

IN THE NAME OF GOD

Assessmentof GFR & ESRD Risk in living kidney Donor C and idates



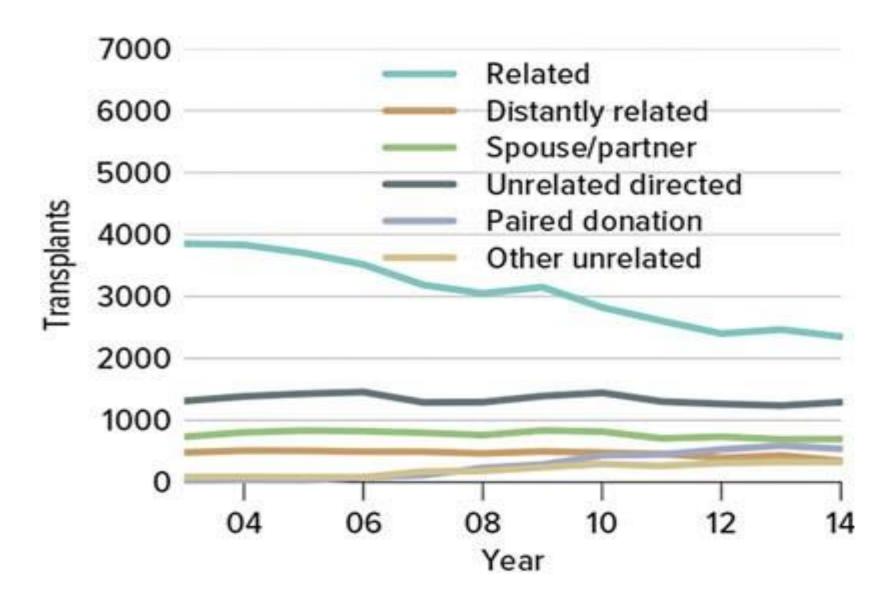
Dr Sahar Vahdat assistant Professor of Nephrology Khorshid Kidney Center IUMS

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Introduction

- ☐ As per UNOS (United Network for Organ Sharing), there are more than 100,000 patients awaiting renal transplantation
- ☐ living donation rates have declined progressively for more than a decade
- ☐ This decrease has largely been driven by a **reduction** in the number of **living related kidney donations**, from 4340 in 2004 to 2693 as per 2014
- ☐ the Organ Procurement and Transplant Network (OPTN) has defined policies which outline the minimum general and kidney-specific requirements for suitability as aliving kidney donor



PREDONATION KIDNEY FUNCTION

1. Evaluation

2. Selection

3. Counseling

Question 1

 Which of the following methods is recommended for the initial evaluation of kidney function in a living donor?

- a. eGFRcr CKD-EPI
- b. eGFR cr-cys
- c. mGFR
- d. mCrCl

Answer

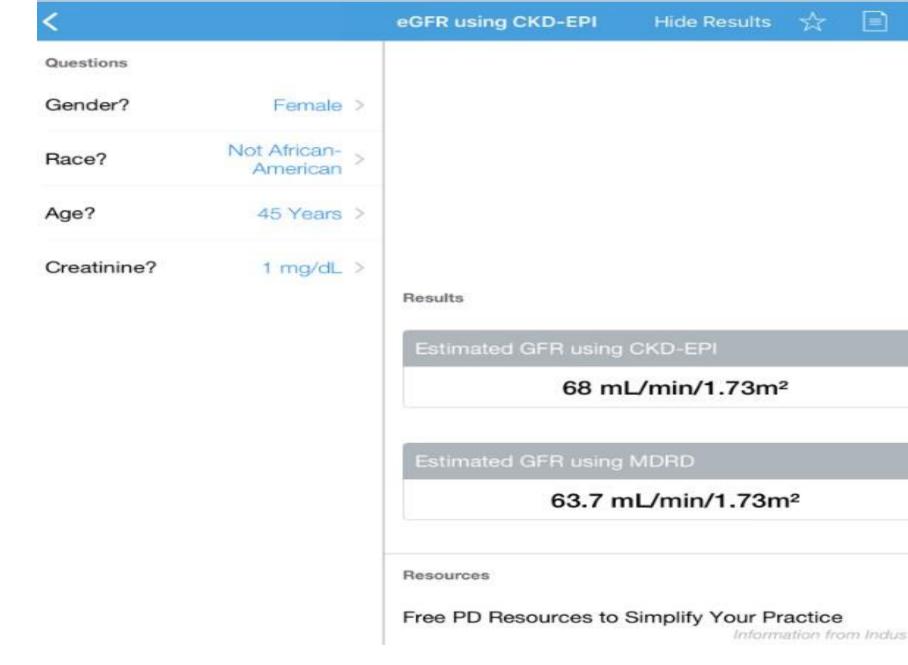
a.eGFRcr CKD-EPI

b.eGFRcr- cys

c. mGFR

d.mCrCl





Evaluation

- ☐ Donor kidney function should be expressed as **GFR** & **not as serum creatinine** concentration
- □ Donor GFR should be expressed in mL/min per 1.73 m2 rather than mL/min
- ☐ Donor GFR should be estimated from **serum creatinine** (**eGFRcr**) for **initial** assessment

Evaluation (cont...)

Donor GFR should be confirmed_using one or more of the following:

- Measured GFR (mGFR) using an exogenous filtration marker: preferably urinary or plasma clearance of inulin, iothalamate, 51Cr-EDTA, iohexol or urinary clearance of 99mTc-DTPA
- Measured Cr clearance (mCrCl): The most commonly used measure of evaluating GFR in clinical practice is based on a 24-h creatinine clearance
- Estimated GFR from the combination of serum Cr & cystatin C (eGFRcr-cys)
- Repeat eGFRcr
- Both eGFRcr and eGFRcys are imprecise at high levels of GFR, so confirmatory testing is recommended for all donor candidates

Evaluation (cont...)

