CKD and CDM FYI

Dr. Maggie Watt

Mr. H. E.

- 72 year old retired truck driver
- New patient in April 2002
- PMHx
 - MI 1983
 - Pituitary Tumour 1983 resected and 6/12 XRT
 - Panhypopituitarism (on Cortisone, Levothyroxine & Testosterone replacement)
 - Renal insufficiency following TUPR
 - bilateral ureteral obstruction Oct 2001
 - Thickened bladder wall and outlet obstruction
 - Creat 150 (no GFR reported yet)

Mr. H.E. (cont'd)

- 55 pack year smoker multiple attempts to quit
- Past alcoholic (quit 1983)
- HTN
- Hypercholesterolemia
- Type 2 DM (dx May 2004)
- Obesity (BMI 39.2)

Mr. H.E. cont'd

- May 2003 BP 180/110
 - Start Ramipril 2.5 mg daily, titrated to 10 mg over 2/12
 - Cough on ACEI change to losartan
- Dec 2003 BP 150/50
 - Add HCTZ
- April 2004
 - Creat 162, GFR 39 (Stable)
 - Enroll in PROMIS (Kidney Care Initiative)

Mr. H.E. cont'd

- May 2004...my chart notes change
 - Review bloodwork (FBS 12.7 = Type 2 DM)
 - "Stage 3 CKD" (GFR 41)
 - Hyper PTH (secondary)
 - Urine ACR elevated (2.67) (normal < 2 males)
 - Plan Renal U/S, Refer Nephrology
 - New goal for Lipids in view of DM 2 and CKD
 - LDL < 2.5 and TC/HDL < 4

Further Investigations

- Renal ultrasound June 2004
 - Bilateral mild symmetric cortical thinning
 - Left kidney 11.4 cm, right kidney 9.2 cm
 - No hydronephrosis
 - Bladder normal

And then he sees the nephrologist

- Nephrology Consult October 2004
 - 3 page consult
 - CKD moderate in severity
 - Small vessel renovascular disease
 - Possibly component of macrovascular dz (asymmetric kidney size on u/s)
 - Twice yearly ACR and renal function
 - Follow up 1 year

- Nephro Follow up November 2005
 - Stable moderate impairment in kidney function
 - Query right renal artery stenosis
- Nuclear renal scan with furosemide (Dec 2005)
 - "asymmetry of kidney function raises possibility of right renal artery stenosis"
- Feb 2006 acute decline in renal fxn GFR 16
 - Book MRA, possible dialysis, D/C antihypertensives, ASA
 - Renal MRA March 2006
 - Severe stenosis at origin of right renal artery
 - Nov. 2006 Angioplasty and Stent placement in Right Renal Artery
 - 70% stenosis
 - Renal function unchanged but felt almost instantly better

Ongoing Management Mr. H.E.

- Q 3/12 Diabetes Check, CKD Check
 - HTN, Sugars, Renal Fxn, Lipids, Self Care, etc.
 - Motivated re: self care
- Stable renal function
- TKR May 2008
- Died 2 days postop of massive UGI Bleed

BC CKD GUIDELINES – Sept 2008

- Recommendation 1 Identify populations at high risk:
 - Diabetes
 - HTN +/- Cardiovascular Disease
 - Family history of kidney disease
 - High risk ethnicity (First Nations, S. Asian, African Descent, Pacific Islanders)
 - \blacksquare (age >60)

BC CKD Guidelines

 Recommendation 2 – Screen Populations at Increased Risk (q1-2 years)

