

KIDNEYSEQ™ VERSION 4 – 312 GENES

IOWA INSTITUTE OF HUMAN GENETICS
CLINICAL TEST REQUISITION FORM
CLIA ID 16D2053873 | CAP ID: 8864771

431 Newton Road, 140 EMRB
Iowa City, IA 52242-1078
Phone: 319-335-3688 Fax: 319-335-3484
Email: iihg@uiowa.edu
www.medicine.uiowa.edu/humangenetics/

Patient Information	Specimen Information
- Place patient ID sticker here -	Specimen Collection Date: ____/____/____ MM/DD/YYYY
Name: _____ Last First	Specimen Type: <input type="checkbox"/> ~6mL EDTA whole blood <input type="checkbox"/> ~3mL EDTA whole blood (pediatric) <input type="checkbox"/> 10µg DNA @110ng/ml (minimum) CLIA ID: _____
DOB: ____/____/____ MM/DD/YYYY	*DNA must be extracted in a CLIA certified laboratory. For labs outside of the USA, equivalent requirements apply.
Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female	MRN: _____
	Accession #: _____

Clinical Indication for Testing (required)

1. What is your working clinical diagnosis?
2. What are the salient clinical features?
3. What are the relevant laboratory data?
4. Is there a family history? Yes No (If yes, provide details and attach pedigree)

***Providing more detailed and relevant clinical records will help in the interpretation of genetic findings. Please include a phone number for the treating clinician as the IIHG may need to discuss findings or obtain additional clinical information.

Clinical information can be emailed to iihg@uiowa.edu or faxed to 319-335-3484

Patient Race	Patient Ethnic Group/Identification
<input type="checkbox"/> Asian <input type="checkbox"/> African / African American <input type="checkbox"/> Caucasian <input type="checkbox"/> Native American (American Indian or Alaska Native) <input type="checkbox"/> Native Hawaiian/Pacific Islander	<input type="checkbox"/> Unknown/Unspecified <input type="checkbox"/> Multiracial/Two or More Races <input type="checkbox"/> Declined <input type="checkbox"/> Other _____
	<input type="checkbox"/> Non-Hispanic <input type="checkbox"/> Hispanic or Latino <input type="checkbox"/> Jewish – Sephardic <input type="checkbox"/> Jewish – Ashkenazi <input type="checkbox"/> Eastern Indian <input type="checkbox"/> Mediterranean
	<input type="checkbox"/> Unknown <input type="checkbox"/> Declined

Billing Information (required prior to testing)

Institutional billing ONLY at this time. The IIHG does not do direct insurance billing or direct patient billing at this time.

Institution Name:	Billing Contact:
Institution Code:	Billing Contact Email:
Street Address:	Phone:
City: State: Zip:	Fax:

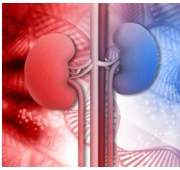
Reporting Information (fax# required to send results)

Reporting Information (fax# required to send results)	Additional Report Recipient (fax# required to send results)
Health Care Provider:	Health Care Provider:
NPI:	NPI:
Institution:	Institution:
Department	Department
Email:	Email:
Phone: Fax:	Phone: Fax:

Test Menu – select one or more options

KidneySeq™: <input type="checkbox"/> Comprehensive testing (312 genes) <input type="checkbox"/> CAKUT (56 genes) <input type="checkbox"/> Ciliopathies/tubulointerstitial diseases (85 genes) <input type="checkbox"/> Disorders of tubular ion transport (72 genes) <input type="checkbox"/> Glomerulopathies (78 genes) <input type="checkbox"/> Nephrolithiasis/ nephrocalcinosis (35 genes)	Single gene tests: <input type="checkbox"/> APOL1 test: G1 and G2 alleles <input type="checkbox"/> Familial testing (segregation analysis) <ul style="list-style-type: none"> • Patient information on page 1 should match the accompanying sample • Family member information on page 2 should identify other members being tested, including the index patient. CPT codes on page 2
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IIHG ID:
(Office use only)



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Patient Name: _____ Last First M		DOB: ____ / ____ / ____ MM / DD / YYYY	IIHG ID: (Office use only)
Familial Testing (segregation analysis) (testing to determine if other family members carry the same genetic variant identified in the index person)			
Family member #1		Family member #2	
Last Name: _____		Last Name: _____	
First Name: _____		First Name: _____	
Relationship to patient listed on page 1: _____		Relationship to patient listed on page 1: _____	
DOB: ____ / ____ / ____ MM / DD / YYYY		DOB: ____ / ____ / ____ MM / DD / YYYY	
<input type="checkbox"/> Asymptomatic <input type="checkbox"/> Symptomatic		<input type="checkbox"/> Asymptomatic <input type="checkbox"/> Symptomatic	
If more than two persons are to be tested, please include a pedigree. For assistance in constructing a pedigree please see https://medicine.uiowa.edu/humangenetics/resources/how-draw-pedigree			
Sample Requirements			
<ul style="list-style-type: none"> Samples must be labeled with the patient's full name and date of birth. Samples received without both pieces of information cannot be processed and will be discarded. Biological parent or other relative samples must be labeled with that person's full name and date of birth, NOT the index patient's information. Samples received without both pieces of information cannot be processed and will be discarded. 6 mL whole blood in lavender EDTA tube (3 mL pediatric minimum). OR 10µg DNA (A260/A280 1.8-2) resuspended in 0.1mM EDTA (10mM Tris HCl, 0.1mM EDTA, pH 8, Teknova Cat #T0220) *DNA must be extracted by a CLIA certified laboratory. For laboratories outside of the USA, equivalent requirements apply. Please include a copy of the accreditation standards and certificate of accreditation. 			
Shipping Requirements			
<ul style="list-style-type: none"> Samples may be shipped at ambient temperature or with an ice pack. DO NOT freeze sample. If the sample is collected on a Friday, refrigerate the sample over the weekend for shipping the following week. Samples should be shipped for next day delivery, Tuesday-Friday. Samples are not received on weekends or US holidays. Please contact us at iihg@uiowa.edu or 319-335-3688 with any questions or concerns. Ship overnight to: Iowa Institute of Human Genetics University of Iowa 431 Newton Road, 140 EMRB Iowa City, IA 52242 Phone: 319-335-3688 			
CPT Codes:			
<ul style="list-style-type: none"> KidneySeq™ <ul style="list-style-type: none"> Comprehensive testing: 81404, 81405, 81406, 81407, 81408, 81479 CAKUT: 81405, 81406, 81407, 81479 Ciliopathies/tubulointerstitial diseases: 81404, 81405, 81406, 81407, 81408, 81479 Disorders of tubular ion transport: 81404, 81405, 81406, 81407, 81479 Glomerular diseases: 81404, 81405, 81406, 81407, 81408, 81479 Nephrolithiasis/nephrocalcinosis: 81404, 81405, 81406, 81407, 81479 APOL1 test: 81479 Familial Testing: 81479 			