☐ If there are parenchymal, vascular or urological abnormalities or asymmetry of kidney size:

(As specified by KDIGO guidelines, considered as a difference in kidney size >10% e.g., a difference in kidney length >1.2 cm or kidney volume >30 ml on renal imaging)

In these situations, most centers would prefer to transplant the kidney with lesser function and leave the donor with the kidney with greater function after all technical considerations

Single kidney GFR should be assessed using radionuclides or contrast agents that are excreted by glomerular filtration (eg, 99mTc-DTPA)

Confirmatory test

mGFR using an exogenous filtration marker is the most accurate confirmatory test

- mGFR is not available in all centers, so alternatives are acceptable mClcr is not as accurate as mGFR
- ➤ mClcr overestimates mGFR because of Cr secretion, with the magnitude of overestimation exceeding 15% at normal GFR, & is prone to error because of inaccurate urine collections
- > eGFRcr-cys is generally recommended over eGFRcr or eGFRcys
- > Repeat eGFRcr can be used if no other confirmatory tests are available

Selection

Classification of GFR category

Not acceptable Intermediate Acceptable for donation range for donation

<u>></u>90

Figure 3.

< 60

KDIGO classification of GFR categories and use in decision-making for donor candidates. Colors are blended together to signify that the threshold for decision-making is imprecise.

60-89

PREDONATION ALBUMINURIA Evaluation

- ☐ Donor proteinuria should be measured **as albuminuria**, not total urine protein
- □ Initial evaluation of donor albuminuria screening should be performed using urine albumin-to-creatinine ratio (ACR) in a random urine specimen(<0.2mg alb/mg creatinine)

Donor albuminuria should be confirmed using:

- . Albumin excretion rate (AER, mg/day [mg/d]) in a timed urine specimen
 - . Repeat ACR if AER cannot be obtained

PREDONATION ALBUMINURIA Selection

- ☐ Urine **AER less than 30 mg/d** should be considered an acceptable level for donation
- ☐ The decision to approve donor candidates with AER 30 to 100 mg/d should be individualized based on demographic and health profile in relation to the transplant program's acceptable risk threshold
- ☐ Donor candidates with urine **AER greater than 100 mg/d** should not donate



Transplantation Proceedings

Volume 38, Issue 9, November 2006, Pages 2796-2797



Transplant Proc. 2006 Nov;38(9):2796-7.

Evaluation of proteinuria in healthy living kidney donor candidates.

Leischner MP1, Naratadam GO, Hou SH, Singh AK, Leehey DJ.

Author information

Abstract

BACKGROUND: Evaluation of living kidney donor candidates includes careful assessment for the presence or absence of kidney disease. Kidney donation has been considered to be at least relatively contraindicated if urinary total protein excretion is above the normal range. However, at the present time, there is no uniformly accepted level of urine total protein excretion that would exclude donation. Albumin excretion instead of total protein excretion as a criterion has not previously been evaluated.

MATERIALS AND METHODS: This was a prospective observational study over a 3-year period in a single tertiary care center designed to assess current selection criteria for kidney donation with respect to urine total protein and albumin excretion.

RESULTS: Twenty four percent (25 of 105) of healthy adult kidney donor candidates had elevated urinary total protein excretion rates (150 to 292 mg/24 h). Of these 105 candidates, 39 had simultaneous measurements of both urinary total protein and albumin. Although one-third (13/39) had elevated 24-hour urine total protein values, none had elevated urine albumin excretion.

CONCLUSION: Measurement of albumin, the most common single protein found in urine, appears to be helpful in the evaluation of proteinuria in donor candidates. Many healthy adult kidney donor candidates have mildly elevated total protein excretion but normal albumin excretion. We believe that such patients should not be excluded from donation.

PMID: 17112832 DOI: 10.1016/j.transproceed.2006.08.126

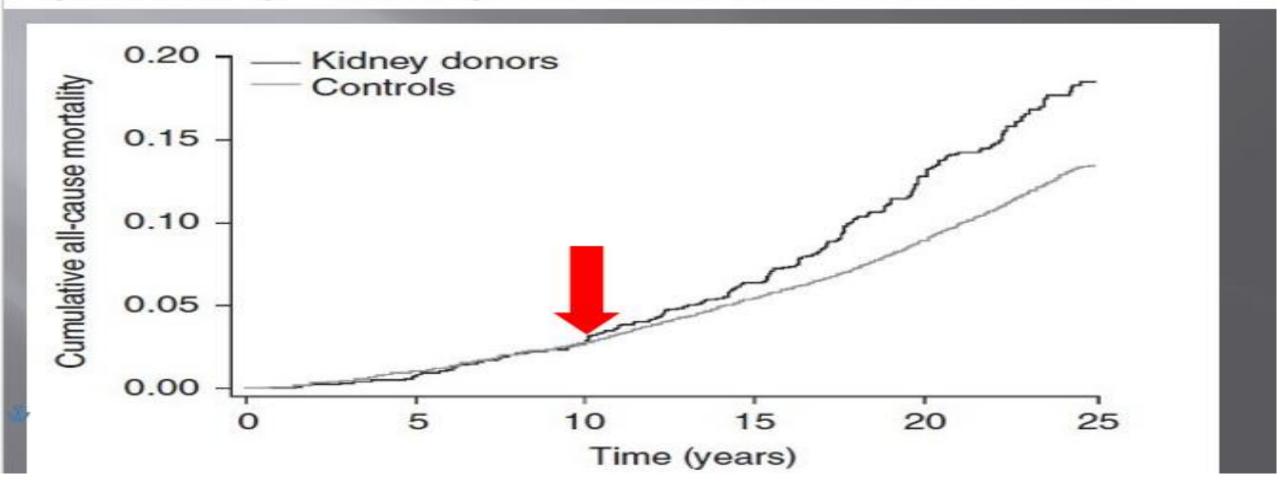
Counseling

- We suggest that donor candidates be informed that the future risk of developing kidney failure necessitating treatment with dialysis or transplantation is slightly higher because of donation
- □ average absolute risk in the 15 years following donation remains low