- Serum creatinine and eGFR
- Urine ACR
- Urinalysis macro and micro
 - (to detect protein, WBC's, RBC's, cellular casts)

eGFR

- Computed from serum creatinine
- Automatically reported by most labs in BC
- Persistent (>3mos) eGFR < 60 mL/min indicates substantial reduction in kidney function
- Patients with eGFR 60-100 with normal u/a and ultrasound do NOT have kidney disease

eGFR

- Accuracy of eGFR:
 - **Age** > 75 years old may **underestimate** true kidney function
 - eGFR 45-60 may reflect normal variation in absence of other conditions
 - Caution recommended with meds, dye and risk of acute kidney injury with severe illnesses
 - Age >85 calculation more problematic
 - Risk of progression not known
 - Conservative approach recommended

eGFR

- eGFR calculation also unreliable in:
 - Very large / very small body habitus
 - Specific diets (very high/low protein)
 - Meds that interfere with creatinine excretion
 - Trimethoprim / Sulfamethoxazole
 - Ciprofloxacin
 - Fenofibrate

Urinalysis (Random)

- Significant abnormalities include:
 - Persistent RBC's or WBC's in absence of instrumentation
 - Cellular casts

ACR

- Albumin / Creatinine Ratio
 - Random urine test no need for 24 hour collection
 - Abnormal (CDA Standard)
 - Men > 2mg/mmol
 - Women >2.8 mg/mmol
 - on 2 out of 3 serial tests between 1 week and 2 months apart (i.e. persistent)
 - Indicates micro-vascular disease +/- glomerular disease

PROTEINURIA - Definitions

MICROALBUMINURIA

- 24 hour urinary albumin excretion 30 300 mg
- Urine ACR
 - < 2.0 mg/mmol (M)
 - < 2.8 mg/mmol (F)
 - Sustained (ie. 2/3 samples)
- PROTEINURIA ('overt')
 - 24 hour urine protein excretion > 150 mg/day
 - Transient, orthostatic, or persistent
- NEPHROTIC RANGE PROTEINURIA
 - \sim > 3 grams/day
 - Typically associated with glomerular disease

Recommendation 3 – Evaluate patients with sustained impairments

- \blacksquare eGFR < 60 mL/min for > 3mos
- Evidence of **Kidney Damage**
 - "Pathologic abnormalities or markers of damage, including abnormalities in blood or urine tests, or imaging studies"

Recommendation 3 – Evaluate patients with sustained impairments

- Determine Stage of CKD based upon:
 - eGFR
 - Urinalysis
 - ACR

See guidelines for National Kidney Foundation
 (US) Staging system

TIPS ON EARLY DIAGNOSIS OF KIDNEY DISEASE

Who's at risk?

Patients with:

- diabetes:
- hypertension (with or without cardiovascular disease);
- a family history of kidney disease; and
- belonging to specific high-risk ethnic groups (First
- Nations, Asian, South Asian, Hispanic, African American and Pacific Islanders)

How can I confirm my diagnosis?

- Measure blood pressure (hypertension is common).
- Measure serum creatinine levels and estimated GFR.
- Do a urinalysis to detect protein, white or red cells.
- Measure electrolytes.
- Check hemoglobin levels.

Produced by the BC Provincial Renal Agency, the Kidney Foundation of Canada (BC Branch) and the Ministry of Health Services. For more information, contact the BCPRA: 604-806-8845, bcpra@cheos.ubc.ca or www.bcrenalagency.ca.



B C R e n a l A g e n c y
An agency of the Provincial Health Services Authority

STAGES OF REDUCED KIDNEY FUNCTION

	/	
Stage	Description	eGFR (mL/min)
1	Kidney damage with normal or ① GFR (but urine test abnormalities)	≥ 90
2	Kidney damage with mild ${\mathbb J}$ GFR	60-89
3	Moderate	30-59
4	Severe GFR	15-29
5	Kidney failure	<15 or dialysis

The CKD Clinical Practice Guideline provides recommendations for identification, evaluation and management of patients with chronic kidney disease. Copies are available through www.healthservices.gov.bc.ca/cdm or by calling 250-952-1347.

