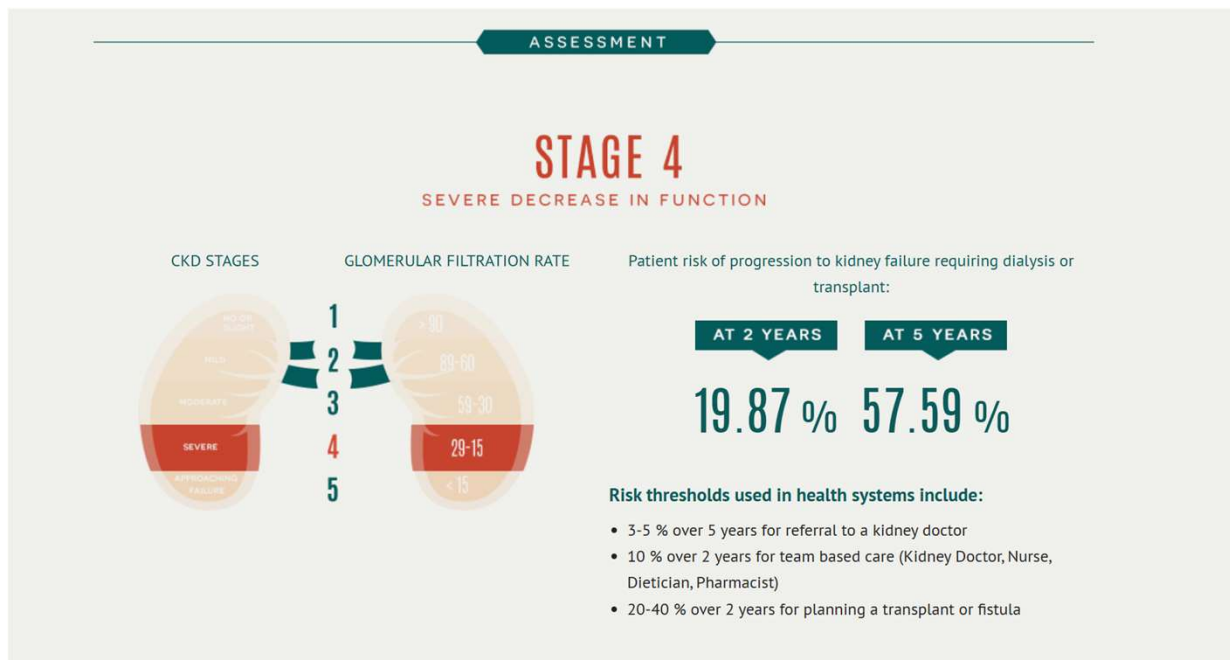
0 2.5% 5% 10% 15% 20% 30% 100%

Print

5 year ESKD risk 55.3%

The Kidney Failure Risk Equation

- <https://kidneyfailurerisk.com>



What would you do for him?

- ACE inhibitor and SGLT-2,
- Adjust diabetes meds
- Stop metformin?
- Stop any Egfr dependent regular medications
 - -> need to look up, nephrologists don't carry in their heads *
- Avoid NSAIDs
- Refer to Renal



Drugs to be mindful of when eGFR <30



DOACs

Dabigatran, Rivoroxiban 15mg od, stop at egfr 15.

Metformin – stop, risk of lactic acidosis, esp if intercurrent illness

Sulphonylureas – dose reduce,

Insulin – often need dose reduction when eGFR goes lower

Empagloflozin dose reduce 25mg-> 10mg

ACEi/ARBs – try hard to continue max dose if potassium allows

Bendrofluazide ineffective -> change to chlorthalidone (effective anti-HTN in CKD4 in CLICK Trial) or use frusemide for diuretic

Clexane – treatment dose 1mg/kg once daily (from bd)

Morphine – avoid

Colchicine – dose reduce

Gabapentin- dose reduce

Digoxin – dose reduce, check levels

****In practice, always check NZF when prescribing for eGFR <30.****

DCKD Treatment Targets

Maximum RAS Blockade possible

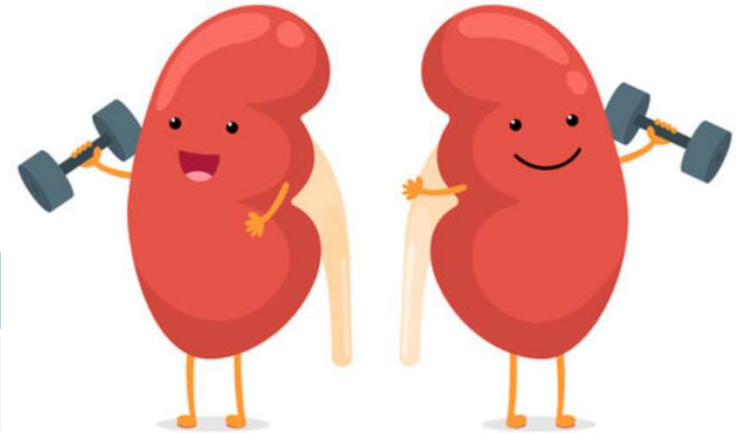
SGLT2 Inhibitor

Any other meds to achieve **BP 130/80**

Hba1c <55

Statin to keep Tot. **Chol <5**

Finerenone



When to refer to local renal service?

