

KIT (Mol.)PDGFRa (Mol.)

Single Biomarkers & Consults Solid Tumor Pathology Requisition

Client Information		Patient Information			
Required Information		Last Name [.]		Ale □ Female	
Account #: Account Name:		First Name:	□ N M.I Other Pt II	D/Acct #:	
Street Address:			/ yyyy Medical R		
City, ST, ZIP: Client represents it has obtained informed consent from patient to perform the services described herein.					
Phone:Fax:		Specimen Information	n		
Additional Reporting Fax:					
Requisition Completed by: Date: Ordering Physician: NPI #:		Fixative/Preservative:	Specimen ID: Block ID: Fixative/Preservative:		
(nlease print: Last First):		Collection Date: mm / c	d/ yyyyCollection T	ïme: □ AM □ PM	
[please print Last, First]:		Retrieved Date: mm / c			
The undersigned certifies that he/she is licensed to order the test(s) listed bel	ow and that such test(s) are medi-	Body Site:	/ dd / yyyy		
cally necessary for the care/treatment of this patient. Authorized Signature: Date:		Primary Metastasis – If N	letastasis, list Primary:		
			Peripheral Blood: Green Top(s) Purple Top(s) Other Fresh Tissue (Media Type required):		
Billing Information		□ Fluid: CSF Pleu	□ Fluid: CSF Pleural Other		
Required: Please include face sheet and front/back of card for both primary and secondary insurance.		FNA cell block:			
Patient Status (Must Choose 1): 🗖 Hospital Patient (in) 🗖 Hospital Patien	Smears: Air Dried	Fixed Stained (type of stained	ain)		
Bill to: 🗆 Client Bill 🔹 Insurance 🔤 Medicare 🔤 Medicaid 🔤 Patient/Self-Pay		Silues # Unsta Paraffin Block(s) #:	□ Paraffin Block(s) #:		
□ Split Billing - Client (TC) and Insurance (PC) □ OP Molecular to MCR, all other testing to Client					
Bill charges to other Hospital/Facility:		Choose best block (for global molecular/NGS testing only) Submit 44 blocks. Blocks will be combined for molecular testing when necessary.			
Prior Authorization # See NeoGenomi	Perform IHC testing on all blocks, unless otherwise noted. For all other testing, specify which block to use for each if sending multiple blocks. See back for details.				
		Predictive Marker Fixation (C	AP/ASCO Requirement):		
Clinical Information		*Indicated markers/profiles/panels require fixation information Cold ischemic duration (mins): □ ≤ 1 hour □ Unknown			
Required: Please attach patient's pathology report (required), clinical hist ICD-10 (Diagnosis) Code/Narrative (Required):	ory, and other applicable report(s).	Fixative: 10% NBF Other		. 🗖 Unknown	
Reason for Referral:		Fixation duration (hours):		. 🗖 6-72 hours 🗖 Unknown	
New Diagnosis Relapse In Remission Mon	G - Global G-IA - Global with Imag	G - Global G-IA - Global with Image Analysis T - Tech-Only/Stain-Only T-IA - Tech-Only with Image Analysis			
Staging: 0 0 0 0 0 A 0 B 0 V Note: T-SOnt - Tech-Only with Semi-Quantitative interpretation by client				- Icen only with intege Analysis	
Reflex options are available with global test orders only. Tech-only clients mus	t use the test add-on process.	T-Qual - Tech-Only with Qualitative	interpretation by client		
Consultation		Head and Neck Cancer			
Consultation A NeoGenomics pathologist will select medically necessary tests (with any exception noted below by the client) to provide comprehensive analysis and professional Colon Cancer & Lynch Syndrome MMR IHC G-IA T-IA T-Qual		SQnt 🔲 T-Qual	G T G □ □ p16 (IHC) □	HPV DNA (Mol.)	
interpretation for the materials submitted.	Reflex to BRAF (Mol.) if MLH1 IHC is not expressed				