Symptoms of CKD

- Foamy / bloody urine Nausea
- Headaches
- Nocturia
- Edema (eyes, ankles)Polyuria
- Fatigue

- Restless legs
- Anorexia
- Pruritus

Complications of Reduced GFR

- Anemia
- Hypertension
- Decreased Calcium absorption
- Dyslipidemia
- Heart Failure
- Volume Overload

- Hyperkalemia
- Hyperparathyroidism
- Hyperphosphatemia
- LVH
- Metabolic Acidosis
- Malnutrition (late)

Recommendation 4 Determine Cause of CKD

- Impared kidney function often multifactorial
- Renal ultrasound
 - Polycystic Kidney Disease
 - Stones
 - Cancer
 - Obstruction
 - Discrepant size Kidneys (? Renal artery stenosis)

Causes of CKD

- Diabetes (Type 1 and Type 2)*
- Hypertension*
- Other vascular diseases
 - Large vessel disease, microangiopathy
- Glomerular diseases:
 - Autoimmune, systemic infection, drugs, neoplasia
- Tubulointerstitial Disiases
 - UTI, stones, obstruction, drug toxicity
- Polycystic Kidney Disease

(*account for 2/3 of CKD and ESRD)

Figure 1. Evaluation and Management of Suspected Kidney Disease

Identify & screen high-risk populations (Recommendations 1 & 2) · Ascertain the risk factors Systems review and physical exam Laboratory tests: · serum creatinine (for estimated GFR) random urine – macro/micro urinalysis & ACR (microalbumin) Repeat tests to confirm abnormal results **TESTS NORMAL TESTS ABNORMAL** Determine stage of CKD Monitor eGFR, urinalysis, based on eGFR, urinalysis, ACR, BP every two years. ACR. Yearly monitoring indicated in selected patient populations. Determine cause of kidney disease. Ongoing follow-up and referral as per Table 1. Meet care objectives as per Recommendation 5.

WHEN TO REFER

- Sustained decline in GFR < 30mL/min</p>
- Acute renal failure
- Subacute decline in kidney function
 - >10 mL/min annually
- Sustained proteinuria > 1gram/24 hrs
- Active urine sediment
 - Cellular casts, sustained hematuria &/or proteinuria

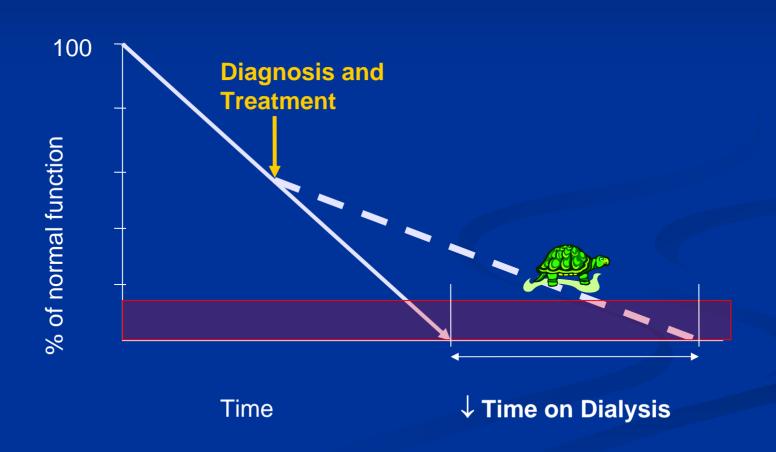
DEFINITIONS / CLARIFICATION

- Certain kidney diseases often require <u>specific</u> management:
 - Glomerulonephritis
 - Obstructive uropathy
 - Acute interstitial nephritis
 - Renal artery stenosis
- Non-disease specific therapies aimed at slowing progressive nephropathy, regardless of:
 - Disease etiology
 - Stage of CKD

