

Laboratory Testing for Chronic Kidney Disease Diagnosis and Management

Chronic kidney disease is defined as abnormalities of kidney structure or function, present for greater than 3 months, with implications for health.¹ Diagnostic criteria include a decreased glomerular filtration rate (GFR) or presence of 1 or more other markers of kidney damage.¹ Markers of kidney damage include a histologic abnormality, structural abnormality, history of kidney transplantation, abnormal urine sediment, tubular disorder-caused electrolyte abnormality, or an increased urinary albumin level (albuminuria).

This Test Guide discusses the use of laboratory tests that may aid in identifying chronic kidney disease and monitoring and managing disease progression, comorbidities, and complications. The tests discussed include measurement and estimation of GFR as well as markers of kidney damage. A list of applicable tests is provided in the **Appendix**. The information is provided for informational purposes only and is not intended as medical advice. A physician's test selection and interpretation, diagnosis, and patient management decisions should be based on his/her education, clinical expertise, and assessment of the patient.

ESTIMATION OF GFR

Direct measurement of GFR may be problematic. The gold standard for measured GFR (mGFR) is the inulin clearance method, but this test is difficult to perform in clinical practice. Clearance measurements using iohexol, ¹²⁵I-iothalamate, and several chelated isotopic radiotracers provide alternatives to the inulin reference method for mGFR, but each has limitations.² These methods are usually reserved for specific circumstances, such as determining correct drug dosing in therapy that requires the most accurate assessment of kidney function, or when clinical decision-making is affected by having disparate results for estimated GFR (eGFR, see below).

Direct measurement of creatinine clearance and urea clearance is also an alternative to assess kidney function. Notably, the average of the 2 values is similar to GFR measured by inulin clearance (<20 mL/min/1.73 m²) in patients not on dialysis.³ In these patients, the overestimation due to creatinine secretion is offset by underestimation due to urea reabsorption, leading to an average value that accurately reflects GFR.³ However, this method of calculating GFR is prone to error due to inaccurate timing of blood sampling, incomplete urine collection over 24-hours, or over collection of urine beyond 24-hours.^{2,3}

Given that direct measurement of GFR may be problematic, eGFR, using either creatinine- or cystatin C-based measurements, is most commonly used to diagnose CKD in clinical practice.

Creatinine-Based eGFR

GFR is typically estimated using the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation.⁴ The CKD-EPI equation uses serum-creatinine measurements and the patient's age (≥18 years old), sex, and race (African American vs non–African American). Creatinine-based eGFR is recommended by the Kidney Disease Improving Global Outcomes (KDIGO) 2012 international guideline for initial assessment of GFR.¹

Creatinine-based eGFR may be imprecise for certain individuals (see *Cystatin C-based eGFR* below). In addition, confirmation of GFR using another method may be required to avoid misclassification of some patients. Cystatin C-based eGFR provides an alternative when creatinine-based estimates are not appropriate.

Cystatin C-Based eGFR

Being less influenced by diet and muscle mass, cystatin C-based eGFR testing is appropriate for patients in whom creatinine-based results may be misleading.⁵ These patients include pregnant women, patients with acute illness, patients with serious comorbid conditions, people with extremes of muscle mass (eg, bodybuilders, patients with amputation, paraplegia, muscle-wasting disease, or a neuromuscular disorder), patients suffering from malnutrition, those with a vegetarian or low-meat diet, and those taking creatine dietary supplements.

In addition, the KDIGO guideline recommends using cystatin C-based eGFR, or creatinine plus cystatin C-based eGFR, to confirm chronic kidney disease when creatinine-based eGFR indicates a mild to moderately high risk of chronic kidney disease progression (45 to 59 mL/min/1.73 m²) in a patient without albuminuria (**Figure 1**).¹

Figure 1. Frequency of Monitoring Chronic Kidney Disease Based on Risk of Disease Progression Assessed Using eGFR and Urine Albumin-Creatinine Ratio (Kidney Profile, test code 39165)

			Albuminuria categories and ACR ranges (mg/g creatinine)		
			Normal to mildly increased	Moderately increased	Severely increased
			<30	30-300	>300
CKD stage and eGFR range (mL/min/1.73 m²)	1 and 2	≥60	1	1	2,R
	3A	45-59	1,C	2	3,R
	3В	30-44	2		3,R
	4	15-29	3,R	3,R	≥4,R
	5	<15	≥4,R	≥4,R	≥4,R

Low risk: monitor yearly if evidence of kidney damage (eg, indicated by imaging or biopsy). The NKDEP recommends that actual values above 60 mL/min/1.73m² be reported only as >60 due to variability near the upper limit of the reference range.¹²

Moderately high risk: monitor yearly

High risk: monitor 2 times yearly

Very high risk: monitor 3 times yearly

Very high risk: monitor ≥4 times yearly

ACR, albumin-creatinine ratio; C, confirm using eGFR based on (1) cystatin C (test code 94588) or (2) creatinine plus cystatin C; CKD, chronic kidney disease; eGFR, estimated glomerular filtration rate; NKDEP, National Kidney Disease Education Program; R, refer to specialist. This figure was adapted from references 1 (with permission) and 12, is provided for informational purposes only as a guide for using laboratory tests, and is not intended as medical advice. A physician's test selection and interpretation, diagnosis, and patient management decisions should be based on his/her education, clinical expertise, and assessment of the patient.



