labcorp

Testing will be performed at a Labcorp laboratory, including formerly branded Diagon Pathology

Dianon Systems, Inc. 1 Forest Parkway Shelton, CT 06484 800-328-2666 203-926-7100

ENDOCRINE PATHOLOGY

formerly branded Dianon Pathology.				
ACCOUNT INFORMATION		PATIENT INFORM	MATION	
ACCOUNT NO. TELEPHONE NO.				
Acceptance.				
		CHART NUM	IBER	PATIENT D.O.B.
ACCOUNT NAME AND ADDRESS				
		PATIENT LAST NAME	FIRST NAME	M.I.
		PATIENT LAST NAME	FIRST NAME	M.I.
		CTDEET ADDRESS		
		STREET ADDRESS		
REQUESTING PHYSICIAN (PLEASE PRINT) PHYSICIA	AN/AUTHORIZED SIGNATURE	CITY		STATE ZIP CODE
REQUESTING THISICIAN (LEDGE TRAIN)	NAME THORIZED SIGNATURE	SEX M F		() -
REQUESTING PHYSICIAN NPI REFERRIN	NG PHYSICIAN	RACE:	MRN / PATIENT ID#	PATIENT TELEPHONE NO.
BILLING INFORMATION				
Diagnosis/Signs/	Symptoms in ICD-CM format in effect at Da	ate of Service (Highest Specificity	Required) REQUIRED	
			ICD-CM CODE	(S):
BILL: ☐ PRACTICE/FACILITY ☐ PATIENT ☐ MEDICARE				
POLICY/ID#	GROUP #	2 ND INS POLICY/ID# _		GROUP #
INSURANCE CARRIER		INSURANCE CARRIER		
CLAIM ADDRESS		CLAIM ADDRESS		
CITY STATE				
PATIENT HOSPITAL STATUS \square INPATIENT \square OL	TPATIENT NON-PATIENT		HIP TO INSURED: SPOU	INSURED'S DOB
CUNICAL DATA		FATIENT 3 KELATIONSE	III TO INSURED: LA SPOL	OSE IN CHIED IN OTHER
CLINICAL DATA				
Collection Date:		Mark site(s) collected for	fine needle aspirate based on l	location on the thyroid illustration:
Fixative: Cytolyt® / 95% EtOH Other:		_	A	A
RNARetain® Reflex Testing Vial included				
MYARetain Renex resting via included		☐ Right Lobe		☐ Left Lobe
		#	A Park Trans	#
Additional Clinical Data:			CV 30-65	"
			10.00	
		_	1	
			-	
		_	☐ Isthmu	IS
			11	
		_	#	
CYTOLOGY SPECIMEN #1 (use separate the	yroid FNA kit for each specimen)	CYTOLOGY SPEC	CIMEN #2 (use separate thy	roid FNA kit for each specimen)
☐ FNA Site:	Number of Slides:	_ ☐ FNA Site:		Number of Slides:
☐ FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate		☐ FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate		
☐ FNA, reflex to ThyGeNEXT®* if FNA results a	FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate, reflex to			
ThyraMIR®v2 if mutation is negative or not f	ThyraMIR®v2 if mutation is negative or not fully indicative of malignancy			
(ThyGeNEXT [®] includes markers for BRAF, HRAS, KRAS, NRAS, PIK3CA#, ALK, GNAS, RET, TERT, PTEN, NTRK, PPARgamma, THADA, and PAX8; ThyraMIR [®] includes miRNA markers)@		(ThyGeNEXT® includes markers for BRAF, HRAS, KRAS, NRAS, PIK3CA#, ALK, GNAS, RET, TERT, PTEN, NTRK, PPARgamma, THADA, and PAX8; ThyraMIR® includes miRNA markers)@		
*Molecular testing requires sample in RNARetain® vial	Training includes mixtor markers/		ires sample in RNARetain® vial	THYTANIK INCIDES THIN IV Harkers,
CYTOLOGY SPECIMEN #3 (use separate the	vroid ENA kit for each specimen)	CYTOLOGY SPEC	IMEN #4 (use separate thy	roid FNA kit for each specimen)
☐ FNA Site:	•			Number of Slides:
FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate		FNA, reflex to ThyGeNEXT [®] * if FNA results are indeterminate		
☐ FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate, reflex to		FNA, reflex to ThyGeNEXT®* if FNA results are indeterminate, reflex to ThyraMIR®v2 if mutation is negative or not fully indicative of malignancy		
ThyraMIR [®] v2 if mutation is negative or not f	ully indicative of malignancy	ThyraMIR [®] v2 if m	utation is negative or not	fully indicative of malignancy
(ThyGeNEXT $^{\otimes}$ includes markers for BRAF, HRAS, KRAS, N				NRAS, PIK3CA#, ALK, GNAS, RET,
TERT, PTEN, NTRK, PPARgamma, THADA, and PAX8; Th	ıyraMIR® includes miRNA markers)@			ΓhyraMIR [®] includes miRNA markers)@
*Molecular testing requires sample in RNARetain® vial	or in the probability	Molecular testing requ	iires sample in RNARetain [®] vial	
Molecular thyroid testing performed by Interpace E	Jiagnostics, LLC, Pittsburgh, PA			
ADDITIONAL TESTS				
© 2024 Laboratory Corporation of America® Holdings	When and mine and the last	Modicaro or Modicaria	C. 4-1 -®	is a registered trademark of Cytyc Corporation
, .	will be sought, physicians shou	Medicare or Medicaid reimbursem Ild order only those tests that are		is a registered trademark of Cytyc Corporation. ain® is a registered trademark of Asuragen, Inc.
CT Lic. #: CL-0356	, , , ,	gnosis or treatment of the patient.		EXT® and ThyraMIR® are registered service
1193 REV. 11/05/2024	WHITE COPY TO DIANON	N PINK COPY TO PHYSICIAN	√ marks of	Interpace Diagnostics, LLC.
	_			
tefer to Determining Necessity of ABN completion on reverse.	Name: N	lame:	Name:	Name:
ymbols Legend				
e = Subject to Medicare medical necessity guidelines	Coll. Date: C	Coll. Date:	Coll. Date:	Coll. Date:
 Medicare deems investigational. Medicare does not pay for services it deems investigational 	Site: S	Site:	Site:	Site:
SPECIMEN LABEL	Sito S		JII.O	
INSTRUCTIONS:				
1.) Complete the requisition with all	+			
requested information.	Name: N	lame:	Name:	Name:
2.) Remove the required number of labels from the front of this sheet.				
3.) Place one (1) label on each specimen	Coll. Date:	Coll. Date:	Coll. Date:	Coll. Date:
container (not on the lid).	Site: S	Site:	Site:	Site:
PLEASE DISPOSE OF UNUSED LABELS.				

Test Combination/Panel Policy

Labcorp's policy is to provide physicians, in each instance, with the flexibility to choose appropriate tests to assure that the convenience of ordering test combinations/panels does not distance physicians who wish to order a test combination/profile 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 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 applicable 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.

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 the 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 Medicare or the test(s) is being performed more frequently than Medicare allows, an ABN should be completed.

*An ABN should be completed for all tests that are considered investigational (experimental or for research use) 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 <u>and</u> dated by the beneficiary or his/her representative <u>prior to</u> the service being rendered.

Symbols used to designate Medicare medical review as of 10/01/2024

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

ThyGeNEXT®
CPT Code 0245U
ThyraMIR®v2
CPT Code 0018U

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