esting abcor	Dianon Systems,Inc 840 Research Parkway Oklahoma City, OK 73104 Phone: (800) 411-1839 or (405) 290-4000 Fax: (800) 211-0442 or (405) 290-4046	0			art #_ st Re		ition Date Color Volume	
HYSICIAN	Physician/Authorized Signature: Copy To: Name							
	Address							
	City State Zip		Requesting Phys		MDN			0 <mark>0B</mark>
	Name (Last, First)Address			IVII			State	7in
PAT							White □ Hispanic □ Oth	er.
	Bill: ☐ Medicare ☐ Medicaid ☐ Insurance ☐ Patient	☐ Ordering Ph	nysician 🗆 Fac	ility (Account) Authoriza	ation #			
ILLING	Insurance Carrier Claim Address			Insured's Name (if not patient check one - □ 2nd Insurance Carrier	l spouse 🗆	child 🗆		
m	City State ZipPI Patient Status □ Hospital Inpatient □ Hospital Outpatient □ Hospital Patient			Claim Address Insured's DOB				
	▲ Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Servi							ling Information Attache
Exam	ICD-CM ⁴ Collection Date Collection Time	ICD-CM ⁴ Collection		Collection Time	AM □		CD-CM ^A	
oscopic	Specimen Type				PM 🗆		ollection Date	Collection Time AM D
Micro	Number of Vials Submitted (UroScore® requires a sextant (6+ vials) biopsy & a PSA Value)		ta □ Hematuria rrent □ TCC, His	tory Dx Date:			Fasting? Yes □ No □	PM C
= Gross &	☐ Prostate Histology ☐ Prostate Histology w/UroScore® Prostate Histology, if Gleason 6 or 7 (3+4), Reflex to: ☐ PTEN IHC ☐ PTEN/ERG IHC	Specimen 1	Type (REQUIRED) Jrine (Bladder)				S = Serum U	= 24 Hr. Urine
შ,	☐ Prostate Histology, Reflex to ConfirmMDx®@ on Non-Cancer Prostate Histology, Reflex to Genomic Prostate Score®@ on Gleason:	☐ Post-Cys	sto Void	☐ Bladder Wash		Ei	ndocrinology	Individual Serum/ 24 Hr. Urine Chemistry
OLO	\square All \square 3+3 or 3+4 \square 4+3 or higher (excluding GG5)	☐ Renal W		☐ Urethral Wash ☐ Renal Wash - Right			∃ Total PSA@% ∃ Total PSA@%/ Rflx Free PS	☐ Alkaline Phosphatase SA ☐ Albumin (S)
	Patient has Life Expectancy of ≥ 10 years? ☐ Yes ☐ No ☐ Bladder Histology Biopsy ☐ Bladder Histology TUR		/ash - Left	☐ Ureter Wash - Right			with free/total PSA ratio I Total PSA@% and Free PS	☐ ALT A ☐ Ammonia (U)
HIS	☐ Vas Deferens (Sterilization) Histology	INDIVIDUA		ordered or added to profile)		> -	with free/total PSA ratio Testosterone	☐ AST ☐ BUN
	□ Other Histology: Consultation:	- / '	Stain (only) Urine e Needle Aspiratio	n) Site:		FIC	☐ Unbound Testosterone ☐ Testosterone/Unbound	☐ Calcium (S, U) ☐ Chloride (S, U)
	Consultation: PSA Date PSA Date PSA ng/mL PSA Ng/mL PSA Date Normal (T1c) ABNL. Bilateral (T2c)	□ K600D B □ β2 Micro		SH (Pathologist Review) ‡ croalbumin * 🗖 Total Pro			Testosterone with % Free I FSH	☐ Cholesterol@% ☐ Citrate (U)
	☐ Suspicious ☐ ABNL, Unilat ≤ 50% lobe (T2a)	· ·		OLOGY PROFILES		浜 ロ	□ LH □ Prolactin	☐ CO2 ☐ Creatinine (S, U)
	Multi Nodules ABNL, Unilat > 50% lobe (T2b)		with Pap and Feulgen lobulin, Total Protein.	ating Cytology (by concentration stain), Urine Dipstick, Microalbu 994 only on void, catheterized, o methods processed as VU3 cyto	min, r] AFP@] Beta HCG@%] TSH@%	☐ Cystine (U) ★ ☐ Direct Bilirubin ☐ Glucose@% ☐ HDL@%
	ICD-CM ^A	□ VU3 Cyte	ology Plus Monit		gy.	Pa	anels	☐ Magnesium (S, U) ☐ Oxalate (U)
	Total Specimen Volume mls Type	□ VU1D Bla	ology (Pap stain) adder Cancer FISH	Cytology Pathodiagnostic F	Profile ‡		∃ Electrolyte Panel ∃ Lipid Panel@%	□ pH (U) □ Phosphorus (S, U)