A large meta-analysis has shown cystatin C-based eGFR improves risk classification for adverse outcomes (death, cardiovascular disease-related death, and end-stage renal disease) across diverse populations.⁶ However, cystatin C-based eGFR may be more affected by some non-GFR determinants such as thyroid disorders, corticosteroid use, and smoking.² In addition, small but significant associations of cystatin C levels with diabetes, obesity, and inflammation have been reported.^{6,7} For these reasons, creatinine-based eGFR is recommended for patients without contraindications.

Finally, interpretation of eGFR is challenging for patients with severe obesity (BMI >40); for this group, creatinine-based eGFR may be overestimated and the utility of cystatin C-based eGFR needs to be further investigated.⁸

ALBUMINURIA AND PROTEINURIA

Albuminuria indicates increased glomerular permeability, a characteristic of chronic kidney disease, and is assessed with either the urine albumin-creatinine ratio or albumin excretion rate over 24 hours. The urine albumin-creatinine ratio obtained from a random sample is more convenient and appropriate in the context of chronic kidney disease. However, carefully performed 24-hour specimen collection is more accurate and appropriate in some circumstances (eg, in glomerular disease when small discrepancies between the random and 24-hour results may influence high-risk therapeutic dosing).⁹ A urine albumin-creatinine ratio result of \geq 30 mg/g (albumin excretion rate \geq 30 mg/24 hours) is evidence of albuminuria (30 to 300 mg/g was formerly referred to as "microalbuminuria" and >300 mg/g as "macroalbuminuria").¹

Proteinuria (urinary total protein-creatinine ratio \geq 150 mg/g) may also indicate increased glomerular permeability and chronic kidney disease, but may have other causes that are distinguished by elevated nonalbumin proteins.¹ Examples of other causes of proteinuria include defective tubular resorption (elevated α 1- and β 2-microglobulin) and tubular resorption capacity being exceeded, as observed in myeloma (elevated immunoglobulin light chains).¹

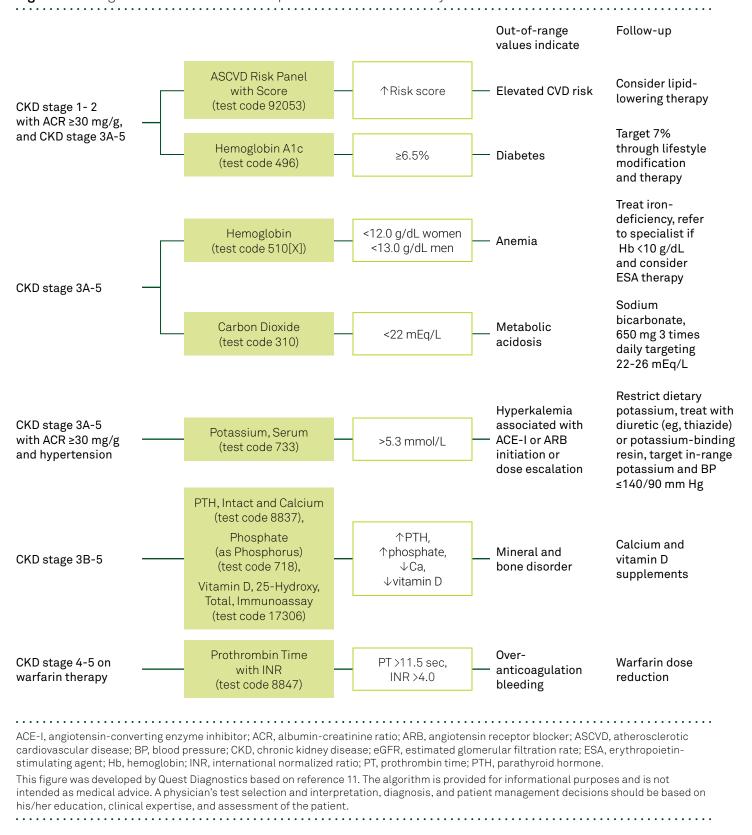
GFR, ALBUMIN-CREATININE RATIO, AND DISEASE MANAGEMENT

A GFR <60 mL/min/1.73 m² for >3 months and/or urine albumin-creatinine ratio ≥30 mg/g for >3 months define chronic kidney disease.¹ Combined, these test results provide a "Kidney Profile" recommended by the National Kidney Foundation for diagnosing and managing chronic kidney disease in at-risk patients (**Figure 1**).^{1,10-12} The results are important independent risk predictors of major adverse cardiovascular events (myocardial infarction or stroke).¹³