Surgical Pathology Consult (FFPE only) Add NeoTYPE® Profile if indicated			Melanoma G T G		
Differential Diagnosis:	Reflex to MMR (IHC) if MSI is high	Microsatellite Instability (MSI) Non-tumor tissue required. Reflex to MMR (IHC) if MSI is high		NeoSITE® Melanoma FISH T KIT (Mol.)	
			IHC is not expressed Molar Pregnancy □ Molar Preg. Comprehensive Consultation (includes p57 IHC □ p57 (IHC, tech-only)		
Bladder Cancer G T	Reflex to BRAF (Mol.) if MLH1 IHC is not expressed				
Bladder Cancer (FISH, urine only)					
Brain Cancer	BRAF (Mol.) Reflex to MLH1 Promoter Methylation if BRAF neg. KRAS (Mol.) NRAS (Mol.)			Ploidy FISH for Molar Preg.	
GT GT			All PD-L1 IHC G T***		
□ 1p/19q Deletion (FISH) □ N/A IDH1/IDH2 (Mol.) □ N/A ATRX (Mol.) □ *Ki67 (IHC)*	MLH1 Promoter Methylation		PD-L1 22C3 FDA (KEYTRUDA®) [‡]		
$N/A \square ATRX (IHC)$	HER2 (Except Breast)		E ESCC (Esophageal)		
N/A 🗖 Beta Catenin (IHC) Methylation (Mol.)	G T		HNSCC (Head & Neck)		
BRAF (FISH)	BRAF V600E (IHC)* (FISH) CDKN2A/B (p16) Deletion for N/A p53 (IHC) Mesothelioma or Glioma (FISH) PTEN (FISH) PTEN (FISH) PTEN (FISH)		TNBC (Breast) PD-L1 22C3 FDA for NSCLC [‡] PD-L1 28-8 FDA for NSCLC (HNSCC, Urothelial Carcinoma) [‡] PD-L1 22C3 FDA (KEYTRUDA [®]) for Gastric/GEA [‡] PD-L1 22C3 FDA (KEYTRUDA [®]) for Gastric/GEJ/EAC [‡] PD-L1 28-8 (OPDIVO [®]) for Gastric/GEJ/EAC [‡]		
Mesothelioma or Glioma (FISH) 🗖 🗖 PTEN (FISH)					
Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests. Control of the performed w/o image analysis unless client requests.		🗖 🗖 PD-L1 SP142 FDA TECENTRIQ® f			
		PD-L1 SP263 FDA for NSCLC [#] D PD-L1 LDT [#]			
			***Ordering Pathologist listed has received to perform the professional interpretation f	d the required competency training	
Breast Cancer G-IA T-IA T	**For global HER2 IHC with result 2+, NeoGenomics will add global HER2 FISH unless marked here:		Prostate Cancer	G T	
□* □ ER/PgR/HER2***			Androgen Receptor (Mol.)	D D PTEN (FISH)	
□* □ ER/PgR/HER2**/Ki67* Lung Cancer □ □ Individual Stains: ER* PgR* HER2*** Ki67* G T			Sarcoma FISH		
Reflex to HER2 FISH G G T if global HER2 IHC is:	□ □ ALK, D5F3 IHC (lung, FDA) [‡]			GT	
	ALK Lung (FISH)*:			2 🗖 🗖 PDGFB Rearr [‡]	
* Reflex to global PD-L1 22C3 FDA (KEYTRUDA®) for TNBC if global ER/PgR/HER2 panel is negative	 Reflex to ROS1 (FISH) if global / N/A BRAF (Mol.) 	ALK IS NEGATIVE 🗖 G 🗂 T			
**For global HER2 IHC with result 2+, NeoGenomics will add global HER2 FISH unless marked here: Do not reflex 2+		r Mesothelioma or Glioma	Thyroid Cancer BRAF (Mol.)	G T □ □ RET (FISH) [‡]	
G T	N/A Early-stage NSCLC Panel		NRAS (Mol.)		
C HER2 (FISH)* Reflex to HER2 IHC G G-IA T-IA T if global HER2 FISH N/A EGER (Mol incl			KRAS (Mol.)		
result is Group 2, 3, or 4 (see back)	 N/A EGFR (Mol., includes T790M) N/A KRAS (includes G12C mutation) 		Other /Pan-Cancer Testing G T		
 For global HER2 FISH: Send path report. If HER2 IHC has been interpreted elsewhere: Send IHC report and also send HER2 IHC slide if result is 2+. 	□ INVA KNAS (Includes 6126 Inutatio	FGFR2 Rearr. FISH			
N/A Breast NGS Fusion Panel (Global Only)	N/A MET Exon 14 Deletion (Mol.)		□ □ FOLR1 (IHC) [#] □ □ NTRK 1,2,3 FISH [#]		
	RET (FISH)*			Reflex to NTRK NGS Fusion Panel	
GI Cancer G	□ □ ROS1 (FISH) [‡] □ □ ROS1 (IHC) [‡]		If IHC is expressed/equivocal: • Re	flex to NTRK 1, 2, 3 FISH 🗖 G 🗖 T	
-			D D Other		

D Other_

FlexREPORT[®] Please add summary report.

