

1A
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1C
Item# 0050659 Form # 260N.32

1A
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1D

PHYSICIAN	Physician/Authorized Signature: _____		Requesting Physician & NPI
	Copy To:		
	Name _____		
	Address _____		
City _____ State _____ Zip _____			

PATIENT	Name (Last, First) _____ MI _____ MRN _____ DOB _____
	Address _____ City _____ State _____ Zip _____
	Home # () _____ Work # () _____ <input type="checkbox"/> Male <input type="checkbox"/> Female Race: <input type="checkbox"/> Black <input type="checkbox"/> White <input type="checkbox"/> Hispanic <input type="checkbox"/> Other: _____

BILLING	Bill: <input type="checkbox"/> Medicare <input type="checkbox"/> Medicaid <input type="checkbox"/> Insurance <input type="checkbox"/> Patient <input type="checkbox"/> Ordering Physician <input type="checkbox"/> Facility (Account) Authorization # _____
	Policy/ID # _____ Group # _____ 2nd Insurance Policy/ID # _____ Group # _____
	Insurance Carrier _____ Insured's Name _____ (if not patient check one - <input type="checkbox"/> spouse <input type="checkbox"/> child <input type="checkbox"/> other)
	Claim Address _____ 2nd Insurance Carrier _____
	City _____ State _____ Zip _____ Phone # _____ Claim Address _____ Insured's DOB _____

Patient Status Hospital Inpatient Hospital Outpatient Hospital Non-Patient Insured's DOB _____
 ▲ **Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Service (Highest Specificity Required)** Billing Information Attached

HISTOLOGY = Gross & Microscopic Exam	ICD-CM▲
	Collection Date _____ Collection Time _____
	Specimen Type _____
	Number of Vials Submitted _____ (UroScore® requires a sextant (6+ vials) biopsy & a PSA Value)
	<input type="checkbox"/> Prostate Histology
	<input type="checkbox"/> Prostate Histology w/UroScore®
	Prostate Histology, if Gleason 6 or 7 (3+4), Reflex to: <input type="checkbox"/> PTEN IHC <input type="checkbox"/> PTEN/ERG IHC
	<input type="checkbox"/> Prostate Histology, Reflex to ConfirmMDx@ on negative/HGPIN
	<input type="checkbox"/> Bladder Histology Biopsy
	<input type="checkbox"/> Bladder Histology TUR
<input type="checkbox"/> Vas Deferens (Sterilization) Histology	
<input type="checkbox"/> Other Histology: _____	
<input type="checkbox"/> Consultation: _____	
PSA Date _____ PSA _____ ng/mL	
DRE Finding <input type="checkbox"/> Suspicious <input type="checkbox"/> Normal	
<input type="checkbox"/> Isolated Nodule <input type="checkbox"/> Multiple Nodules	
Previous Biopsy <input type="checkbox"/> Positive <input type="checkbox"/> Negative	
<input type="checkbox"/> PIN <input type="checkbox"/> Suspicious	
Therapy <input type="checkbox"/> Cryosurgery <input type="checkbox"/> Chemotherapy	
<input type="checkbox"/> Hormone Therapy <input type="checkbox"/> Radiation Therapy	