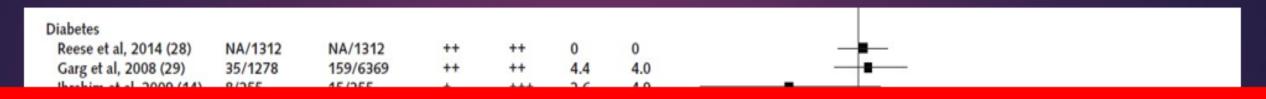


Long-term risks for kidney donors

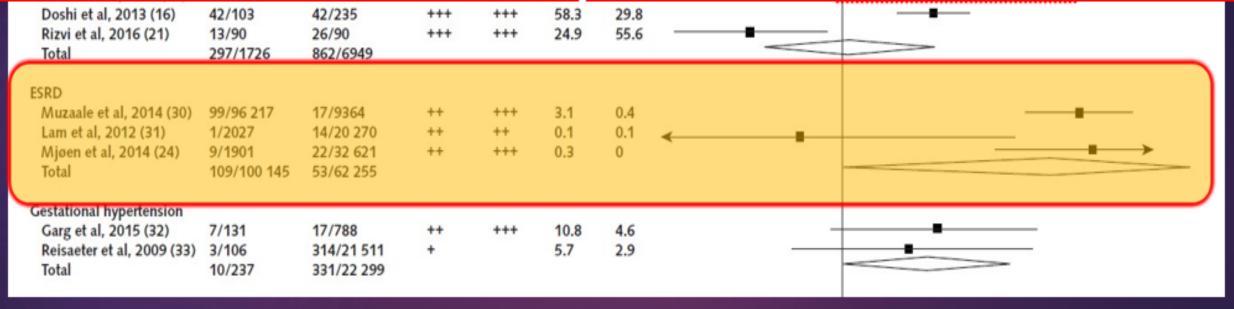
Geir Mjøen¹, Stein Hallan^{2,3}, Anders Hartmann¹, Aksel Foss¹, Karsten Midtvedt¹, Ole Øyen¹, Anna Reisæter¹, Per Pfeffer¹, Trond Jenssen¹, Torbjørn Leivestad⁴, Pål- Dag Line¹, Magnus Øvrehus², Dag Olav Dale¹, Hege Pihlstrøm¹, Ingar Holme⁵, Friedo W. Dekker⁶ and Hallvard Holdaas¹



Mid- and Long-Term Health Risks in Living Kidney Donors: A Systematic Review and Meta-analysis.



kidney donation was associated with a relative risk for ESRD of 8.83 compared with nondonors



Question 2

 Do more complicated GFR approval procedures beyond eGFR for all candidates?

a.Yes

b.No

1397/12/14

20

Answer

- Do more complicated GFR approval procedures beyond eGFR for all candidates?
- a. Yes

b. No

Assessment of GFR Range

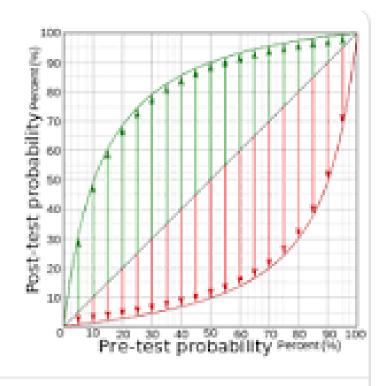
A web-based calculator has been developed to compute post-test probabilities for mGFR above or below various threshold probabilities

(http://ckdepi.org/equations/donor-candidate-gfr-calculator/)

Accessed March 1, 2017

Pre- and post-test probability





Pre-test probability and post-test probability are the probabilities of the presence of a condition before and after a diagnostic test, respectively. Post-test probability, in turn, can be positive or negative, depending on whether the test falls out as a positive test or a negative test, respectively.

Cont...

web-based tool to compute the **probability of mGFR** in living donor candidates based on demographic characteristics obtained from the National Health and Nutrition Examination Survey (NHANES) and the test performance of **eGFR** from **CKD-EPI**

This tool was subsequently validated in a French cohort of 311 living donor candidates, with demonstration of good diagnostic performance

if the post test probability that mGFR is greater than the GFR threshold for decision-making based on this tool (eg, .80 or 90 ml/min per 1.73 m²) is extremely high and if urine ACR is very low, then these tests(eGFR) could simply be repeated for confirmation without mGFR, mCrCl, or timed AER

Assessment of GFR Range (cont..)

Donor candidates with eGFR in <u>intermediate</u> ranges would require confirmatory tests with mGFR, mCrCl

Diagnostic Post Test Probability of Disease Calculation	
Enter the value of pretest probability	_
	%
Enter the value of likelihood ratio	
+ _	
Calculate	
Calculate Clear For a positive diagnostic test, the post test probability is	
	%
	%
For a positive diagnostic test, the post test probability is	%

CKD-EPI

CKD-EPI (Chronic Kidney Disease Epidemiology Collaboration)

Search

Home

says

FR CKD Prevalence

CKD Prognosis

Surrogate Endpoint

CKD-EPI

Links



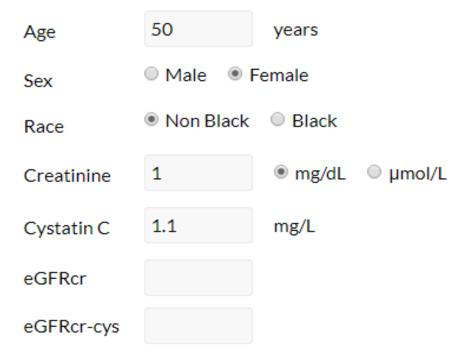
Donor Candidate GFR Calculator: Determining Probability of GFR Above or Below Certain Threshold

The data below are derived from Huang N, Foster M, Lentine K et al. Estimated GFR for living kidney donor evaluation.

Donor Candidate GFR Calculator: Determining Probability of GFR Above or Below Certain Threshold

The data below are derived from Huang N, Foster M, Lentine K et al. Estimated GFR for living kidney donor evaluation. American Journal of Transplantation [epub ahead of print Nov 23 2015].

Step 1: Describe your patient



Step 2: Determine pre-test probability

Calculate pre-test probability



Result (Pre-test probability based on age, sex and race from US population data)

Measured GFR Thresholds			
<60	<70	≥80	≥90
6	12	77	64

Percentage of NHANES participants who had eGFRcr-cys less or above the specified thresholds. If you are interested in the opposite of the threshold (eg < 80), then subtract the value given from 100.

You can use the numbers above or a different number based on knowledge of your patient's medical history. For example, if your patient has a strong family history of CKD, you may wish to alter the pretest probability. If so, write the number to be used as pre-test probability in table below.