Request non-acute nephrology assessment if:

- eGFR declining by more than 10 within previous 12 months.
- eGFR less than 30 in a patient without diabetes.
- *eGFR less than 45 in a patient with diabetes and persistent significant albuminuria.*
- uACR >70
- uACR > 30 and haematuria
- **Diagnostic or management uncertainty**
- More than 30% increase in creatinine after starting ACE inhibitor or ARB therapy, or on dose change

What is the purpose of the referral?

- To exclude a reversible cause,
- To manage sequelae of kidneys that don't work properly,
- To discuss whether RRT would be suitable or if Conservative care is a better choice
- To prepare for dialysis and transplant work up

What to include on referral to local renal service

- What is their functional status
- Your opinion about whether they would be a dialysis candidate,
- Consider using a risk calculator [The Kidney Failure Risk Equation](#)
- **Hydration status**
- **MSU**, uACR, Blood Pressure
- Information about: prostate symptoms, systemic symptoms (weight loss, arthralgia, rash etc), excessive NSAIDs

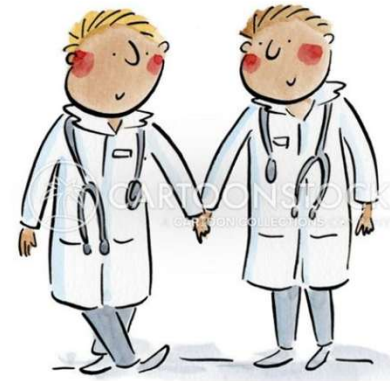
Case Study update – 18 months later

- Recurrent admissions for fluid overload
- eGFR dropped to 12 mL/min/1.73m²
- Poor diabetes control, anaemia, bone disease
- **Symptoms:**
- Nausea, severe tiredness, pruritus
- Struggles to keep up with work



GP Shared Cares in CKD4/5

- Reinforce dietary messages– Fluid, salt, potassium restriction
- Adjust Binocrit dose
- Psycho-social support
- Immunisations *
- Help with Blood pressure medication adjustment
- Help with diuretic and fluid balance management
- Ongoing Management of CV risk (lipid control, smoking cessation support, lifestyle eg, green prescription)



Immunisation for adults pre-dialysis, on dialysis or pre-/post-kidney transplantation



Vaccine	Additional notes	Recommended schedule	Pre-dialysis	On dialysis	Pre-transplant	Post-transplant
<i>Haemophilus influenzae</i> type b Hib (Act-HIB/Hiberix)		Administer one dose	Recommended NOT FUNDED	FUNDED	FUNDED	FUNDED
Hepatitis A (Havrix)	Give on the advice of renal specialist or transplant team	Administer two doses at least 6 months apart	NOT FUNDED	NOT FUNDED	FUNDED	FUNDED
Hepatitis B (Engerix-B)	Give on the advice of renal specialist or transplant team Check serology 4 weeks after 3rd dose: if non-immune, seek advice from renal specialist or transplant team	If pre-dialysis (20mcg) - Administer one 20mcg dose at each visit at 0, 1, 6 month intervals If on dialysis or pre-/post-transplant (40mcg) - Coadminister two doses of 20mcg/mL (i.e. 40 mcg) at each visit at 0, 1, 6 month intervals If accelerated schedule requested by specialist or transplant team pre-transplant e.g. active on deceased donor list(40mcg) - Coadminister two doses of 20mcg/mL (i.e. 40mcg) at each visit - Administer three doses at 0, 1, 2 months	Recommended NOT FUNDED	FUNDED	FUNDED	FUNDED
Herpes zoster Recombinant ZV (Shingrix)	Recommended for adults from the age of 18 years.	Administer two doses, at least 2–6 months apart	FUNDED - Aged 65 years - Aged 18 years+ for those with end-stage kidney disease (CKD 4 or 5) or pre- or post- solid organ transplant			