Approaches to monitoring and managing cardiovascular risk, diabetes, and other comorbidities and complications associated with the various stages of chronic kidney disease are provided in **Figure 2.**¹¹



Figure 2. Testing for Comorbidities and Complications of Chronic Kidney Disease





APPENDIX

Quest Diagnostics offers many tests and panels for diagnosis and management of chronic kidney disease. Test offerings range from health screenings for abnormal eGFR, proteinuria, and/or albuminuria, to tests for management of CKD and its

comorbidities and complications. For example, the Diabetes, Advancing Chronic Kidney Disease Management Panel (test code 91713) combines many of these tests according to the recommendations of the American Diabetes Association.¹⁴

APPENDIX TABLE

Tests Used in Diagnosis and Management of Chronic Kidney Diseasea,b

Test code	Test name	Primary clinical use and differentiating factors
Screening, di	agnosis, and monitoring	
6517	Albumin, Random Urine with Creatinine	Detect albuminuria using albumin-creatinine ratio
	Includes albumin and creatinine, random urine 8459(X).	
15281	Albumin, 24-Hour Urine with Creatinine	Detect albuminuria using albumin excretion rate corrected for
	Volume measurement adds additional CPT code and charge.	creatinine excretion
4555	Albumin, 24-Hour Urine without Creatinine	Detect albuminuria using albumin excretion rate
	Volume measurement adds additional CPT code and charge.	
10165	Basic Metabolic Panel	Screen for acute and chronic kidney disease
	Includes BUN/creatinine ratio (296), calcium (303), carbon dioxide (310), chloride (330), creatinine (375), eGFR (calculated), glucose (483), potassium (733), and sodium (836).	Assess blood glucose, calcium, and electrolyte balance
296	BUN/Creatinine Ratio	Screen for acute and chronic kidney disease
	Includes BUN (294) and creatinine and calculated BUN/creatinine ratio and eGFR.	
10231	Comprehensive Metabolic Panel	Screen for acute and chronic kidney disease and liver
	Includes albumin (223), albumin/globulin ratio (calculated), alkaline phosphatase (234), ALT (823), AST (822), BUN/creatinine ratio (296), calcium (303), carbon dioxide (310), chloride (330), creatinine (375), eGFR (calculated), globulin (calculated), glucose (483), potassium (733), sodium (836), total bilirubin (287), and total protein (754).	dysfunction Assess blood glucose, calcium, and electrolyte balance
375	Creatinine	Screen for CKD°
	Includes serum creatinine and eGFR calculation.	Monitor CKD therapy and/or progression in adults
7943	Creatinine Clearance	Screen for CKD°
	Includes serum (375), 24-hour urine (381) creatinine, creatinine clearance and eGFR calculation. Volume measurement adds additional CPT code and charge.	Confirm CKD using 24-hour creatinine measurement
94588	Cystatin C with Glomerular Filtration	Screen for CKD ^d
	Rate, Estimated (eGFR)	Confirm a diagnosis of CKD
		Monitor CKD therapy and/or progression in adults
		(Continued

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APPENDIX TABLE (Continued) Tests Used in Diagnosis and Management of Chronic Kidney Disease^{a,b}