	Collection Date Collection Time AM PM	integrated by patholo	ancer FISH ASSAY and cytomolecular diagno: igist (MD)	Cytology (Pap Stain) Including stic interpretation with clinical con	relation		Hepatic Function Panel Basic Metabolic Panel	□ Potassium (S, Ú) ´ □ PTH
Щ	24 Hr Urine Chemistry Profiles (Dianon 24hr Urine Kit REQUIRED) Select Profile below or individual tests available in Chemistry section.	Cytology (F	Pap stain); reflex to Blad	eflex/Cytology Pathodiagnostic der Cancer FISH (Pathologist	Profile ‡		Renal Function Panel Comp. Metabolic Panel	☐ Sodium (S, U) ☐ Sulfate (U)
URIN	☐ UroStone® 24 Uric Acid (Uric Acid/Creatinine/Sulfate) ☐ UroStone® 24 Cystine★ (Creatinine/Qualitative Cystine★)	‡ Bladder Can	atypical cytology results cer FISH/Urine Cytolog	y Kit (Alcohol Fixative)		Pa	anel components on back	☐ Total Bilirubin☐ Total Protein (S, U)
Œ	□ UroStone® 24 Calcium (Creatinine/Calcium/Sodium/pH)	Vurine Cytopa See reverse for	athology Kit (Tablet Pre r collection method req	servative) uirements and CPT codes				☐ Triglyceride@% ☐ Uric Acid (S, U)
24 H	☐ UroStone® 24 Citrate (Citrate/Creatinine) ☐ Creatinine Clearance (Serum Creatinine/Urine Creatinine)	ICD-CM4					1 Other:	
	requires serum & urine specimens <u>and</u> Patient Height: Inches & Weight: Ibs.			Collection Time	AM 🗆			
	☐ UroStone® 24★ (Calcium/Citrate/Creatinine/Magnesium/	1			РМ 🗆	► Ind	1 Labcorp performed venipunct licate previous Urinary Tract/Syst sults, and current Medications:	ture & PST Initialstemic Disorders, Biopsy or Therap
	☐ UroStone® Max24★ (Ammonia/Calcium/Chloride/Citrate/Creatinine/ Magnesium/Oxalate/pH/Phosphorus/Potassium/Sodium/Sulfate Uric Acid/Qualitative Cystine★)	Urinary	nalysis - Tract Calculus nalysis Kit)	☐ Spontaneously Pass☐ Lithotripsy☐ Surgically Removed	- 1	HISTOF	our our out moundiblis.	
/hen o	ntitative Cystine performed on positive Qualitative Cystine at additional charge. ordering tests for which Medicare or Medicaid reimbursement will be sought, physicians sh arately billable stains may be added by pathologist when medically necessar	hould only order tes ry to render a di	sts that are medically nagnosis.	ecessary for the diagnosis or treat Dianon Systems,	ment of the pa Inc. is a sub	atient. sidiary of	Laboratory Corporation of America	(260N) Rev 11/29/20. Holdings, using the brand Labcorp.
efer '	to Determining Necessity of ABN Completion on reverse.			_				-
<u> </u>	ols Legend Subject to Medicare medical necessity guidelines. SITE, IF APP	PLICABLE					ianon	LEFT BASE Dianon
! = I	Subject to Medicare frequency guidelines. Medicare deems investigational. Medicare does not			+		System	ns, Inc T MID	Systems, Inc
	pay for services it deems investigational. PLEASE ENSURE REQUESTING PHYSICIAN IS SITE, IF APP						ianon	Dianon Systems, Inc
	DICATED AND THE TEST REQUESTED IS MARKED.					LLAT		LEFT APEX
	Specimen Label ZONE Dianon Sys						ianon	Dianon Systems, Inc
	. Complete the requisition with all RTRANS	SITION				RLATE	BASE ianon	RIGHT BASE Dianon
2	Remove the required number of Dianon Sys	stems, Inc				System		Systems, Inc
	labels from the front of this sheet.					R LA	TMID	RIGHT MID
3	B. Place one (1) label on each specimen container (not on lid).						ianon	Dianon Systems, Inc