MicrocytePLUS® URINE CYTOLOGY	ICD-CM▲
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	Clinical Data <input type="checkbox"/> Hematuria
	<input type="checkbox"/> TCC, Current <input type="checkbox"/> TCC, History Dx Date: _____
	<input type="checkbox"/> Other: _____
	Specimen Type (Required)
	<input type="checkbox"/> Voided Urine (Bladder) <input type="checkbox"/> Catheterized Urine
	<input type="checkbox"/> Post-Cysto Void <input type="checkbox"/> Bladder Wash
	<input type="checkbox"/> Ileal Conduit/NeoBladder <input type="checkbox"/> Urethral Wash
	<input type="checkbox"/> Renal Wash - Left <input type="checkbox"/> Renal Wash - Right
<input type="checkbox"/> Ureter Wash - Left <input type="checkbox"/> Ureter Wash - Right	
<input type="checkbox"/> Other _____	
INDIVIDUAL TESTS: (May be ordered or added to profile)	
<input type="checkbox"/> VU6 Pap Stain (only) Urine Cytology ‡	
<input type="checkbox"/> FNA (Fine Needle Aspiration) Site: _____	
<input type="checkbox"/> K600D Bladder Cancer FISH (Pathologist Review) ‡	
<input type="checkbox"/> β2 Microglobulin ✦ <input type="checkbox"/> Microalbumin ✦ <input type="checkbox"/> Total Protein ✦	
MicrocytePLUS® URINE CYTOLOGY PROFILES	
<input type="checkbox"/> 994 Hematuria Profile ✦ Cytodiagnostic Urinalysis Correlating Cytology (by concentration technique, with Pap and Feulgen stain), Urine Dipstick, Microalbumin, β2 Microglobulin, Total Protein. 994 only on void, catheterized, or post-cysto void; other collection methods processed as U03 cytology.	
<input type="checkbox"/> VU3 Cytology Plus Monitoring Profile ‡ Cytology (Pap and Feulgen stain)	
<input type="checkbox"/> VU1D Bladder Cancer FISH/Cytology Pathodiagnostic Profile ‡ Bladder Cancer FISH Assay and Cytology (Pap and Feulgen Stain) Including integrated cytomolecular diagnostic interpretation with clinical correlation by pathologist (MD)	
<input type="checkbox"/> VU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic Profile ‡ Cytology (Pap and Feulgen stain); reflex to Bladder Cancer FISH (Pathologist review) on atypical cytology results	
‡ Bladder Cancer FISH/Urine Cytology Kit (Alcohol Fixative) ✦ Urine Cytopathology Kit (Tablet Preservative) See reverse for collection method requirements and CPT codes	

CHEMISTRY	ICD-CM▲
	Specimen Type
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	Fasting? Yes <input type="checkbox"/> No <input type="checkbox"/> Frozen <input type="checkbox"/>
	S = Serum U = 24 Hr. Urine
	Endocrinology
	<input type="checkbox"/> Total PSA@% <input type="checkbox"/> Alkaline Phosphatase
	<input type="checkbox"/> Total PSA@%/Rflx Free PSA with free/total PSA ratio <input type="checkbox"/> Albumin (S)
	<input type="checkbox"/> Total PSA@% and Free PSA with free/total PSA ratio <input type="checkbox"/> ALT
	<input type="checkbox"/> Testosterone <input type="checkbox"/> Ammonia (U)
<input type="checkbox"/> Unbound Testosterone <input type="checkbox"/> AST	
<input type="checkbox"/> Testosterone/Unbound <input type="checkbox"/> BUN	
<input type="checkbox"/> Testosterone with % Free <input type="checkbox"/> Calcium (S, U)	
<input type="checkbox"/> FSH <input type="checkbox"/> Chloride (S, U)	
<input type="checkbox"/> LH <input type="checkbox"/> Cholesterol@%	
<input type="checkbox"/> Prolactin <input type="checkbox"/> Citrate (U)	
<input type="checkbox"/> AFP@ <input type="checkbox"/> CO2	
<input type="checkbox"/> Beta HCG@% <input type="checkbox"/> Creatinine (S, U)	
<input type="checkbox"/> TSH@% <input type="checkbox"/> Cystine (U)*	
<input type="checkbox"/> Direct Bilirubin	
<input type="checkbox"/> Glucose@%	
<input type="checkbox"/> HDL@%	
<input type="checkbox"/> Magnesium (S, U)	
<input type="checkbox"/> Oxalate (U)	
<input type="checkbox"/> pH (U)	
<input type="checkbox"/> Phosphorus (S, U)	
<input type="checkbox"/> Potassium (S, U)	
<input type="checkbox"/> PTH	
<input type="checkbox"/> Sodium (S, U)	
<input type="checkbox"/> Renal Function Panel	
<input type="checkbox"/> Sulfate (U)	
<input type="checkbox"/> Comp. Metabolic Panel	
<input type="checkbox"/> Total Bilirubin	
<input type="checkbox"/> Total Protein (S, U)	
<input type="checkbox"/> Triglyceride@%	
<input type="checkbox"/> Uric Acid (S, U)	
<input type="checkbox"/> Other: _____	
<input type="checkbox"/> Dianon performed venipuncture & PST Initials _____	

