

Title: Molecular Pathology Order Form (NCBH)		Published Date: 07/29/2025	
Division(s):	Area:	Last Review / Revised Date: 07/29/2025	
Entity: NC Baptist Hospital		Dept.: Molecular Pathology	

Name (Last)		(First)	(MI)	Date of Birth	Sex	EPIC MRN	Social Security Number	
Patient Address					City		State	Zip
Name of Insurance Carrier (please attach copy of card(s))			Subscriber Name (Last, First)			Subscriber ID		Group Number
Relationship to Patient	List ICD-10 Code(s):			Collection Date	Collection Time	Collected by		
<p>*Medicare will only pay for services that it determines to be reasonable and necessary under section (a)(1) of the Medicare Law. Tests are only covered under certain ICD10 diagnosis codes specified by Medicare. If ordered with a code not specified by Medicare, an Advance Beneficiary Notice indicating patient responsibility for payment must be obtained.</p>								
Authorizing Provider (Last, First)					Authorizing Provider Signature			
Point of Contact				Phone		Fax		Email
Specimen Type: <input type="checkbox"/> Amniotic Fluid <input type="checkbox"/> Bone Marrow <input type="checkbox"/> Extracted DNA From a CLIA-certified laboratory			<input type="checkbox"/> Peripheral Blood <input type="checkbox"/> FFPE <input type="checkbox"/> Fresh Tissue			Specimen ID		FFPE Number
FISH (Individual Tests)			FISH (Individual Tests) cont'd			Molecular Oncology		
<input type="checkbox"/> CDKN2C-CKS1B, BM, UPB <input type="checkbox"/> ABL2, BM, UPB <input type="checkbox"/> ALK, FFPE <input type="checkbox"/> LSI EVI1 (3q26.2), BM, UPB <input type="checkbox"/> BCL6(3Q27), BM, FFPE <input type="checkbox"/> +4, +10, +17, BM, UPB <input type="checkbox"/> del5q, BM, UPB <input type="checkbox"/> 5, 9, 15, BM <input type="checkbox"/> ROS1, FFPE <input type="checkbox"/> DEK-CAN, BM, UPB <input type="checkbox"/> del7q, BM, UPB <input type="checkbox"/> CEP 8, BM, UPB <input type="checkbox"/> T(8;21) Translocation (RUNX1T1/RUNX1), BM, UPB <input type="checkbox"/> MYC(8Q24), BM, FFPE <input type="checkbox"/> MYC CEP8 IGH, BM, FFPE <input type="checkbox"/> BCR/ABL1/ASS1, BM, UPB <input type="checkbox"/> ATM TP53, BM, UPB <input type="checkbox"/> ABL1, BM, UPB <input type="checkbox"/> MLL (KMT2A), BM, UPB <input type="checkbox"/> FLI1 EWSR1, FFPE <input type="checkbox"/> NUP98, BM <input type="checkbox"/> MDM2 (12q14)/CEP12, FFPE <input type="checkbox"/> ETV6, BM, UPB <input type="checkbox"/> T(12;21) Translocation (ETV6/RUNX1), BM, UPB <input type="checkbox"/> DDIT3, FFPE <input type="checkbox"/> Rhabdomyosarcoma, FFPE <input type="checkbox"/> TCL1, BM, UPB <input type="checkbox"/> Mantle Cell Lymphoma, BM, UPB <input type="checkbox"/> IGH(14Q), BM, UPB <input type="checkbox"/> IGH MAF, BM, UPB <input type="checkbox"/> IGH MAFB, BM, UPB <input type="checkbox"/> Follicular Lymphoma, BM, UPB <input type="checkbox"/> PML/RARA, BM, UPB <input type="checkbox"/> CBFβ/MYH11, BM, UPB <input type="checkbox"/> BCL2, BM, UPB, FFPE <input type="checkbox"/> MALT1 (18Q), BM, UPB, FF			<input type="checkbox"/> D20S108, BM, UPB <input type="checkbox"/> ERG EWSR1, FFPE <input type="checkbox"/> Trisomy 21 Analysis, PB <input type="checkbox"/> EWSR1, FFPE <input type="checkbox"/> TUPLE1, PB <input type="checkbox"/> SRY, PB FISH Panels <input type="checkbox"/> Acute Myeloid Leukemia (AML) Panel, BM, UPB <input type="checkbox"/> AneuVysion Screen, PB, AF, CVS, POC <input type="checkbox"/> Acute Lymphocytic Leukemia (ALL) Panel, BM, UPB <input type="checkbox"/> Chronic Lymphocytic Leukemia (CLL) Panel, BM, UPB <input type="checkbox"/> Eosinophilia Panel, BM, UPB <input type="checkbox"/> Lymphoma Panel, BM, FFPE <input type="checkbox"/> Multiple Myeloma Panel, BM, UPB <input type="checkbox"/> Myelodysplastic Syndrome (MDS) Panel, BM, UPB Cytogenetics Chromosome Analysis <input type="checkbox"/> Chromosome Analysis, Amniotic Fluid <input type="checkbox"/> Chromosome Analysis, Blood <input type="checkbox"/> Chromosome Analysis, Bone Marrow <input type="checkbox"/> Chromosome Analysis, Chorionic Villi <input type="checkbox"/> Chromosome Analysis, Fresh tissue <input type="checkbox"/> Chromosome Analysis, UPB Molecular Genetics <input type="checkbox"/> CTFR Full Gene Sequencing, PB <input type="checkbox"/> Cystic Fibrosis Carrier Screen, PB <input type="checkbox"/> Fragile X Syndrome, PB <input type="checkbox"/> Spinal Muscular Atrophy SMN1/2, PB <input type="checkbox"/> Pre-Transplant Chimerism Donor, BM, PB <input type="checkbox"/> Pre-Transplant Chimerism Recipient, BM, PB <input type="checkbox"/> Post-Chimerism/Marker analysis, BM, PB <input type="checkbox"/> Post-Chimerism Panel (CD3, CD33), BM, PB <input type="checkbox"/> Maternal Cell Contamination Geno, PB, AF, CVS, POC			<input type="checkbox"/> B-Cell Clonality, IGH, PB, FFPE <input type="checkbox"/> BCR-ABL1 p190, PB, BM <input type="checkbox"/> BCR-ABL1 p210, PB only <input type="checkbox"/> BRAF Mutation Testing, FFPE <input type="checkbox"/> EGFR Mutation Testing, FFPE <input type="checkbox"/> Extended RAS/BRAF Mutation Testing (KRAS, NRAS, BRAF), FFPE <input type="checkbox"/> T-Cell Clonality, TCRG, PB, FFPE <input type="checkbox"/> NGS Myeloid Panel, BM, PB <input type="checkbox"/> NGS Solid Tumor Panel, FFPE <input type="checkbox"/> MYD88 p.L265P variant detection, PB, BM, FFPE Specimen Types Definitions BM (Bone Marrow) UPB (Unprocessed Peripheral Blood (Whole Blood)) FFPE (Formalin Fixed Paraffin Embedded) PB (Peripheral Blood) AF (Amniotic Fluid) CVS (Chorionic Villi Sampling) POC (Products of Conception)		
						Additional Comments For molecular tumor testing on FFPE, the tumor cells or area of interest must be marked on the H&E slide. AND percent tumor in the circled must be added to this form before testing can be performed. % tumor: _____ <input type="checkbox"/> Tumor Cells are circled on H&E <input type="checkbox"/> Percent tumor is included Bill to: <input type="checkbox"/> Client <input type="checkbox"/> Medicare <input type="checkbox"/> Patient		