Dianon Systems, Inc

Please Call Client Services

at 1-800-411-1839

RIGHT APEX Dianon Systems, Inc

R LAT APEX Dianon Systems, Inc

======================================	Labco	IDCOF g will be performed at a prp laboratory, including rly branded Dianon Path
Porm #	Z	Physician/Autho
2B 659	CIA	Сору То:

Dianon Systems,Inc 840 Research Parkway Oklahoma City, OK 73104 **Phone:** (800) 411-1839 or (405) 290-4000 **Fax:** (800) 211-0442 or (405) 290-4046

Chart # Test Requisition	LAB USE	Date Color Volume	> Y
1			

0111101	ny branded Dianon Pathology.			100111	- 4 -				
Z	Physician/Authorized Signature:								
SICIA	Сору То:								
¥	Name								
直	Address								
	City State Zip		Requesting Physic	cian & NPI					
늘	Name (Last, First)			MI MRN		D	OB		
	Address		City			State	_Zip		
٦	Home # ()Work # (_)		☐ Male ☐ Female Race: ☐	Blac	k □ White □ Hispanic □ Oth	er:		
	Bill: ☐ Medicare ☐ Medicaid ☐ Insurance ☐ Patient		l Ordering Physician ☐ Faci	ility (Account) Authorization #_					
	Policy/ID #	_ (Group #	2nd Insurance Policy/ID #	\mathbb{Z}	Gro	oup #		
5	Insurance Carrier		Attach secondary billing into. Insured's Name						
E	Claim Address			(if not patient check one - □ spouse I 2nd Insurance Carrier	⊐ child	□ other)			
層	City State Zip			Claim Address					
	Patient Status □ Hospital Inpatient □ Hospital Outpatient □			Insured's DOB					
	▲ Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Se						ing Information Attached		
E	ICD-CM⁴		ICD-CM▲			ICD-CM ^A			
c Exam	Collection Date Collection Time		Collection Date	Collection Time AM C	j	Specimen Type			
copi	Specimen Type			PM C]	Collection Date	Collection Time AM		
cros	Number of Vials Submitted		Clinical Data				PM □		
Σ. Mi	(UroScore® requires a sextant (6+ vials) biopsy & a PSA Value) □ Prostate Histology □ Prostate Histology w/UroScore®		☐ TCC, Current ☐ TCC, Hist	ory Dx Date:	-	Fasting? Yes □ No □	Frozen □		
Gross &	Prostate Histology, if Gleason 6 or 7 (3+4), Reflex to:		Specimen Type (REQUIRED)		=	S = Serum U	= 24 Hr. Urine		
Ę.	□ PTEN IHC □ PTEN/ERG IHC	_	□ Voided Urine (Bladder)	☐ Catheterized Urine			Individual Commo		
75	= 1 restate rineresgy, rising, to committee @ cirrien cancer	ָם	□ Post-Cysto Void	☐ Bladder Wash		Endocrinology	Individual Serum/ 24 Hr. Urine Chemistry		
ပြု	Prostate Histology, Reflex to Genomic Prostate Score® @ on Gleason:		☐ Ileal Conduit/NeoBladder	☐ Urethral Wash		☐ Total PSA@%	☐ Alkaline Phosphatase		
6	□ All □ 3+3 or 3+4 □ 4+3 or higher (excluding GG5) Patient has Life Expectancy of ≥ 10 years? □ Yes □ No	20	☐ Renal Wash - Left	☐ Renal Wash - Right		☐ Total PSA@%/ Rflx Free PS	A □ Albumin (S)		
ΙË	☐ Bladder Histology Biopsy ☐ Bladder Histology TUR	☐ Ureter Wash - Left ☐ Ureter Wash - Right ☐ Other				with free/total PSA ratio ☐ Total PSA@% and Free PS.	☐ ALT A ☐ Ammonia (II)		
I≌	☐ Vas Deferens (Sterilization) Histology	Е				with free/total PSA ratio	□ AST		
1	□ Other Histology:	Z	INDIVIDUAL TESTS: (May be o UV6 Pap Stain (only) Urine		≿	☐ Testosterone☐ Unbound Testosterone	☐ BUN ☐ Calcium (S, U)		
	□ Consultation:	☐ FNA (Fine Needle Aspiration) Site:			STE	☐ Testosterone/Unbound			
	PSA DatePSA) (®/(adder Cancer FISH (Pathologist Review) ‡ lobulin ∻ □ Microalbumin ∻ □ Total Protein ∻		Testosterone with % Free ☐ FSH	☐ Cholesterol@%		
	DRE Finding □ Normal (T1c) □ ABNL, Bilateral (T2c)						☐ Citrate (U) ☐ CO2		
	☐ Suspicious ☐ ABNL, Unilat ≤ 50% lobe (T2a)	ᆸ	•	rtePLUS® URINE CYTOLOGY PROFILES Hematuria Profile ÷ iagnostic Urinalysis Correlating Cytology (by concentration que, with Pap and Feulgen stain), Urine Dipstick, Microalbumin, croglobulin, Total Protein. 994 only on void, catheterized, or		☐ Prolactin	☐ Creatinine (S, U)		
	☐ Multi Nodules ☐ ABNL, Unilat > 50% lobe (T2b) Previous Bx ☐ Positive ☐ Negative ☐ PIN ☐ Suspicious	/te				☐ AFP@ ☐ Beta HCG@%	☐ Cystine (U)★ ☐ Direct Bilirubin		
	Therapy ☐ Chemo ☐ Cryo ☐ Hormone ☐ Radiation	crocytel	β2 Microglobulin, Total Protein. 9			☐ TSH@%	☐ Glucose@%		
		cr		methods processed as VU3 cytology.			☐ HDL@% ☐ Magnesium (S, U)		
	ICD-CMA REQUIRED	Σ	Urine Cytology (Pap stain)	orning Prome +		Panels	☐ Oxalate (U)		
