

Dianon Systems, Inc 840 Research Parkway Oklahoma City, OK 73104 Phone: (800) 411-1839 or (405) 290-4000 Fax: (800) 334-5590 or (405) 290-4046

Patient Support Services

Order Number:

_				Order	Tuilibe	••			
P	atient and Billing Infor	mation							
	Patient Name			MRN		_			
	Shipping Address	Home Phone ()							
ED	City, State, Zip	Work Phone () Race							
E	Sex M F Dat								
ם מ	Bill: Practice/Facility Medicare Medicaid Insurance Patient								
REQUIR	Insurance Carrier				Phone Number ()				
			Policy #						
					Group # PIES OF BOTH CARDS, FRONT AND BACK □ Billing Information Attached				
Re	questing Physician/NPI		Physician/Au	thorized Signatu	ire		Kit To Be Sent (1	within 3 months)	
_				_			Month/Yr:	/	
	UroStone [®] Metabolic	Manage	ement Sys	stem					
	ICD-CM Code ^A	Ending Collection Date							
	Collection Number of 24-hour Urine Collection Numb			ction Kits reque	tion Kits requested: Patient has i			increased urinary output (>4 L/day)	
	Monitoring Profiles								
ST				oStone® 24 Cys	stine * (Creatinine, Qual Cystine)*	Creatinine C	learance (serum creatinine)	
QUES	UroStone® 24 Citrate (Cre	rate (Citrate) Uro			Stone® 24 Urio Acid (Creatinine,			Requires serum within 48 hrs of urine collection Required:	
П	UroStone® 24 * Calcium, Citrate, Creatinine, Magnesium, Oxalate, pH, Phosphorus, Qualitative Cystine*, Sodium, Uric Acid Wtlbs								
<u>~</u>	UroStone® Max24 * UroSton	e [®] 24 + Ammon	ia, Chloride, Potassi	ium, Sulfate					
EST	Individual Ammonia Citrate Tests Calcium Creatinine Cystine, Company Cystine, Company				☐ Magnesium ☐ Phosphorus ☐ Sodium ☐ Oxalate ☐ Potassium ☐ Sulfate				
					D pH	แลเย	Total Protein	Uric Acid	
	*Quantitative Cystine performed o	n positive Qua	litative Cystine at	t additional charge					
E	ladder Cytology								
IC	D-CM Code [*]			Microcyte	PLUS®/	Urine Cytolo	ogy Profiles		
Collection Date				☐ 994 Hematuria Profile (Tablet Preservative Kit) Cytodiagnostic Urinalysis Correlating Cytology (by concentration technique, includes Pap and					
Sp	ecimen Type:			-				oalbumin, and Total Protein	
☐ Voided Urine ☐ Post Cysto Void Date of last cysto			□ VU3 Cytology Plus Monitoring Profile						
IN	DIVIDUAL TESTS:	Cytology* (Pap and Feulgen stain)							
□ VU6 Pap Stain (only) Cytology				☐ VU1D Bladder Cancer FISH/Cytology Pathodiagnostic Profile					
Require Monitoring Kits			Bladder Cancer FISH Assay and Cytology* (Pap and Feulgen Stain); including integrated cytomolecular diagnostic interpretation with clinical correlation by pathologist (MD)						
	(600D Bladder Cancer FISH Ass	•	g diagnostic	□ VU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic Profile					
	nterpretation by Pathologist (M	D)		Cytology* (Pap and Feulgen stain), reflex to Bladder Cancer FISH (Pathologist review) on					
	equire Tablet Preservative Kits 32 Microglobulin 🔲 Microalbu	ımin 🗖 To	ital Protein	atypical cytol	ogy result		lective cellular enhance	ment, see reverse for CPT codes	
				cificity Required)		30			
	gnosis/Signs/Symptoms in ICD-CM format in e ordering tests for which Medicare or Medicaio				er tests that	are medically necess		Necessity of ABN Completion on reve tment of the patient.	
۱ny	Questions? Call 1-800-280-8	484	F	ax Form to 1-8	00-334-!		22 Laboratory Corporation o	f America® Holdings. All rights reserv 261N Rev 07/19/20	
	Dear Patient,	_							
	n an effort to identify the	,		_		•	•	. ,	
urine be analyzed. Your doctor has chosen Labcorp's laboratory specializing in urology, to p								3	
	ou will be asked to provic	le a urine	specimen.	Labcorp will	be shi	pping a urir	ne collection kit	with	
İ	nstructions to your home.								

At this time, you do not need to do anything. Your physician's office will fax the request to Labcorp.

please feel free to contact Patient Support Services at 1-800-280-8484.

Your collection container and instructions will be shipped to you automatically. If you should have any questions

Test Combination/Panel Policy

Labcorp's policy is to provide physicians, in each instance, with flexibility to choose appropriate tests to assure that the convenience of ordering test combinations/panels do not distance physicians who wish to order a test combination/panel from making deliberate decisions regarding which tests are truly medically necessary. All the tests offered in test combinations/panels may be ordered individually using the Labcorp request form. Labcorp encourages clients to contact their local Labcorp representative or Labcorp location if the testing configurations shown here do not meet individual needs for any reason, or if some other combination

In an effort to keep our clients fully informed of the content, charges and coding of its test combinations/panels when billed to Medicare, we periodically send notices concerning customized test combination/panels, as well as information regarding patient fees for all Labcorp services. We also welcome the opportunity to provide, on request, additional information in connection with our testing services and the manner in which they are billed to physicians, health care plans, and patients.