24 HR URINE	ICD-CM▲
	REQUIRED
	Total Volume _____ mls Specimen Type _____
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	24 Hr Urine Chemistry Profiles (Dianon 24hr Urine Kit REQUIRED)
	Check profile below or individual tests available in Chemistry section.
	<input type="checkbox"/> UroStone®24 Uric Acid (Uric Acid/Creatinine/Sulfate)
	<input type="checkbox"/> Creatinine Clearance (Serum Creatinine/Urine Creatinine) requires serum & urine specimens and
	Patient Height: _____ Inches & Weight: _____ lbs.
	<input type="checkbox"/> UroStone®24 Cystine* (Creatinine/Qualitative Cystine*)
<input type="checkbox"/> UroStone®24 Calcium (Creatinine/Calcium/Sodium/pH)	
<input type="checkbox"/> UroStone®24 Citrate (Citrate/Creatinine)	
<input type="checkbox"/> UroStone®24* (Calcium/Citrate/Creatinine/Magnesium/Oxalate/pH/Phosphorus/Qualitative Cystine*/Sodium/Uric Acid)	
<input type="checkbox"/> UroStone®Max24* (Ammonia/Calcium/Chloride/Citrate/Creatinine/Magnesium/Oxalate/pH/Phosphorus/Potassium/Sodium/Sulfate/Uric Acid/Qualitative Cystine*)	

STONES	ICD-CM▲
	Specimen Type
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	<input type="checkbox"/> Stone Analysis - Urinary Tract Calculus (Stone Analysis Kit)
	<input type="checkbox"/> Spontaneously Passed
	<input type="checkbox"/> Lithotripsy
	<input type="checkbox"/> Surgically Removed

HISTORY	<input type="checkbox"/> Other: _____
	<input type="checkbox"/> Dianon performed venipuncture & PST Initials _____
	Indicate previous Urinary Tract/Systemic Disorders, Biopsy or Therapy Results, and current Medications:

* Quantitative Cystine performed on positive Qualitative Cystine at additional charge.
 When ordering tests for which Medicare or Medicaid reimbursement will be sought, physicians should only order tests that are medically necessary for the diagnosis or treatment of the patient.
 † Separately billable stains may be added by pathologist when medically necessary to render a diagnosis.
 (260N) Rev 11/10/2021
 Dianon Systems, Inc. is a subsidiary of Laboratory Corporation of America Holdings, using the brand Labcorp.

Refer to Determining Necessity of ABN Completion on reverse.

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

PLEASE ENSURE REQUESTING PHYSICIAN IS INDICATED AND THE TEST REQUESTED IS MARKED.

Specimen Label Instructions . . .

1. Complete the requisition with all requested information.
2. Remove the required number of labels from the front of this sheet.
3. Place one (1) label on each specimen container (not on lid).

Any Questions?
Please Call Client Services at 1-800-411-1839

SITE, IF APPLICABLE
 Dianon Pathology

SITE, IF APPLICABLE
 Dianon Pathology

L TRANSITION ZONE
 Dianon Pathology

R TRANSITION ZONE
 Dianon Pathology

LEFT
 Dianon Pathology

RIGHT
 Dianon Pathology

L LAT BASE
 Dianon Pathology

L LAT MID
 Dianon Pathology

L LAT APEX
 Dianon Pathology

R LAT BASE
 Dianon Pathology

R LAT MID
 Dianon Pathology

R LAT APEX
 Dianon Pathology

LEFT BASE
 Dianon Pathology

LEFT MID
 Dianon Pathology

LEFT APEX
 Dianon Pathology

RIGHT BASE
 Dianon Pathology

RIGHT MID
 Dianon Pathology

RIGHT APEX
 Dianon Pathology

2A
2B
2C
Item# 0050659 Form # 260N.32

2A
2B
2C

PHYSICIAN	Physician/Authorized Signature: _____		Requesting Physician & NPI _____
	Copy To: _____		
	Name _____		
	Address _____		
City _____ State _____ Zip _____			