	Total Specimen Volume mls Type	☐ VU1D Bladder Cancer FISH/Cytology Pathodiagnostic Profile ‡ Bladder Cancer FISH Assay and Cytology (Pap Stain) Including integrated cytomolecular diagnostic interpretation with clinical correlation				☐ Electrolyte Panel☐ Lipid Panel@%	□ pH (U) □ Phosphorus (S, U)		
	Collection Date Collection Time AM PM					☐ Hepatic Function Panel	□ Potassium (S, U)		
	24 Hr Urine Chemistry Profiles (Dianon 24hr Urine Kit REQUIRED)		by pathologist (MD)	flov/Cutology Pathodiagnostic Profile +		☐ Basic Metabolic Panel☐ Renal Function Panel	□ PTH □ Sodium (S, U)		
URINE	Select Profile below or individual tests available in Chemistry section.		□ WU4D Bladder Cancer FISH Reflex/Cytology Pathodiagnostic Profile ‡ Cytology (Pap stain); reflex to Bladder Cancer FISH (Pathologist review) on atypical cytology results ‡ Bladder Cancer FISH/Urine Cytology Kit (Alcohol Fixative)			☐ Comp. Metabolic Panel	☐ Sulfate (Ù)		
2	☐ UroStone® 24 Uric Acid (Uric Acid/Creatinine/Sulfate)					Panel components on back	☐ Total Bilirubin ☐ Total Protein (S, U)		
2	☐ UroStone® 24 Cystine★ (Creatinine/Qualitative Cystine⋆) ☐ UroStone® 24 Calcium (Creatinine/Calcium/Sodium/pH)		 Urine Cytopathology Kit (Tablet Pres See reverse for collection method requ 	servative)			☐ Triglyceride@%		
품	☐ UroStone® 24 Citrate (Citrate/Creatinine)		Oct for the composition method requ	montonico una or i ocuco	1	☐ Uric Acid (S, U)			
24	☐ Creatinine Clearance (Serum Creatinine)		ICD-CM▲	M▲		□ Other:			
1	requires serum & urine specimens <u>and</u>	S							
	Patient Height: Inches & Weight: Ibs.	z	Collection Date	Collection Time AM		□ Labcorp performed venipuncture & PST Initials			
	2 of octorio 2 rx (calcium) of auto/or out in into/magnicolarity	10		PM □		Indicate previous Urinary Tract/Syst			
	Oxalate/pH/Phosphorus/Qualitative Cystine*/Sodium/Uric Acid) UroStone® Max24* (Ammonia/Calcium/Chloride/Citrate/Creatinine/	S	☐ Stone Analysis -	☐ Spontaneously Passed		Results, and current Medications:			
	Magnesium/Oxalate/pH/Phosphorus/Potassium/Sodium/Sulfate		Urinary Tract Calculus (Stone Analysis Kit)	☐ Lithotripsy ☐ Surgically Removed					
* Qua	Uric Acid/Qualitative Cystine*) antitative Cystine performed on positive Qualitative Cystine at additional charge.		, , , , , , , , , , , , , , , , , , , ,	,	أتنار				
When	ordering tests for which Medicare or Medicaid reimbursement will be sought, physicians parately billable stains may be added by pathologist when medically necess			ecessary for the diagnosis or treatment of the Dianon Systems, Inc. is a si	patient ubsidiar	PHYSICIAN'S COPY y of Laboratory Corporation of America I	(260N) Rev 11/29/2023 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.
- 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 that 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.

- Be executed on the CMS approved ABN form (CMS-R-131)
 Identify the Medicare Part B Beneficiary, using the name as it appears on the patient's red, white and blue Medicare card Indicate the test(s)/procedure(s) which may be denied within the relevant reason column Include an estimated cost for the test(s)/procedure(s) subject to the ABN Have 'Option 1', 'Option 2', or 'Option 3' designated by the beneficiary

 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 10/01/2023

- @ = 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 Pane	(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								
<u>Test</u>	<u>Urine Collection Method</u>	<u>CPT</u>						
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 Hematuria Profile.

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

HELPFUL HINTS

24 Hour Urine Specimens - Do Not Collect First Morning Void

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

<u>Lipid Panel</u> 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 and GPS testing performed and billed by MDxHealth® at Irvine, CA ConfirmMDx® and Genomic Prostate Score® are registered trademarks of MDxHealth SA.

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