Step 3: Calculate the post-test probability

The calculator will look up the likelihood ratio based on the eGFR calculated from the creatinine and cysatin provided above and will compute post-test probability.

Calculate post-test probability

	Measured GFR Thresholds				
	<60	<70	≥80	≥90	
Post-test probability from eGFRcr [†]	4	15	52	18	
Post-test probability from eGFRcr-cys‡	2	19	20	2	

Summary report

Based on the information supplied:

Age 50

Sex Female

Race Non Black

Creatinine 1 mg/dL

Cystatin C 1.1 mg/L

Estimated GFR from creatinine 66 ml/min per 1.73 m²

Estimated GFR from creatinine-cystatin C 66 ml/min per 1.73 m²

	Measured GFR Thresholds			
	<60 <70 ≥80 ≥9			
Pre-test probability	6	12	77	64
Post-test probability from eGFRcr†	4	15	52	18
Post-test probability from eGFRcr-cys‡	2	19	20	2

Assessment of GFR Range(cont...)

❖ A recent study suggested that eGFR may be sufficiently accurate for decision-making without the need for mGFR or mClcr in many donor candidates

In that study, 53% of recent donors in the United States had eGFR sufficiently high to provide a ≥95% post-test probability that mGFR was ≥90 ml/min per 1.73 m²

Step 3: Calculate the post-test probability

The calculator will look up the likelihood ratio based on the eGFR calculated from the creatinine and cysatin provided above and will compute post-test probability.

Calculate post-test probability

	Measured GFR Thresholde				70
	<60	<70	≥80	≥90	١
Post-test probability from eGFRcr ⁺	0	0	100	98	J

Post-test probability fro

[†]Post test probability of mGFR

the threshold (eg < 80), then sul

No need for confirmatory test if post probability is ≥95%

tes

33

Summary report

Based on the information supplied:

Age 70

Sex

Race Non Black

Creatinine 1.3 mg/dL

Cystatin C 1.1 mg/L

Estimated GFR from creatinine 42 ml/min per 1.73 m²

Estimated GFR from creatinine-cystatin C 51 ml/min per 1.73 m²

	Measured GFR Thresholds				
	<60	<70	≥80	≥90	
Pre-test probability	27	47	34	16	
Post-test probability from eGFRcr [†]	91	98	1	О	
Post-test probability from eGFRcr-cys [‡]	97	100	О	О	

Necessity for confirmatory test

Donor Candidate GFR Calculator: Determining Probability of GFR Above or Below Certain Threshold

National	Kidney	Foundation*
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The data below are derived from Huang N, Foster M, Lentine K et al. Est American Journal of Transplantation (epub ahead of print Nov 23 2015)

Step 1: Describe your patient

Age	50	years	
Sex	O Male ® F	emale	
Race	® Non Black	O Black	
Creatinine	0.7	● mg/dL	○ µmol/L
Cystatin C		mg/L	
eGFRcr			
eGFRcr-cys			

Hoton	(Commented to the control of the con	Minney Offsonia	Part	mente.	Organia Decembrican III. Franciscophical Addison
GER CAL	CULATOR				
to age, say		nd declines with	ege, The	e reations	draw function. Marro at Eddney Foundatio
Sterum Cre	arthertree :	0.7	-	macet.	C Lement/L
Serum Cvs	datin Ci		100	m/L	
distant 1		240			
Conder-		C) Harte		residen	
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CALE	*****				
Reputts					
CKD-EFT O	eathers equation validities (equation (canages (5015)	100 h 100 m 100 m		eL/min/1.72m* si_/min/1.72m* si_/min/1.72m*

Calculate post-test probability

	Measured GFR Thresholds				
	<60	< 70	≥80	≥90	
Post-test probability from eGFRcr [†]	0	1	96	89	
Post-test probability from eGFRcr-cys ^a	N/A	N/A	N/A	N/A	

*Post test probability of mGFR above or below the threshold based on eGFRcr. If you are interested in the opposite of the threshold (eg = 50), then subtract the value given from 100.

³ If you also indicated the eGFRcr-cys, the calculator will use the post-test probability from the eGFRcr as the pre-

Necessity for mGFR in spite of eGFR 101 ml/min

American Journal of Transplantation 2016; 16: 3024–3032 Wiley Periodicals Inc. Copyright 2016 The American Society of Transplantation
 and the American Society of Transplant Surgeons