Test code	Test name	Primary clinical use and differentiating factors	
91713	Diabetes, Advancing Chronic Kidney	Screen for CKD in patients with diabetic nephropathy	
	Disease Management Panel	Monitor CKD therapy in patients with diabetes mellitus	
	Includes electrolyte panel (34392) (includes sodium [836], potassium [733], chloride [330], carbon dioxide [310]), hemoglobin (510[X]), intact PTH and calcium (8837), phosphate (phosphorus [718]), total 25-hydroxyvitamin D by immunoassay (17306), serum creatinine (375); albumin, random urine with creatinine (6517).		
91712	Diabetes, Newly Diagnosed and	Screen for CKD in patients recently diagnosed with diabetes	
	Monitoring Panel	mellitus	
	Includes glucose (483); HbA1c (496); hepatic function panel (10256) (includes total protein [754], albumin [223], globulin [calculated], albumin/ globulin ratio [calculated], total [287], direct [285], and indirect [calculated] bilirubin, alkaline phosphatase [234], AST [822], and ALT [823]); lipid	Establish baseline measurements for patients recently diagnosed with diabetes mellitus Monitor patients with diabetes mellitus	
	panel (total [334], HDL [608], and LDL [calculated] cholesterol; triglycerides [896] with reflex to direct LDL [8293] if triglycerides >400 mg/dL; cholesterol/ HDL ratio [calculated]; and non-HDL [calculated]); serum creatinine (375); albumin, random urine with creatinine (6517).		
39165	Kidney Profile	Screen for CKD°	
	Includes albumin, random urine with creatinine (6517) and creatinine with calculated eGFR (375).	Detect albuminuria	
		Monitor CKD therapy and/or progression in adults	
1715	Protein, Total, Random Urine	Detect proteinuria	
	with Creatinine		
10314	Renal Function Panel	Screen for acute and chronic kidney disease	
	Includes albumin (223), BUN/creatinine ratio (296), calcium (303), carbon dioxide (310), chloride (330),	Detect albuminuria	
	creatinine (375), eGFR (calculated), glucose (483), phosphate (as phosphorous) (718), potassium	Assess blood glucose, calcium, phosphorus, and electrolyte	
	(733), and sodium (836).	balance	
7329(X)	Urea Clearance	Screen for CKD°	
	Includes BUN (294), urea nitrogen 24-hour creatinine (973[X]), and calculated urea clearance. Volume measurement adds additional CPT code and charge.	Confirm CKD using 24-hour creatinine measurement	
Other tests fo	or screening, diagnosis, and monitoring	•••••••••••••••••••••••••••••••••••••••	
NA	Dipstick test	Rapidly check for out-of-range urine acidity, protein, glucose, bilirubin, as well as WBC, blood in the urine	
NA	Measured GFR	Confirm a diagnosis of CKD	
		Determine correct drug dosing for individuals with abnormal	
		kidney function and GFR resulting in altered pharmacokinetic	
		-	
		or pharmacodynamic relationships (Continue)	

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APPENDIX TABLE (Continued)

Tests Used in Diagnosis and Management of Chronic Kidney Diseasea,b

Test code	Test name	Primary clinical use and differentiating factors
Tests for mor	nitoring comorbidities and complications ¹¹	
92053	ASCVD Risk Panel with Score	Assess risk for ASCVD in patients with CKD stage 1- 2 with ACR
	Includes total (334), HDL (608), and LDL (calculated) cholesterol; triglycerides (896); cholesterol/HDL ratio (calculated); non-HDL (calculated); 10-year and lifetime atherosclerotic cardiovascular risk scores (calculated); and reflex to direct LDL (8293).	≥30 mg/g or CKD stage 3A-5
310	Carbon Dioxide	Monitor metabolic acidosis in patients with CKD stage 3A-5
510(X)	Hemoglobin	Monitor anemia in patients with CKD stage 3A-5
496	Hemoglobin A1c	Monitor diabetes in patients with CKD stage 1- 2 with ACR
		≥30 mg/g or CKD stage 3A-5
718	Phosphate (as Phosphorus)	Monitor mineral and bone disorder in patients with CKD stage
		3B-5
733	Potassium, Serum	Monitor hyperkalemia associated with ACE-I or ARB initiation
		or dose escalation in patients with hypertension and CKD
		stage 3A-5, ACR ≥30 mg/g
8847	Prothrombin Time with INR	Monitor over-anticoagulation bleeding in patients on warfarin
		therapy with CKD stage 4-5
8837	PTH, Intact and Calcium	Monitor mineral and bone disorder in patients with CKD stage
		3B-5
17306	Vitamin D, 25-Hydroxy, Total,	Monitor mineral and bone disorder in patients with CKD stage
	Immunoassay	3B-5

ACE-I, angiotensin-converting enzyme inhibitor; ACR, albumin-creatinine ratio; ALT, alanine aminotransferase; ARB, angiotensin receptor blocker; ASCVD, atherosclerotic cardiovascular disease; AST, aspartate aminotransferase; BUN, blood urea nitrogen; CKD, chronic kidney disease; eGFR, estimated glomerular filtration rate; HDL, high-density lipoprotein; INR, international normalized ratio; LDL, low-density lipoprotein; NA, not available through Quest Diagnostics; PTH, parathyroid hormone; WBC, white blood cell.

^a Panel components may be ordered separately.

^b Reflex tests are performed at an additional charge and are associated with additional CPT codes.

^c In at-risk adults (>18 years old) including those who have diabetes, cardiovascular disease, hypertension, previous kidney damage, systemic disease with potential kidney involvement (eg, systemic lupus erythematosus), or a family history of CKD, as well as individuals who are moderately obese or ≥65 years old.

^d In patients for whom creatinine-based results may lead to an incorrect diagnosis.



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