PATIENT	Name (Last, First) _____ MI _____ MRN _____ DOB _____
	Address _____ City _____ State _____ Zip _____
	Home # (_____) _____ Work # (_____) _____ <input type="checkbox"/> Male <input type="checkbox"/> Female Race: <input type="checkbox"/> Black <input type="checkbox"/> White <input type="checkbox"/> Hispanic <input type="checkbox"/> Other: _____

BILLING	Bill: <input type="checkbox"/> Medicare <input type="checkbox"/> Medicaid <input type="checkbox"/> Insurance <input type="checkbox"/> Patient <input type="checkbox"/> Ordering Physician <input type="checkbox"/> Facility (Account) Authorization # _____
	Policy/ID # _____ Group # _____ 2nd Insurance Policy/ID # _____ Group # _____
	Insurance Carrier _____ Attach secondary billing info. Insured's Name _____
	Claim Address _____ (if not patient check one - <input type="checkbox"/> spouse <input type="checkbox"/> child <input type="checkbox"/> other) 2nd Insurance Carrier _____
	City _____ State _____ Zip _____ Phone # _____ Claim Address _____
	Patient Status <input type="checkbox"/> Hospital Inpatient <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Hospital Non-Patient Insured's DOB _____

▲ Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Service (Highest Specificity Required) _____ Billing Information Attached

HISTOLOGY = Gross & Microscopic Exam	ICD-CM[▲] _____
	Collection Date _____ Collection Time _____
	Specimen Type _____
	Number of Vials Submitted _____ (UroScore [®] requires a sextant (6+ vials) biopsy & a PSA Value)
	<input type="checkbox"/> Prostate Histology
	<input type="checkbox"/> Prostate Histology w/UroScore [®]
	Prostate Histology, if Gleason 6 or 7 (3+4), Reflex to: <input type="checkbox"/> PTEN IHC <input type="checkbox"/> PTEN/ERG IHC
	<input type="checkbox"/> Prostate Histology, Reflex to ConfirmMDx@ on negative/HGPIN
	<input type="checkbox"/> Bladder Histology Biopsy
	<input type="checkbox"/> Bladder Histology TUR

MicrocytePLUS[®] URINE CYTOLOGY	ICD-CM[▲] _____
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	Clinical Data <input type="checkbox"/> Hematuria
	<input type="checkbox"/> TCC, Current <input type="checkbox"/> TCC, History Dx Date: _____
	<input type="checkbox"/> Other: _____
	Specimen Type (Required)
	<input type="checkbox"/> Voided Urine (Bladder) <input type="checkbox"/> Catheterized Urine
	<input type="checkbox"/> Post-Cysto Void <input type="checkbox"/> Bladder Wash
	<input type="checkbox"/> Ileal Conduit/NeoBladder <input type="checkbox"/> Urethral Wash
	<input type="checkbox"/> Renal Wash - Left <input type="checkbox"/> Renal Wash - Right

CHEMISTRY	ICD-CM[▲] _____
	Specimen Type
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	Fasting? Yes <input type="checkbox"/> No <input type="checkbox"/> Frozen <input type="checkbox"/>
	S = Serum U = 24 Hr. Urine
	Endocrinology
	<input type="checkbox"/> Total PSA@% <input type="checkbox"/> Alkaline Phosphatase
	<input type="checkbox"/> Total PSA@%/Rflx Free PSA with free/total PSA ratio <input type="checkbox"/> Albumin (S)
	<input type="checkbox"/> Total PSA@% and Free PSA with free/total PSA ratio <input type="checkbox"/> ALT
	<input type="checkbox"/> Testosterone <input type="checkbox"/> Ammonia (U)