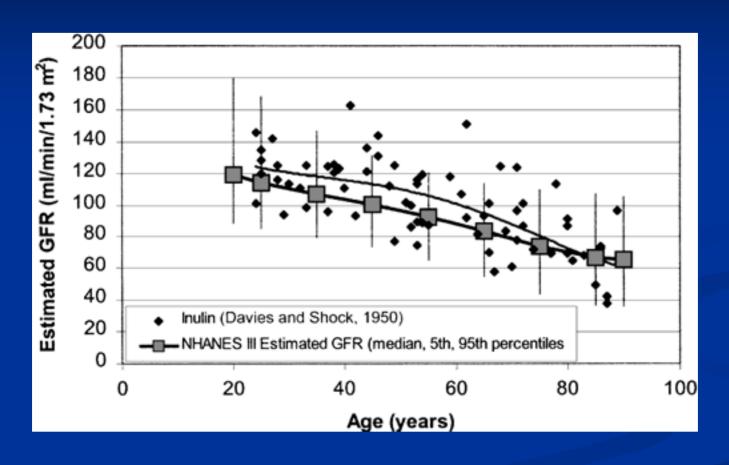
A BRIEF REVIEW – CKD Treatment

- Consider reversible factors
- Avoid nephrotoxins
 - NSAIDs, contrast, aminoglycosides
- Slow CKD progression:
 - BP <130/80 (or 125/75 if proteinuria >1 gram/day)
 - Consider ACEi or ARB therapy
 - Control BG in diabetics (HgA1c <7%)
 - +/- dyslipidemia therapy
 - +/- dietary protein restriction
- Follow CHEP, CDA, CCS guidelines for secondary cardiovascular prevention

END-STAGE KIDNEY DISEASE CAN BE PREVENTED (OR SLOWED)



GFR DECLINES WITH AGE



Normal decline 1 % per year

IMPLICATIONS

- Patients need information on CVD / mortality risk not just progressive nephropathy
- Patients with progressive disease need info on preparation for RRT
- Older patients may benefit less than younger from intensive therapeutic efforts
- Male patients may require more aggressive evaluation, treatment, follow-up, and earlier referral
- More predictors of progressive CKD required

PROTEINURIA - SUMMARY

- Proteinuria is significant when
 - Sustained (>3mos)
 - High-grade
 - Always warrants nephrology referral

- Treatment
 - Lower BP (ACEi or ARB first line)!
 - Treat diabetes to target
 - Attend to other CV risk factors

ANEMIA - SUMMARY

- Increasing prevalence with reduced kidney function
- Transferrin Saturation better gauge of iron stores than Ferritin at low GFR
- Prescribe erythropoietin therapy (Nephrology)
 - After other causes of anemia ruled out
 - After iron stores replete
- Monitor response to therapy monthly
 - Therapy usually well tolerated but watch for HTN with rapid increases in Hgb
- Maintain target hemoglobin 110-130 increased mortality outside that range

Bone Mineral Metabolism Objectives for Stage 3 CKD

Disease State:

- Hyperphosphatemia
- Hypocalcemia
- Decreased Calcitriol (activated Vit D)

■ all increase PTH

Treatment sequence (Not a medical emergency)

- 1. Dietary Phosphate restriction (target normal PO4 level)
- 2. Calcium-based binders with meals (target normal Ca and P04 levels)
 - Start TUMS 1 tab with each meal (decrease P04 and increase Ca2+)
- 3. Alpha Calcidiol (if PTH > 7.7 pmol / L)
 - One-alpha 0.25 mg daily
- Monitor labs q 6 mos in treatment phase

Hyper PTH in CKD

- Need to target progressively higher PTH to maintain normal bone turnover as CKD progresses
- Caused by skeletal resistance to PTH

CKD Stage	GFR	Target PTH*
3	30-60	3.8-7.7
4	15-29	7.7 - 12
5	< 15 (dialysis)	16.5 - 33