doi: 10.1111/ajt.1390k

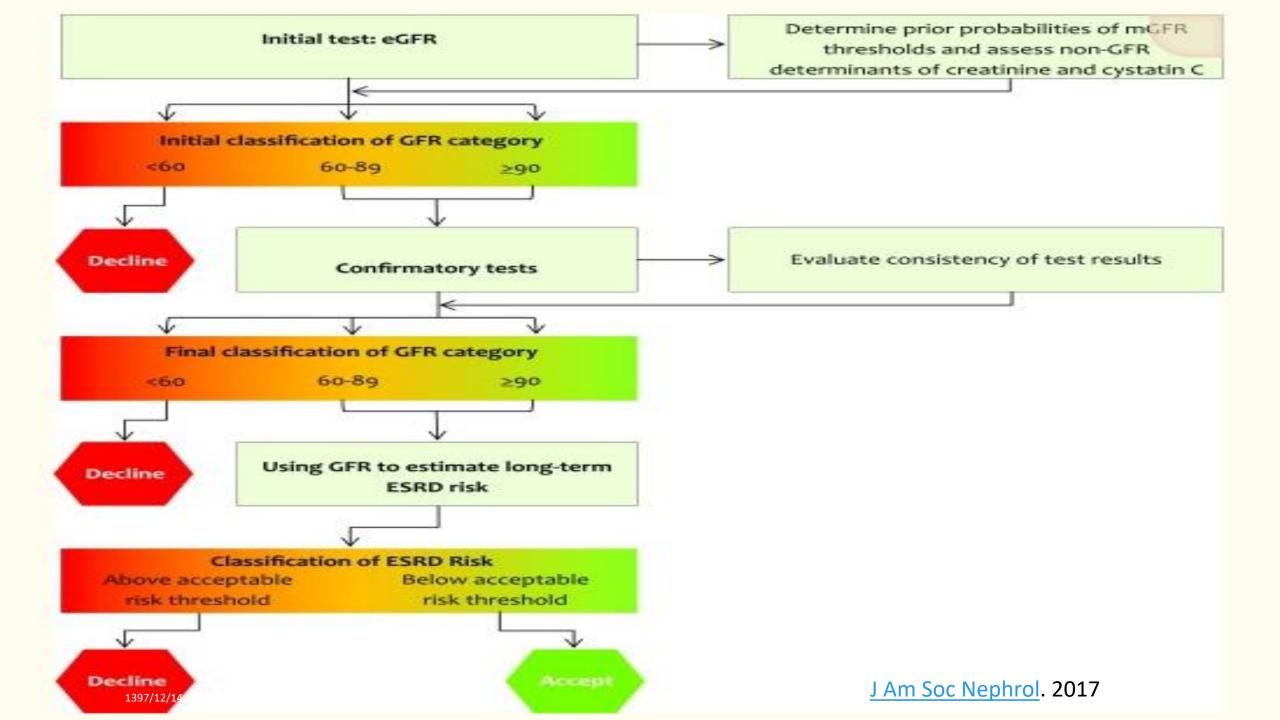
Brief Communication

Estimated or Measured GFR in Living Kidney Donors

Work-up?

Physiology Department, Paris Descartes University, and INSERM, Unit 1151, Paris, France ¹²AP-HP, Hopital Europ[^] een Georges Pompidou,!

In conclusion, we recommend calculating posttest 90 for each potential kidney donor



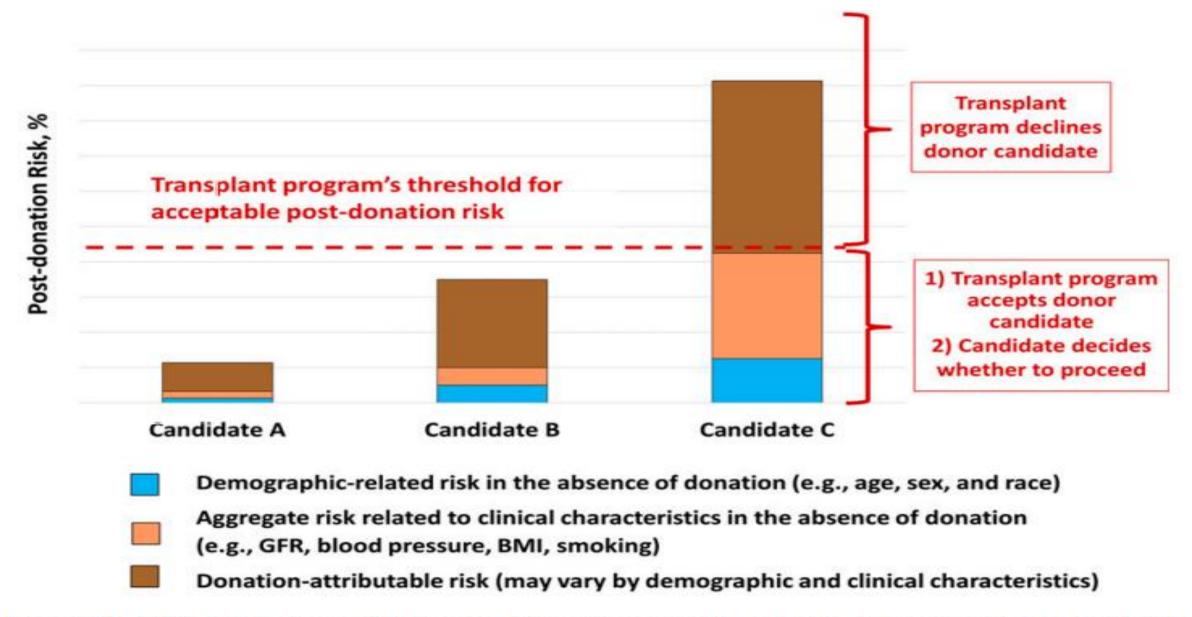
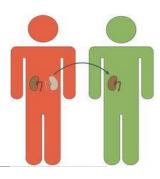


Figure 1. KDIGO proposed framework for a transplant center to accept or decline a donor

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Long-term ESRD risk



A. Long-term risk in the absence of donation can be computed from demographic and clinical characteristics, including GFR (http://www.transplantmodels.com/esrdrisk/)

B. Additional risk attributable to donation is likely to be 3.5–5.3 times higher than risk in the absence of donation depending on sex and race, but there is substantial uncertainty, especially in younger donor candidates, and we suggest caution in decision-making

J Am Soc Nephrol. 2017 Apr; 28(4): 1062–1071

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Prediction Models

This model projects 15 year and lifetime predonation ESRD risks (ie, without donation-attributable risk) in donor candidates based on age, sex, race, GFR, ACR, diabetes, smoking, blood pressure, antihypertensive drug use, and BMI

➤ The tool was developed from a meta-analysis of data from nearly 5 million healthy persons from 7 cohorts, with calibration to annual ESRD incidence in the US healthy population, and is now available online: http://www.transplantmodels.com/esrdrisk

<u>Grams ME, Sang Y, Levey AS, et al.</u> N Engl J Med 2016;374(5):411-421

Use of prediction tool to estimate ESRD risk in donor candidates

- Use the online tool (http://www.transplantmodels.com/esrdrisk) to estimate the projected lifetime risk of kidney failure in the absence of donation according to baseline demographic and health characteristics included in the online tool
- Multiply the projected predonation risk by the best available estimate for donationattributable risk to obtain the projected postdonation risk. For example, Grams et al report a relative risk of 3.5-5.3 for 15-year ESRD risk, according to sex and race
- 3. Compare the projected risk estimate to the program's postdonation threshold of acceptable risk
- 4. Exercise caution when there is concern that the individual has risk factors not captured in the online tool (eg, familial or genetic risk) and for younger candidates

KDIGO Clinical Practice Guideline on the Evaluation and Care of Living Kidney Donors.

Transplantation. 2017;101(8S Suppl 1):S1-S109

ESRD Risk Tool for Kidney Donor Candidates

Projected Incidence of End-Stage Renal Disease:

0.04%

Pre-Donation 15-Year*

0.30%

Pre-Donation Lifetime*

?