24 HR URINE	ICD-CM[▲] _____
	REQUIRED
	Total Volume _____ mls Specimen Type _____
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	24 Hr Urine Chemistry Profiles (Dianon 24hr Urine Kit REQUIRED) Check profile below or individual tests available in Chemistry section.
	<input type="checkbox"/> UroStone [®] 24 Uric Acid (Uric Acid/Creatinine/Sulfate)
	<input type="checkbox"/> Creatinine Clearance (Serum Creatinine/Urine Creatinine) <i>requires serum & urine specimens and</i>
	Patient Height: _____ Inches & Weight: _____ lbs.
	<input type="checkbox"/> UroStone [®] 24 Cystine★ (Creatinine/Qualitative Cystine*)
	<input type="checkbox"/> UroStone [®] 24 Calcium (Creatinine/Calcium/Sodium/pH)

STONES	ICD-CM[▲] _____
	Specimen Type
	Collection Date _____ Collection Time _____ AM <input type="checkbox"/> PM <input type="checkbox"/>
	<input type="checkbox"/> Stone Analysis - Urinary Tract Calculus (Stone Analysis Kit)
	<input type="checkbox"/> Spontaneously Passed
	<input type="checkbox"/> Lithotripsy
	<input type="checkbox"/> Surgically Removed
	<input type="checkbox"/> VU3 Cytology Plus Monitoring Profile ‡ Cytology (Pap and Feulgen stain)
	<input type="checkbox"/> VU1D Bladder Cancer FISH/Cytology Pathodiagnostic Profile ‡ Bladder Cancer FISH Assay and Cytology (Pap and Feulgen Stain) Including integrated cytological diagnostic interpretation with clinical correlation by pathologist (MD)
	<input type="checkbox"/> VU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic Profile ‡ Cytology (Pap and Feulgen stain); reflex to Aladder Cancer FISH (Pathologist review) on atypical cytology results

HISTORY	Individual Serum/ 24 Hr. Urine Chemistry
	<input type="checkbox"/> Calcium (S, U)
	<input type="checkbox"/> Chloride (S, U)
	<input type="checkbox"/> Cholesterol@%
	<input type="checkbox"/> Citrate (U)
	<input type="checkbox"/> CO2
	<input type="checkbox"/> Creatinine (S, U)
	<input type="checkbox"/> Cystine (U)★
	<input type="checkbox"/> Direct Bilirubin
	<input type="checkbox"/> Glucose@%
<input type="checkbox"/> HDL@%	
<input type="checkbox"/> Magnesium (S, U)	
<input type="checkbox"/> Oxalate (U)	
<input type="checkbox"/> pH (U)	
<input type="checkbox"/> Phosphorus (S, U)	
<input type="checkbox"/> Potassium (S, U)	
<input type="checkbox"/> PTH	
<input type="checkbox"/> Sodium (S, U)	
<input type="checkbox"/> Sulfate (U)	
<input type="checkbox"/> Total Bilirubin	
<input type="checkbox"/> Total Protein (S, U)	
<input type="checkbox"/> Triglyceride@%	
<input type="checkbox"/> Uric Acid (S, U)	
<input type="checkbox"/> Other: _____	
<input type="checkbox"/> Dianon performed venipuncture & PST Initials _____	
Indicate previous Urinary Tract/Systemic Disorders, Biopsy or Therapy Results, and current Medications:	

* Quantitative Cystine performed on positive Qualitative Cystine at additional charge.
 When ordering tests for which Medicare or Medicaid reimbursement will be sought, physicians should only order tests that are medically necessary for the diagnosis or treatment of the patient.
 † Separately billable stains may be added by pathologist when medically necessary to render a diagnosis.

Refer to Determining Necessity of ABN Completion on reverse.
Symbols Legend
 @ = Subject to Medicare medical necessity guidelines.
 % = Subject to Medicare frequency guidelines.
 # = Medicare deems investigational. Medicare does not pay for services it deems investigational.