The CPT Code(s) listed here are in accordance with the current edition of Current Procedural Terminology, a publication of the American Medical Association. CPT codes are provided for the convenience of our clients; however, correct coding often varies from one carrier to another. Consequently, the codes presented here are intended as general guidelines and should not be used without confirming with the appropriate payor that their use is appropriate in each case. All laboratory procedures will be billed to third-party carriers (including Medicare and Medicaid) at fees billed to patients and in accordance with the specific CPT coding required by the carrier. Microbiology CPT code(s) for additional procedures such as susceptibility testing, identification, serotyping, etc. will be billed in addition to the primary codes when appropriate. Labcorp will process the specimen for a Microbiology test based on source.

Medical Necessity

Determining Necessity of Advance Beneficiary Notice of Non-coverage (ABN) Completion*

- Determining Necessity of Advance Beneficiary Notice of Non-coverage (ABN) Completion*

 1. Diagnose. Determine your patient's diagnosis.

 2. Document. Write the diagnosis code(s) on the front of this requisition.

 3. Verify. Determine if the laboratory test(s) ordered for the patient is subject to Local Coverage Determination or National Coverage Determination. This information can be located in the policies published by your Medicare Administrative Contractor (MAC), CMS, or www.labcorp.com/MedicareMedicalNecessity.

 4. Review. If the diagnosis code for your patient <a href="https://docs.not/does.not

How to Complete an Advance Beneficiary Notice of Non-coverage (ABN)
Medicare is very specific in requiring that all of the information included on the ABN be completed. Additionally, Labcorp requests that the specimen number or bar code label be included on the form. To be valid an ABN must:

- the form. To be valid an ABN must:

 1. Be executed on the CMS approved ABN form (CMS-R-131)

 2. Identify the Medicare Part B Beneficiary, using the name as it appears on the patient's red, white and blue Medicare card

 3. Indicate the test(s)/procedure(s) which may be denied within the relevant reason column

 4. Include an estimated cost for the test(s)/procedure(s) subject to the ABN

 5. Have 'Option 1', 'Option 2', or 'Option 3' designated by the beneficiary

 6. Be signed and dated by the beneficiary or his/her representative prior to the service being rendered

Symbols used to designate Medicare medical review as of 07/01/2022

- @ = Subject to Medicare medical necessity guidelines
 % = Subject to Medicare frequency guidelines
 # = Medicare deems investigational. Medicare does not pay for services it deems investigational.

	UroStone®	Testing CPT Code	:s
Ammonia	82140	Phosphorus	84105
Calcium	82340	Potassium	84133
Chloride	82436	Qualitative Cystine	82127
Citrate	82507	Quantitative Cystine	82131
Creatinine	82570	Sodium	84300
Magnesium	83735	Sulfate	84392
Oxalate	83945	Total Protein	84156
рН	83986	Uric Acid	84560

Explanation of MicrocytePLUS®/Urine Cytology Testing

Hematuria Profile I - Urine Cytology for directing further evaluations of patients currently not monitored for TCC who present with hematuria or other signs of urinary tract or renal disease. Feulgen performed on all urine CYTOLOGY PROFILES.

Urine Volume – Provide a minimum of 50mL urine for optimum cellularity.

Urine Viability - Hematuria to 5 days, Bladder Cancer FISH to 7 days, Cytology to 8 days.

Bladder Cancer FISH Cytology Pathodiagnostic Profile for therapeutic monitoring of patients with a history of TCC and for initial diagnosis of patients presenting with hematuria with suspicions of TCC.

Bladder Cancer FISH Pathodiagnostic Test is FISH Assay, including diagnostic interpretation with clinical correlation by pathologist (MD).

Bladder Cancer FISH results are intended for use as a method for monitoring for tumor recurrence in patients previously diagnosed with bladder cancer

MicrocytePLUS®/Urine Cytology CPT Codes					
994 Hematuria Profile - Urine Cytology	88108, 88313, 81003, 82232, 82043, 84156				
K600D Bladder Cancer FISH Pathodiagnostic	88120				
VU1D Bladder Cancer FISH / Cytology Pathodiagnostic Profile	88112, 88120				
VU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic	88112, if reflexed 88120				
VU3 Cytology Plus Monitoring Profile	88112				
VU6 Cytology Pap Stain Only	88112				

Explanation of Reflex Testing Quantitative Cystine

When a qualitative cystine is positive, a quantitative cystine will be performed at an additional

Specimen Collection Information

- All Urine Cytologies, FNA's, and Fluid Aspirates must be submitted in the cytology alcohol fixative provided.
- Hematuria Specimens must be collected in a preservative tablet kit
- Do not collect first morning void for 24-hour urine specimens