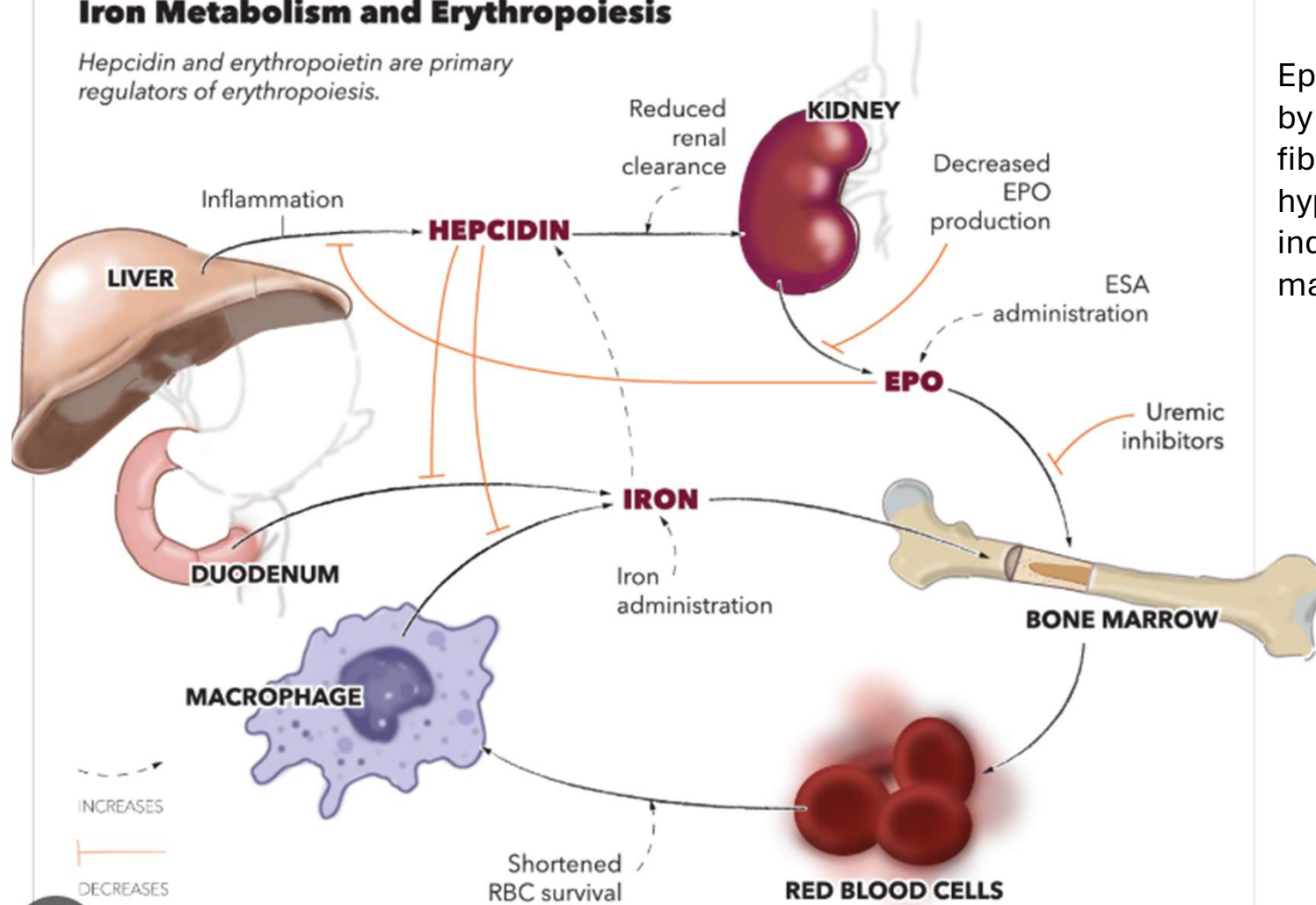
Vaccine	Additional notes	Recommended schedule	Pre-dialysis	On dialysis	Pre-transplant	Post-transplant
Human papillomavirus HPV (Gardasil 9)	Recommended for males and females 18–45 years of age inclusively	If pre-dialysis, on dialysis, or pre-/post-transplant: - Administer three doses at 0, 2, and 6 month intervals If accelerated schedule requested by specialist or transplant team pre-transplant (funded <27 years), e.g. active on deceased donor list: - Administer three doses at 0, 1, and 2 month intervals	FUNDED Up to 27 years of age (standard or accelerated schedule)			
			Recommended NOT funded 27–45 years			
Influenza	Annually, during the funded Influenza Immunisation Programme Two doses are recommended in the first year post-transplant but only the first dose is funded Influenza vaccination is recommended but not funded for household and other close contacts of pre-dialysis, dialysis, pre- and post-transplant patients	If pre-dialysis, on dialysis, or pre-transplant: - Administer one dose annually If post-transplant: - Wait until 3 months post-transplant unless at high-risk of infection: - If at high risk of infection, e.g. during influenza epidemic, wait until 1 month post-transplant - Administer one funded dose annually - In the first-year post-transplant a second purchased influenza vaccination can be administered 4 weeks later to maximise the person's immune response to the vaccine - In subsequent years only one dose is required annually	FUNDED	FUNDED	FUNDED	FUNDED
Meningococcal A,C,W,Y (MenQuadfi)	Prescription required for second primary dose	- Administer two doses at least 8 weeks apart - Schedule a precall for a booster dose every 5 years	NOT FUNDED unless on immunosuppressive therapy for longer than 28 days	NOT FUNDED unless on immunosuppressive therapy for longer than 28 days	FUNDED	FUNDED
Meningococcal B 4CMenB (Bexsero)	Can be co-administered with any other vaccine	- Administer two doses 8 weeks apart - Schedule a precall for a booster dose every 5 years +	NOT FUNDED unless on immunosuppressive therapy for longer than 28 days	NOT FUNDED unless on immunosuppressive therapy for longer than 28 days	FUNDED	FUNDED
Pneumococcal PCV13 (Prevenar 13)	If Pneumovax23 has been administered before Prevenar 13, wait one year to give Prevenar 13	Administer one dose	Recommended NOT FUNDED	FUNDED	FUNDED	FUNDED