PLEASE ENSURE REQUESTING PHYSICIAN IS INDICATED AND THE TEST REQUESTED IS MARKED.

Specimen Label Instructions . . .

1. Complete the requisition with all requested information.
2. Remove the required number of labels from the front of this sheet.
3. Place one (1) label on each specimen container (not on lid).

Any Questions?
Please Call Client Services
at 1-800-411-1839



Medical Necessity

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 does not meet the medical necessity requirements set forth by the Medicare carrier or the test(s) is/are being performed more frequently than the carrier allows, an ABN should be completed.
- *An ABN should be completed for all tests that are considered research or investigational by Medicare.

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:

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 local/national medical review as of 10/01/2021

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

TUBE AND SPECIMEN TRANSPORTATION REQUIREMENTS

TEST	TUBE	CPT	SPECIMEN	TEST	TUBE	CPT	SPECIMEN	TEST	TUBE	CPT	SPECIMEN
AFP	(SST)	82105	(S,R)	Comprehensive Metabolic Panel	(SST)	80053	(S,R)	Prolactin	(SST)	84146	(S,R)
Albumin	(SST)	82040	(S,R)	Creatinine	(SST)	82565	(S,R)	PSA	(SST)	84153	(S,R)
ALT	(SST)	84460	(S,R)	Creatinine Clearance	(Urine+SST)	82575	(U,S,R)	PSA, Free	(SST)	84154	(S,R)
Alkaline Phosphatase	(SST)	84075	(S,R)	Direct Bilirubin	(SST)	82248	(S,R)	PTH ♦	(SST)	83970	(S,R)
AST	(SST)	84450	(S,R)	Electrolyte Panel	(SST)	80051	(S,R)	Renal Function Panel	(SST)	80069	(S,R)
Basic Metabolic Panel	(SST)	80048	(S,R)	FSH	(SST)	83001	(S,R)	Sodium	(SST)	84295	(S,R)
Beta HCG	(SST)	84702	(S,R)	Glucose	(SST)	82947	(S,R)	Testosterone	(SST)	84403	(S,R)
BUN	(SST)	84520	(S,R)	Hepatic Function Panel	(SST)	80076	(S,R)	Total Bilirubin	(SST)	82247	(S,R)
Calcium	(SST)	82310	(S,R)	HDL	(SST)	83718	(S,R)	Total Protein	(SST)	84155	(S,R)
CBC with Plt	(LT)	85027	(WB,R)	LH	(SST)	83002	(S,R)	Triglycerides	(SST)	84478	(S,R)
CBC with Plt & Diff	(LT)	85025	(WB,R)	Lipid Panel	(SST)	80061	(S,R)	TSH	(SST)	84443	(S,R)
Chloride	(SST)	82435	(S,R)	Magnesium	(SST)	83735	(S,R)	Unbound Testosterone	(SST)	84402	(S,R)
Cholesterol	(SST)	82465	(S,R)	Phosphorus	(SST)	84100	(S,R)	Uric Acid	(SST)	84550	(S,R)
CO ₂	(SST)	82374	(S,R)	Potassium	(SST)	84132	(S,R)				

TUBE REQUIREMENTS: SST-Serum Separator Tube LT-Lavender Top

SPECIMEN REQUIREMENTS: F-Frozen S-Serum R-Refrigerate U-Urine WB-Whole Blood
♦ Must be processed within 48 hours of collection if not received frozen

