

## UriFind® Urothelial Carcinoma Test Requisition Form

### PART 1. PATIENT INFORMATION (REQUIRED)

Last Name	First Name	Middle Initial	Date of Birth
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Street Address	City	State	Zip Code
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Preferred Contact Phone Number			
<input type="text"/>			
<input type="checkbox"/> Home <input type="checkbox"/> Mobile <input type="checkbox"/> Work			
Gender <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other		ETHNICITY IDENTIFICATION <input type="checkbox"/> Hispanic or Latino <input type="checkbox"/> Not Hispanic or Latino <input type="checkbox"/> Unknown	
Race Identification <input type="checkbox"/> African American <input type="checkbox"/> Asian <input type="checkbox"/> White <input type="checkbox"/> American Indian or Alaska Native <input type="checkbox"/> Native Hawaiian or Other Pacific Islander <input type="checkbox"/> Other			

### PART 2. SAMPLE COLLECTION TYPE (REQUIRED)

<input type="checkbox"/> Urine	Collection Date & Time <input type="text"/>	I attest that the urine sample was properly collected in a sterile container, mixed with the Preservative provided and placed in refrigerator within 30 minutes of collection.  Signature and Date
<input type="text"/>		

Urine sample should be mixed with the Preservative provided and placed in refrigerator within 30 minutes after collection.

### PART 3. PRACTICE/CLINIC INFORMATION (REQUIRED)

Practice/Clinic Name	Physician's NPI Number		
<input type="text"/>	<input type="text"/>		
Street Address	City	State	Zip Code
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Email	Phone Number		
<input type="text"/>	<input type="text"/>		
Clinic Name or Account Number	Physician Name		
<input type="text"/>	<input type="text"/>		

#### Medical Professional Consent

My signature constitutes a Certification of Medical Necessity, and I hereby authorize and order DiaCarta, Inc. to perform testing for this patient as indicated on this requisition, I have reviewed the medical consent on this form and will provide test interpretation to the patient as appropriate.

Signature and Date

### PART 4. ICD-10 CODES (MOST COMMONLY USED ICD-10 CODES) (REQUIRED)

<input type="checkbox"/> R31.0 Gross hematuria	<input type="checkbox"/> R31.1 Microscopic hematuria	<input type="checkbox"/> R31.2 Other Microscopic hematuria	<input type="checkbox"/> R31.9 Hematuria, unspecified
<input type="checkbox"/> Z12.6 Encounter for screening for malignant neoplasm of bladder		<input type="checkbox"/> Additional ICD-10 code(s)	

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### PART 5. PATIENT INSURANCE INFORMATION (REQUIRED FOR INSURANCE COVERAGE)

Where applicable please include a photocopy of insurance card(s) (both sides); for Credit Card please include a photocopy of a valid credit card (both sides).

Please select a billing option & complete the information below

☐ Insurance
 ☐ Cash Pay
 ☐ Credit Card
 ☐ Check
 ☐ Client Bill

Primary Insurance Carrier

Primary Insurance ID No.

Primary Insurance Group No.

Patient Relationship to Insured

☐ Self
 ☐ Spouse
 ☐ Dependent
 ☐ Other

Secondary Insurance Carrier

Secondary Insurance ID No.

Secondary Insurance Group No.

Patient Relationship to Insured

☐ Self
 ☐ Spouse
 ☐ Dependent
 ☐ Other

### PART 6. PATIENT CONSENT

I \_\_\_\_\_ (Patient or legal guardian name), request and authorize the DiaCarta Clinical Laboratory to perform the requested test(s) for the person(s) listed above. I acknowledge the benefits, risks, and limitations outlined below. I understand that my specimen(s) will be submitted to DiaCarta for the purpose of lab testing. I authorize DiaCarta to store my specimen in case additional testing is necessary. The DiaCarta Clinical Laboratory does not return patient samples. I can request additional tests or send out samples to other institutions if there is enough sample. Once my test result has been released, remaining samples may be de-identified to be used for laboratory quality control or research. I can withdraw my consent at any time by calling the DiaCarta Laboratory at (800) 246-8878. My signature below indicates that I have read the above information. All my questions have been answered and my inquiries regarding the purpose of this test have been discussed and fully understood by me.

Patient Name  
(Print)

Date

Patient Signature

**AFFIX SPECIMEN  
BARCODE HERE**

**THE END OF THE DOCUMENT**