Post-Donation 15-Year**

?

Post-Donation Lifetime**

blue: < 1%, green: 1-2%, yellow: 2-3%, orange: 3-5%, red: >5%

The pre-donation risks represent projections if a person does not donate

Patient Characteristics:

Age (18-80yrs)

Gender

Race (White or Black)

eGFR (mL/min/1.73m²)

Systolic Blood Pressure (mmHg)

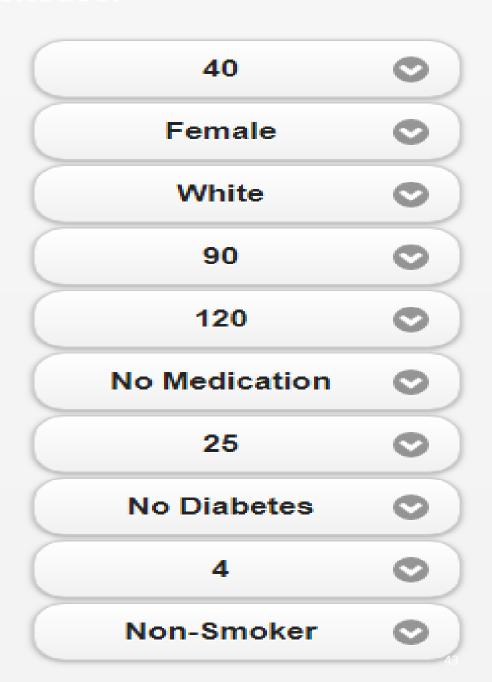
Hypertension Medication

BMI (kg/m²)

Non-Insulin Dependent Diabetes

Urine Albumin to Creatinine (mg/g)
click on units to change between mg/g and mg/mmol

Smoking History



Postdonation Risk of ESRD in Living Kidney Donors

Kidney Donor Risk of ESRD

Select your donor characteristics below. This prediction model is intended for adults who have already donated a kidney in the United States. It provides an estimated risk of developing ESRD.

Patient Characteristics:

Sex	Female	0
Race (African American or non-African American)	Non-African American	0
Age (18-80yrs)	40	0
BMI (kg/m²)	25	0
Donor is 1st degree biological relative to recipient	No	0

1207/12/1/

Question3

Do you accept the following candidate for donation?

a. yes b. No

Projected Incidence of End	i-Stage Renal Disease:	
0.16% Pre-Donation 15-Year	2.35% Pre-Donation Lifetime	,-
?	?	
Post-Donation 15-Year**	Post-Donation Lifetime	
The pre-donation risks represent projet a kidney. Details about estimating posterior print Patient Characteristics.	t-donation risk are provided t summary	
Age (18-80yrs)	30	0
Gender	Male	0
Race (White or Black)	White	0
eGFR (mL/min/1.73m²)	95	0
Systolic Blood Pressure (mmHg)	130	0
Hypertension Medication	No Medication	0
BMI (kg/m²)	30	0
Non-Insulin Dependent Diabetes	No Diabetes	0
Urine Albumin to Creatinine (mg/g) click on units to change between mg/g and mg/mmol	20	0

answer

Do you accept the following candidate for donation?

a. yes

b. No

Projected Incidence of End-S	Stage Renal Disease:	
0.16% Pre-Ponation 15-Year	2.35% Pre-Donation Lifetime	_
Pre-Donation 15-Year	Pre-Donation Elletime	
Post-Donation 15-Year**	Post-Donation Lifetime	
blue: < 1%, green: 1-2%, yellow: 2-3	3%, orange: 3-5%, red: >	5%
The pre-donation risks represent projection a kidney. Details about estimating post-donation reset print s		
Patient Characte	eristics:	
Age (18-80yrs)	30	0
Gender	Male	0
Race (White or Black)	White	0
eGFR (mL/min/1.73m²)	95	0
Systolic Blood Pressure (mmHg)	130	0
Hypertension Medication	No Medication	0
BMI (kg/m²)	30	0
Nen Inculin Dependent Diebetes	No Diabetes	0
Non-insulin Dependent Diabetes	140 Diabetes	
Urine Albumin to Creatinine (mg/g) click on units to change between mg/g and mg/mmol	20	0

Patient Characteristics:

Age (18-80yrs)

Gender

Race (White or Black)

eGFR (mL/min/1.73m²)

Systolic Blood Pressure (mmHg)

Hypertension Medication

Urine Albumin to Creatinine (mg/g)

Post-Donation Lifetime**

BMI (kg/m²)

Non-Insulin Dependent Diabetes

click on units to change between mg/g and mg/mmol

Smoking History

Male White

32

85

145

No Medication

28

No Diabetes

45

Non-Smoker

No, I do'nt accept this

blue: < 1%, green: 1-2%, yellow: 2-3%, orange: 3-5%, red: >5%

Projected Incidence of End-Stage Renal Disease:

Pre-Donation Lifetime*

1397/12/14

0.21%

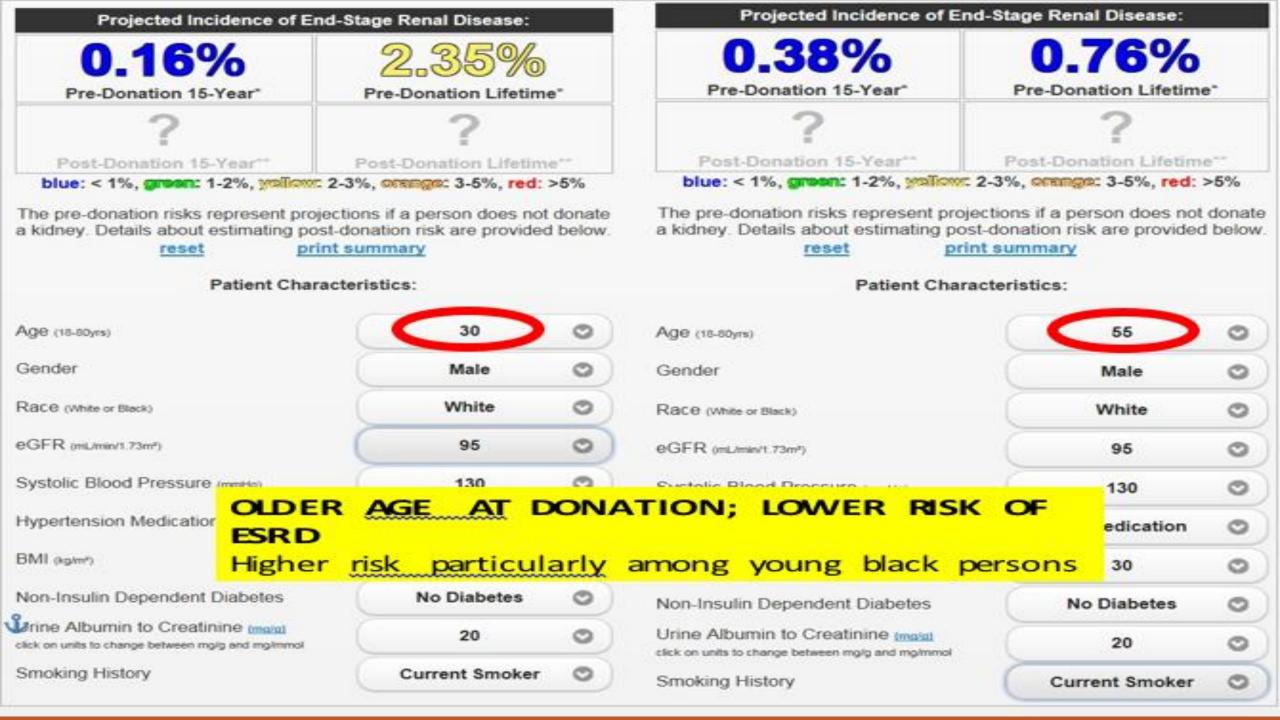
Pre-Donation 15-Year*

Post-Donation 15-Year**

Projected Incidence of En	d-Str⊿e Renal Disease:	
0.19% Pre-Donation 15-Year	0.75% Pre-Donation Lifetin	
Post-Donation 15-Year**	Post-Donation Lifetin	me**
he pre-donation risks represent projekidney. Details about estimating po-		ot dona
a kidney. Details about estimating por	C IC MELY	
Age (18-80yrs)	octeristics:	0
		0
Age (18-80yrs) Gender	50	
Age (18-80yrs)	Male Male	0

For example, if a transplant program sets the acceptable lifetime postdonation

ESKD risk threshold at 5%, and assumes a donation attributable RR of 3.5 to 5.3 according to sex and race, then the acceptable predonation lifetime ESKD risk threshold would be approximately 1.0-1.5.