MicroCytePLUS®/Urine Cytology

Test	Urine Collection Method	CPT
994 Hematuria Profile-Urine Cytology	Voided, Catheterized, Post-Cysto Void	88108, 88313, 81003, 82232, 82043, 84156
K600D Bladder Cancer FISH Pathodiagnostic	Voided, Catheterized, Post-Cysto Void, Wash (Bladder, Renal, Ureter)	88120
VU1D Bladder Cancer FISH/Cytology Pathodiagnostic Profile	Voided, Catheterized, Post-Cysto Void, Wash (Bladder, Renal, Ureter)	88112, 88120
VU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic	Voided, Catheterized, Post-Cysto Void, Wash (Bladder, Renal, Ureter)	88112, if reflexed, 88120
VU3 Cytology Plus Monitoring Profile	Voided, Catheterized, Post-Cysto Void, Wash (Bladder, Renal, Ureter), Ileal Conduit/Neobladder	88112
VU6 Cytology Pap Stain Only	Voided, Catheterized, Post-Cysto Void, Wash (Bladder, Renal, Ureter), Ileal Conduit/Neobladder	88112

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 – Bladder Cancer FISH to 7 days, Cytology or Hematuria 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 Bladder Cancer 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.
Bladder Cancer FISH will not be performed on Ileal Conduit/Neobladder urine specimens.

EXPLANATION OF REFLEX TESTING

Reflex Free PSA Testing

Free PSA will be performed and billed if "Reflex Free PSA" is requested and the total PSA results fall within the requesting physician's previously defined parameters. The default range is 2-10 ng/ml.

Quantitative Cystine

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

Specimen Collection Information

- *Avoid submitting tissue specimens on fibrous materials such as gauze.
- *After tissue has been obtained, place biopsy into 10% Formalin immediately. Do Not allow to air dry.
- *All Urine Cytologies, FNA's and Fluid Aspirates must be submitted in the cytology alcohol fixative provided.
- *Hematuria Profile Specimens must be collected in a preservative tablet kit.
- *All 24 Hour Urine Specimens must be collected in a Dianon 24 Hour Urine Specimen Collection Kit (orange collection container) and submitted in the two vials provided with the kit.

HELPFUL HINTS

24 Hour Urine Specimens - Do Not Collect First Morning Void

Ordering Kits

Kit Orders may be placed through our Client Relations Department at 800-411-1839.

Please do not return unused histology vials or fixative bottles to our lab. Please dispose of unused histology vials in accordance with local laws and regulations regarding formalin disposal.

DESCRIPTION OF PRIMARY LAB TESTING

AMA PANELS

Electrolyte Panel 80051 - Sodium, Potassium, Chloride, Carbon Dioxide

Lipid Panel 80061@% - Chol@%, HDL@%, LDL (calculated)@%, Triglycerides@%

Hepatic Function Panel 80076 - Total Protein, Albumin, Total Bilirubin, Direct Bilirubin, Alkaline Phosphatase, SGOT (AST), SGPT (ALT)

Basic Metabolic Panel 80048 - Calcium, CO₂ (Carbon Dioxide), Chloride, Creatinine, Glucose@%, Potassium, Sodium, Urea Nitrogen (BUN)

Renal Function Panel 80069 - Albumin, Calcium, CO₂ (Carbon Dioxide), Chloride, Creatinine, Glucose@%, Phosphorus Serum, Potassium, Sodium, Urea Nitrogen (BUN)

Comprehensive Metabolic Panel 80053 - SGPT (ALT), Albumin, Total Bilirubin, Calcium, Chloride, Creatinine, Glucose@%, Alkaline Phosphatase, Potassium, Total Protein, Sodium, SGOT (AST), Urea Nitrogen (BUN), CO₂ (Carbon Dioxide)

PROFILES

Hematology 85027@ / 85025@ - CBC with PLT@, CBC with PLT and Diff@

24 Hr Urine CPT Codes

Ammonia 82140, Calcium 82340, Chloride 82436, Citrate 82507, Creatinine 82570, Magnesium 83735, Oxalate 83945, pH 83986, Phosphorus 84105, Potassium 84133, Qualitative Cystine 82127, Quantitative Cystine 82131, Sodium 84300, Sulfate 84392, Total Protein 84156, Uric Acid 84560

ConfirmMDx test performed and billed by MDxHealth® at Irvine, CA.

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 of procedures is desired.

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 combinations/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 here 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 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.