



Regional Pathology Services

University of Nebraska Medical Center
981180 Nebraska Medical Center
Omaha NE 68198-1180
www.reglab.org

Toll Free 1-800-334-0459
Phone (402) 559-6420
FAX (402) 559-9497

MOLECULAR DIAGNOSTICS TEST REQUEST FORM

Lab Use Only

SHADED AREAS FOR PATIENT INFORMATION REQUIRED

PATIENT LAST NAME		FIRST NAME		MI
DOB	GENDER <input type="checkbox"/> MALE <input type="checkbox"/> FEMALE	PT. ID# / ADDITIONAL INFO		
SSN	BILL	<input type="checkbox"/> OFFICE/CLIENT <input type="checkbox"/> PATIENT/PATIENT INSURANCE		
ATTACH COPY OF FRONT AND BACK OF INSURANCE CARD AND ATTACH COPY OF FRONT OF DRIVERS LICENSE IF UNABLE TO OBTAIN COPY OF REQUIRED INFORMATION ALL FIELDS BELOW ARE REQUIRED				
GUARANTOR NAME/DOB (REQUIRED IF PATIENT IS A MINOR)				
ADDRESS		CITY	STATE	ZIP
PRIMARY INSURANCE <input type="checkbox"/> MEDICARE IN-PATIENT <input type="checkbox"/> MEDICARE OUT-PATIENT <input type="checkbox"/> MEDICAID <input type="checkbox"/> INSURANCE				
POLICY ID#		GROUP ID#		
INSURANCE COMPANY			PHONE NUMBER	
INSURANCE COMPANY ADDRESS		CITY	STATE	ZIP
EFFECTIVE DATE / /				
DIAGNOSIS / MEDICAL NECESSITY (ENTER ALL THAT APPLIES)				
ICD-9/10 #1	ICD-9/10 #2	ICD-9/10 #3		

Accession #:	_____
Date Rec'd:	___/___/___ # of Slides: _____
Collection Date	___/___/___ Collection Time _____ AM PM
PHYSICIAN PROVIDER: _____ (Indicate the Supervising Dr./P.A. or N. Pract.)	
SECONDARY / TERTIARY INS - ATTACH INFORMATION	

NOTICE: WHEN ORDERING TESTS FOR WHICH MEDICARE REIMBURSEMENT WILL BE SOUGHT, PHYSICIANS SHOULD ONLY ORDER TESTS THAT ARE MEDICALLY NECESSARY FOR THE DIAGNOSIS OR TREATMENT OF A PATIENT RATHER THAN FOR SCREENING PURPOSES. FOR MORE INFORMATION SEE reglab.org/billingcompliance/