Specimen Requirements

Refrigerate specimen if not shipping immediately and use cool pack during transport. Please call Client Services Team with any questions regarding specimen requirements or shipping instructions at 866.776.5907 option 3. Please refer to the website for specific details on each specimen.

Additional Billing Information

Any organization referring specimens for testing services pursuant to this Requisition Form ("Client") expressly agrees to the following terms and conditions.

- 1. Binding Service Order. This Requisition Form is a contractually binding order for the services ordered hereunder ("Services") and Client agrees that it is financially responsible for all tests billable to Client hereunder.
- 2. Third Party Billing by NeoGenomics and Right to Bill Client. Client agrees to accurately indicate on the front of the Requisition Form that either Client should be billed (e.g., Client receives reimbursement pursuant to a non-fee-for-service basis, including, but not limited to, a capitated, diagnostic related group ("DRG"), per diem, all-inclusive, or other such bundled or consolidated billing arrangement) or NeoGenomics should bill the applicable federal, state or commercial health insurer or other third party payer (collectively, "Payers") for all Services ordered pursuant to this Requisition Form. For all such Services billable to Payers, Client agrees to provide all billing information necessary for NeoGenomics to bill such payer. In the event NeoGenomics: (i) does not receive the billing information required for it to bill any Payers within ten days of the date that any Services are reported by NeoGenomics; (ii) the Services were performed for patients who have no Payer coverage arrangements; or (iii) the Payer identified by Client denies financial responsibility for the Services and indicates that Client is financially responsible, NeoGenomics shall have the right to bill such Services to Client.

Additional Specimen Information

If submitting multiple blocks, clients must indicate either "Choose best block (global molecular/NGS testing only)", "Perform IHC testing on all blocks", or assign the selection of blocks to individual tests. If multiple blocks are sent without a selection, they will be held until clarification is provided. Please call Client Services Team with any questions regarding specimen information.

Test Descriptions

Please see complete test descriptions and all available tests at our website, www.neogenomics.com/test-menu.

Test Notations

Specimen Usage

NeoGenomics makes every effort to preserve and not exhaust tissue, but in small and thin specimens, there is a possibility of exhausting the specimen in order to ensure adequate material and reliable results.

Breast HER2, ER, PgR (IHC) and Breast HER2 (FISH)

Breast specimens undergoing any of these tests should be invasive breast cancer or the invasive component of the breast cancer fixed in 10% neutral buffered formalin for at least 6 hours and no longer than 72 hours.

For global breast HER2 FISH cases, NeoGenomics will (if requested) reflex FISH to HER2 IHC if FISH results are consistent with CAP/ASCO 2018 result Groups 2, 3, or 4 for dualprobe ISH assays.

- Group 2: HER2/CEP17 ratio ≥ 2.0 and average HER2 copy number < 4.0 signals/cell
- Group 3: HER2/CEP17 ratio < 2.0 and average HER2 copy number ≥ 6.0 signals/cell
- Group 4: HER2/CEP17 ratio < 2.0 and average HER2 copy number ≥ 4.0 and < 6.0 signals/cell

If ordering global HER2 FISH after HER2 IHC was already interpreted outside NeoGenomics, please send the HER2 IHC result and the path report. If that IHC result was 2+, please submit the HER2-stained IHC slide to NeoGenomics with the FISH order so that we may correlate our analysis. This includes stain-only cases that were not scanned by NeoGenomics. If outside HER2 IHC results were other than 2+, we do not request the IHC slide but still request the HER2 IHC report.

FlexREPORT[®]

FlexREPORT can be ordered on any global or tech-only testing referred to NeoGenomics. This report template can be used to import data and images collected from testing performed outside of NeoGenomics, and incorporated into a one page summary report. Client logo and contact information will be in the header of the FlexREPORT.