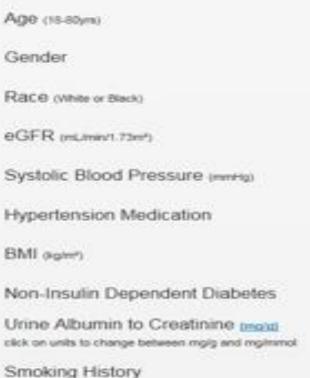
GFR

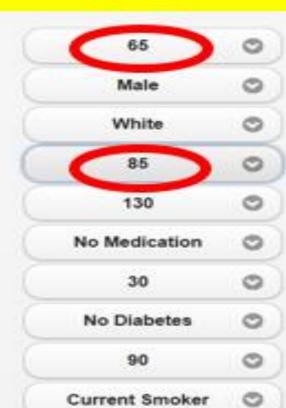


Current Smoker

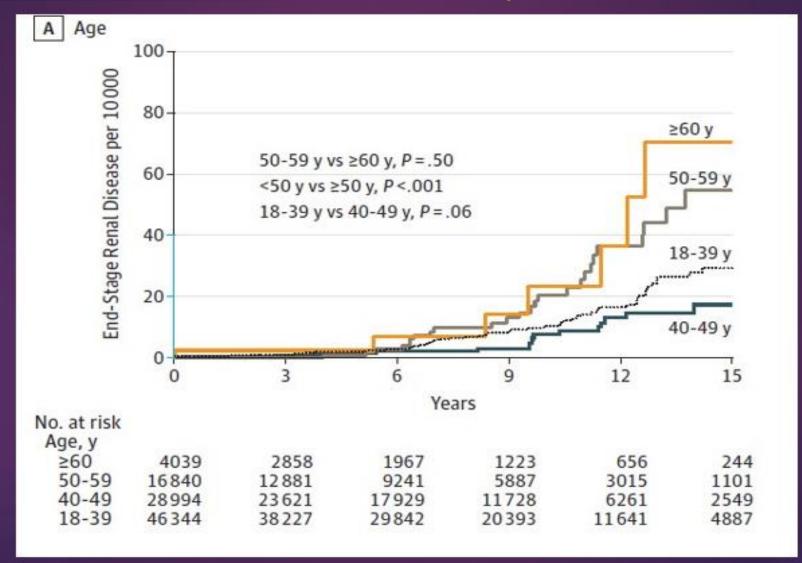
0

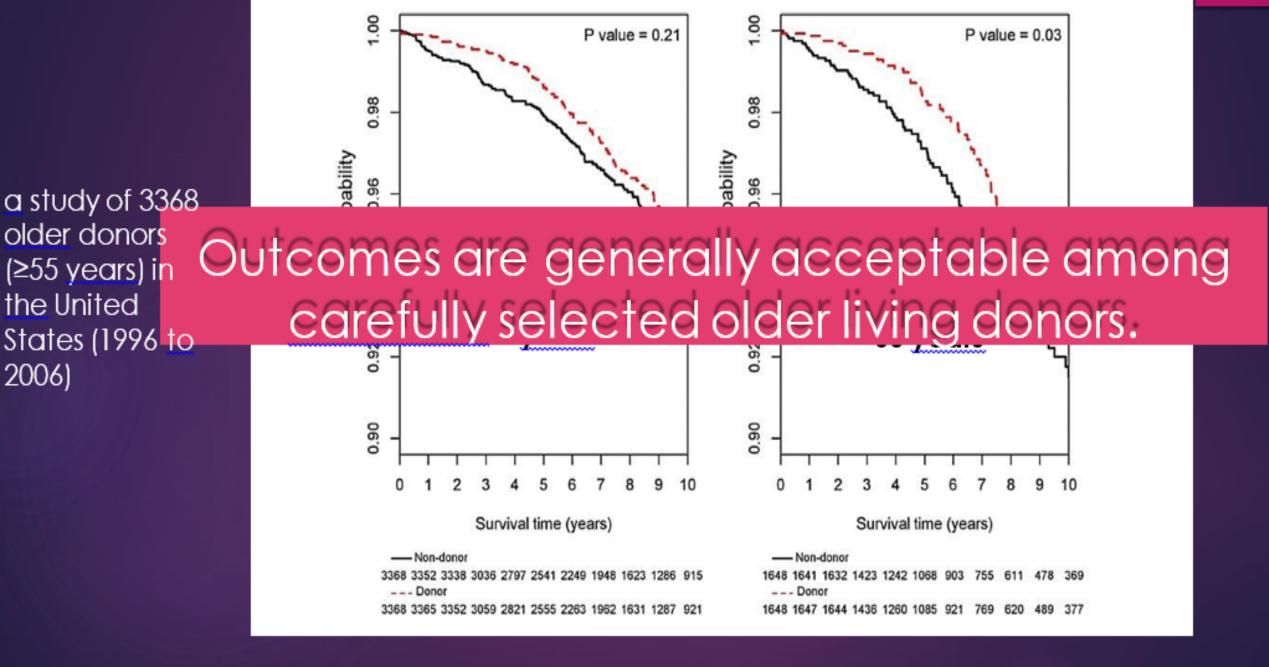
Y, Smoking History





Cumulative Incidence of End-Stage Renal Disease in Live Kidney Donors





Outcome of the living kidney donor



consequences of kidney donation on the living donor health, considering:

- ✓ very short term (linked to the surgery)
- ✓ short term (effect of nephrectomy on glomerular filtration rate)
- ✓ long term (risk of mortality, chronic kidney disease, proteinuria and hypertension)



The following should be performed at least annually postdonation:

- Blood pressure measurement
- BMI measurement
- Serum creatinine measurement with GFR estimation
- Albuminuria measurement
- Review and promotion of a healthy lifestyle including regular exercise, healthy diet and abstinence from tobacco
- Review and support of psychosocial health and well-being

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LONG-TERM POST NEPHRECTOMY ISSUES

- □ Within days to weeks after uninephrectomy, hyperfiltration in the remaining kidney increases the GFR to about 75% to 80% of predonation value
 □ Similar to the nondonating population, an additional 5 mL per minute loss in GFR per decade occurred after donating
- ☐ This acute compensation is, however, **less efficient in elderly** or **obese patients** as it is related to the use of the renal functional reserve

LONG-TERM POST NEPHRECTOMY ISSUES

proteinuria

- ☐ Urine albumin excretion, attributable to single nephron hyperfiltration may be elevated but **is usually low grade** and not associated with a higher risk for renal dysfunction
- ☐ this complication occurs only in a minority of donors
- ☐ The proteinuria will be <1 g/24 h in the vast majority of donors More severe and nephrotic proteinuria are exceptional

LONG-TERM POST NEPHRECTOMY ISSUES



hypertension

- ☐ The incidence of **hypertension requiring treatment** increases with time following kidney donation, but most studies suggest a similar frequency compared with an age-matched population
- ☐ it would appear reasonable to **target a SBP**≤ **130 mm Hg** for long-term follow-up of donors

Meta-analysis: risk for hypertension in living kidney donors

Figure 1. Meta-analysis of controlled studies of systolic blood pressure (SBP) and diastolic blood pressure (DBP) at least 5 years after kidney donation.

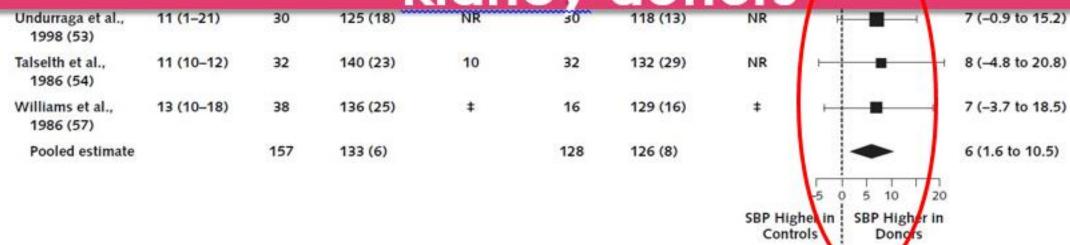
Study, Year (Reference)

Donors, after Donation

Control Participants

Mean Difference in SBP (95% CI), mm Hg

SBP and DBP were 6 and 4 mmHg higher in kidney donors



Long-Term Medical Care

- ➤ In the United States, transplant centers are required to report follow-up donors at discharge (or 6 weeks postdonation, which ever comes first) 6 months, 1 year, and 2 years after donation
- ➤ Routine checkups, cancer screening appropriate for age, regular aerobic exercise, weight reduction, tobacco avoidance, and excessive alcohol abstinence should be emphasized

Long-Term Medical Care(cont..)

- ☐ Kidney donors with established medical issues before donation, such as mild hypertension, history of nephrolithiasis, or obesity, should have more frequent follow-up
- Donors should be **discouraged** from using **high-protein diets** for weight loss or **protein supplements** for body building because they may contribute to hyperfiltration injury
- ☐ They should be advised to avoid **longterm regular use** of nonsteroidal anti-inflammatory drugs

Summary

- ➤ Donor kidney function **should be expressed** as GFR(mL/min per 1.73 m2) & **not** as serum creatinine concentration
- > mGFR using an exogenous filtration marker is the most accurate confirmatory test
- ➤ The **most commonly** used measure of evaluating GFR in clinical practice is based on a 24-h creatinine clearance (**mCrCl**)
- future risk of developing kidney failure necessitating treatment with dialysis or transplantation is slightly higher
- A web-based calculator has been developed to compute post-test probabilities http://ckdepi.org/equations/donor-candidate-gfr-calculator for mGFR above or below various threshold probabilities
- Long-term risk in the absence of donation can be computed from http://www.transplantmodels.com/esrdrisk
- Advanced age can increase the risk for perioperative complications, but there is **no** mandated upper age limit for living kidney donation

thanks