ABN ATTACHED PRIOR AUTHORIZATION ATTACHED

MOLECULAR DIAGNOSTIC TESTING

SOURCE: _____	
Pharmacogenomics <input type="checkbox"/> Abacavir Sensitivity Genotyping (HLA B*5701)	FISH Testing - specify indication(s) or gene(s): Indication: _____
HLA Genotyping for Disease Association <input type="checkbox"/> Ankylosing Spondylitis (B27); HLA-B Genotype <input type="checkbox"/> Antiglomerular basement membrane disease (DR15); HLA-DR Genotyping <input type="checkbox"/> Autoimmune Thyroid Disease (DR3, DR5); HLA-DR Genotyping <input type="checkbox"/> Autoimmune Hepatitis (DR3, DR4); HLA-DR Genotyping <input type="checkbox"/> Behcet's disease (B51); HLA-B Genotyping <input type="checkbox"/> Bird Shot Retinopathy (A29); HLA-A Genotyping <input type="checkbox"/> Celiac Disease (HLA-DQA1*05, HLA-DQB1*02, and HLA-DQB1*03:02); HLA-DQ Genotyping <input type="checkbox"/> Graves Disease (DR3); HLA-DR Genotyping <input type="checkbox"/> Hashimoto's thyroiditis (DR3); HLA-DR Genotyping <input type="checkbox"/> HLA-B*27 Syndromes; HLA-B Genotyping <input type="checkbox"/> Idiopathic Inflammatory Myopathy; HLA-B, DR and DQ Genotyping <input type="checkbox"/> Myelodysplastic syndrome (MDS) (DR15); HLA-DR Genotyping <input type="checkbox"/> Narcolepsy (DQB1*06:02); HLA-DQ Genotyping <input type="checkbox"/> Pemphigus vulgaris; HLA-B and HLA-DR Genotyping <input type="checkbox"/> Rheumatoid Arthritis (DR4); HLA-DR Genotyping <input type="checkbox"/> Systemic Lupus Erythematosus (DRB1*15:01, DRB1*03:01); HLA-DR Genotyping	Gene: _____
HLA Other Testing <input type="checkbox"/> Platelet Support (HLA-A&B Antigen Level) <input type="checkbox"/> HLA Type for suspected TRALI <input type="checkbox"/> Red Blood Cell Genotyping	Infectious Disease Testing <input type="checkbox"/> Adenovirus DNA Detection, Qualitative <input type="checkbox"/> B Pertussis DNA Detection <input type="checkbox"/> BK Virus DNA Detection, Quantitation; Plasma or Urine <input type="checkbox"/> BK Virus DNA Detection, Tissue Qualitative <input type="checkbox"/> CMV DNA Detection, Qualitative <input type="checkbox"/> CMV DNA Detection, Quantitation, Plasma <input type="checkbox"/> EBV DNA Detection, Qualitative <input type="checkbox"/> EBV DNA Detection, Quantitation, Plasma <input type="checkbox"/> Enterovirus RNA Detection <input type="checkbox"/> HBV DNA Detection, Quantitative <input type="checkbox"/> HCV Genotyping <input type="checkbox"/> HCV RNA Detection, Quantitative <input type="checkbox"/> Herpes Virus Panel (HSV, CMV, EBV, VZV, HHV-6) <input type="checkbox"/> HHV-6 DNA Detection, Qualitative <input type="checkbox"/> HHV-8 DNA Detection, Plasma/Other source <input type="checkbox"/> HIV RNA Detection, Quantitation <input type="checkbox"/> HSV DNA Detection <input type="checkbox"/> JC Virus DNA Detection <input type="checkbox"/> Norovirus RNA Detection, stool <input type="checkbox"/> Parvovirus DNA Detection <input type="checkbox"/> VZV DNA Detection
Molecular Solid Tumor <input type="checkbox"/> BRAF c.1799T>A (p.V600k, p.V600R, p.V600E) mutation <input type="checkbox"/> EGFR mutation (29 common mutations) <input type="checkbox"/> KRAS mutation (codons 12, 13, 61) <input type="checkbox"/> Microsatellite Instability Analysis (MSI)	Molecular Hematology <input type="checkbox"/> BCL1 Translocation t(11;14) <input type="checkbox"/> BCL2 Translocation t(14;18) <input type="checkbox"/> BCR-ABL t(9;22) (p190, p210) Qualitative for Diagnosis of CML, ALL <input type="checkbox"/> BCR-ABL t(9;22) (p 210) Quantitative for CML Therapeutic Monitoring* <input type="checkbox"/> FLT3 Internal Tandem Duplication <input type="checkbox"/> IGH Gene Rearrangement by DNA Amplification <input type="checkbox"/> IGH Gene Rearrangement by Southern Analysis <input type="checkbox"/> JAK2 (p.V617F) Mutation, Qualitative <input type="checkbox"/> NPM1 Mutation <input type="checkbox"/> T Cell Receptor Beta Gene Rearrangement by Southern Analysis <input type="checkbox"/> T Cell Receptor Gamma Gene Rearrangement by DNA Amplification <input type="checkbox"/> MPL Mutation
Sarcoma RT-PCR <input type="checkbox"/> Ewing Sarcoma t(11;22) <input type="checkbox"/> Rhabdomyosarcoma, t(1 or 2;13) <input type="checkbox"/> Synovial Sarcoma t(X;18)	
Inherited Disease Testing <input type="checkbox"/> Alpha-1 Antitrypsin Mutation (Z and S Mutations) <input type="checkbox"/> Hemochromatosis Mutation (p.C282Y, p.H63D) <input type="checkbox"/> Factor II Mutation (Prothrombin g.20210G>A) <input type="checkbox"/> Factor V Mutation (Leiden p.R506Q) <input type="checkbox"/> Factor II/V Mutations (Prothrombin g.20210G>A/Leiden p.R506Q) <input type="checkbox"/> Factor II/V and MTHFR Mutations <input type="checkbox"/> MTHFR Mutations (c.677C>T, c.1298A>C) <input type="checkbox"/> Cystic Fibrosis 39 Mutation Panel <input type="checkbox"/> Fragile X Mutation (FMR 1)	
	*Include recent CBC and differential test results.
	Comment: _____

Testing Supplies

Regional Pathology Services furnishes specimen-collection supplies for use by clients that send tests to us.

Supplies are ordered online at reglab.org/customer-service/supply-orders/ testing supplies and log-on information may be obtained by calling client services

Toll Free 800-334-0459

Phone 402-559-6420

Courier Services

Regional Pathology Services offers an extensive courier network, which includes contracted land and air courier services. Contracted land specimen pickup is provided at no charge for specimens tested by Regional Pathology Services or our designated reference laboratories.

To inquire about scheduled stops and after hours courier service, call

Client Services Toll Free 800-334-0459

Phone 402-559-6420

If shipping specimens address to:

Regional Pathology Services

University Of Nebraska Medical Center

668 S 41ST ST MSB3500

OMAHA NE 68105-1180

Transport Instructions:

Specimen Handling/Shipping

To ensure the safety of personnel and the community, proper handling of specimens for shipment is mandatory. Specimens will be rejected if submitted improperly. The shipper is responsible for the proper packaging and shipping of all specimens. Shippers must be trained and certified by their employer to be able to prepare and ship packages containing diagnostic specimens, biological substances and infectious substances. Rules of the various agencies involved may differ, and may change regularly. Specimen must have at least two patient identifiers and be packaged in sealed plastic bags to prevent leakage or contamination. Place accompanying paperwork in the bag pocket, away from the specimen. Practice universal blood and body fluid precautions when handling specimens.

For tests that require scheduling or special arrangements prior to specimen collection, refer to the Online Test Menu at reglab.org for more information.

Questions?

Contact client services at 800-334-0459