^{*} opinion based levels

SUMMARY – MINERAL METABOLISM

- Measure Ca / PO4 / PTH (and albumin) at least yearly
- Restrict dietary PO4 intake
- When hyperphosphatemia occurs:
 - Reinforce dietary PO4 restriction
 - start PO4 binders (typically Ca-based)
- Maintain normal serum Ca levels
- Rx Vitamin D if hypocalcemic or if PTH above target



Strategic Initiatives, Medical and Pharmaceutical Services





BRITISH COLUMBIA MEDICAL

CHRONIC KIDNEY DISEASE COLLABORATIVE FLOW SHEET/ ENCOUNTER FORM

	ATORY FIELDS					♦ PHN#	OR OTHER	UNIQUE PATIENT ID)	♦ DA	TE OF VISIT (DD-MMM-YYYY)
♦ PATIENT	NAME									,
♦ BIRTHD	ATE (DD-MMM-YYYY)	♦ GENDER MALE	FEMALE	PHONE (INCLUDE	AREA COL	DE)	CI	HART NUMBER		
PRACTICE	TEAM ID		1	PROVIDER ID (MS	P PRACTIT	TIONER NU	MBER / NAI	ME)		
CO-MOR	BID CONDITIONS									
ALCOHOL OVERUSE COR. ART. DIE ARTHRITIS CARDIOMYOL ASTHMA COPD ATRIAL FIBRILLATION HYPERTENSI			DPATHY LIVER OBESITY			SMOKING SUBSTANCE ABUSE		☐ DIABETES ☐ KIDNEY ☐ DEPRESSION ☐ CONGESTIVE HEART FAILURE		
	OSIS: TYPE OF KIDNEY DIS		YSTIC KD	OTHER					DATE	E OF DIAGNOSIS (DD-MMM-YYYY)
PATIEN	T ENCOUNTERS, DIA	GNOSTIC/CL	INICAL D	ATA, BY DAT	ΓE					√ = RECALL
REVIEW						recen	T DATA			NEW DATA
≿ 5	BP <130/80 - Every V	'isit						ì		ENTER VALUE
PHYSIOLOGY	Weight BMI (stable) 18.5-24.9 - Every Visi	it .					-			LBS FTIN CM - or -
z	sCr & eGFR (stable)				-			·		ENTER VALUE
KIDNEY FUNCTION	- At Least 6 Months ACR ≥50%Reduction Baseline - At Least 6			Neg						ENTER VALUE OR POS NEG
SUGAR	A1C <0.7 - Every 3 M								•	ENTER VALUE
LIPID S PROFILE	LDL <2.5 - At Least A	nnually								ENTER VALUE
	Ratio <4.0 - At Least	Annually								ENTER VALUE
MIA.	Hgb >120 - At Least A	Annually							,	ENTER VALUE
ANEMIA	TSAT >20% - At Leas	st Annually								ENTER VALUE
SM	Calcium >2.2 - At Lea	ast Annually								
MINERAL	Phosphorus <1.4 - At Least Annually iPTH & Albumin in No	rmal Panga	•							ENTER VALUE
Σ	- At Least Annually	offilial Range								REVIEWED
REMINDERS	More Kidney-Specific	Education?								REVIEWED
	Regular Blood Work Schedule Established	d?								COMPLETED
	Referred to a Nephro	ology Team?								REVIEWED
	Regular Visits Establ	ished?								COMPLETED
IMUNIZATION	Date of Last Influenz	a Vaccine			COLUMN					DATE (DD-MMM-YYYY)
	Date of Last Pneumo	onia Vaccine	,		N. Carlos					DATE (DD-MMM-YYYY)
	Date of Last Hepatitis	s B Vaccine								DATE (DD-MMM-YYYY)
				COMM	IENTS				-	
HLTH 469	1 2005/04/04									

How has Toolkit/CDM been useful

- Learn and follow guidelines
- Planned follow-up
 - need to develop recall system
- CDM visits are MY agenda
- Office visits more organized / less harried
- "Shared care" with nephrologist
- ease of billing