Vaccine	Additional notes	Recommended schedule	Pre-dialysis	On dialysis	Pre-transplant	Post-transplant
Pneumococcal 23PPV (Pneumovax 23)	Administer Pneumovax23 a minimum of 8 weeks after Prevenar 13	If aged 18 years to under 60 years: - Administer one dose - Schedule a precall for the second dose in 5 years - Schedule a precall for the third/final dose 5 years after second dose or at age 65 years, whichever is later If aged 60 years or older: - Administer one dose - Schedule a precall for second/final dose in 5 years	Recommended NOT FUNDED	FUNDED	FUNDED	FUNDED
Polio IPV (Ipol)	Check immunisation history for a primary course of three polio containing vaccines	If unsure of polio immunisation history Administer three doses with a minimum of 4 weeks between each dose	FUNDED	FUNDED	FUNDED	FUNDED
SARS-CoV-2 (COVID-19)	A third primary dose of mRNA-CV is recommended for severely immunocompromised individuals Third dose should be given 8 weeks after second dose See Immunisation Handbook in relation to timing for current/planned immunosuppressive therapies	If pre-dialysis, on dialysis, or pre-transplant: - Administer vaccine doses following the recommended schedule for the available COVID-19 vaccine If post-transplant: - Wait until 3 months post-transplant, unless earlier administration indicated by specialist - Administer vaccine doses following the recommended schedule for the available COVID-19 vaccine	FUNDED	FUNDED	FUNDED	FUNDED
Tetanus/diphtheria/pertussis Tdap (Boostrix)	Check immunisation history for a primary course of three tetanus/diphtheria containing vaccines	If unsure of tetanus/diphtheria immunisation history - Administer three doses with a minimum of 4 weeks between each dose If has a confident recollection of completed tetanus/diphtheria immunisation - Administer one Tdap at age 45 years if less than four documented tetanus containing vaccine doses - Administer one Tdap at age 65 years	FUNDED	FUNDED	FUNDED	FUNDED

Vaccine	Additional notes	Recommended schedule	Pre-dialysis	On dialysis	Pre-transplant	Post-transplant
Measles/mumps/rubella MMR (Priorix)	Individuals born in 1969 or later who do not have two documented doses of MMR vaccine, or on the Advice of renal specialist or transplant team	If less than two documented doses • Complete a documented course of two MMR doses • Administer up to two doses at least 4 weeks apart^{a,b,c} • Doses as advised by renal specialist or transplant team	FUNDED	FUNDED	FUNDED for individuals who meet the eligibility criteria CONTRAINDICATED from 4 weeks pretransplant	CONTRAINDICATED
Varicella (chickenpox) VV (Varivax)	Give on the advice of renal specialist or transplant team	• Administer two doses at least 4 weeks apart^{a,b,c,d}	Recommended NOT FUNDED	Recommended NOT FUNDED	FUNDED for individuals who meet the eligibility criteria CONTRAINDICATED from 4 weeks pretransplant	CONTRAINDICATED

Iron Metabolism and Erythropoiesis

Hepcidin and erythropoietin are primary regulators of erythropoiesis.



Epo is produced by renal interstitial fibroblasts, in a hypoxia inducible manner