

CGx Requisition Form

First Name		Last Name		Middle Initial	Clinic Name
Social Security #	Date of Birth	Sex <input type="checkbox"/> F <input type="checkbox"/> M	Ethnicity <input type="checkbox"/> African American <input type="checkbox"/> Asian <input type="checkbox"/> Ashkenazi Jewish <input type="checkbox"/> Caucasian <input type="checkbox"/> Hispanic <input type="checkbox"/> Other		
Address				City	State Zip Phone
INSURANCE: Please provide a legible copy of the front and back of the patient's insurance card. IF NO INSURANCE: <input type="checkbox"/> Self Pay <input type="checkbox"/> WC/Auto (Date of Injury) <input type="checkbox"/> Other					
Name of Insured		Relationship to Patient		Insurance Company/Provider	Member/ID Number Group Number
Collector Name (Print)		Date Collected		Time Collected Fasting <input type="checkbox"/> Yes <input type="checkbox"/> No	
Specimen Type <input type="checkbox"/> OCD-100 (Buccal)		Specimen Storage <input type="checkbox"/> Room Temperature <input type="checkbox"/> Refrigerated		Specimen Shipping <input type="checkbox"/> Room Temperature <input type="checkbox"/> Cooling/Ice Pack	

MOLECULAR DIAGNOSTICS TESTING OPTIONS

Cancer Genomics (CGx) Please select the Panel to be tested. Please attach patient Medication List.

<input type="checkbox"/> Comprehensive Cancer	<input type="checkbox"/> Nervous System/Brain Comprehensive
<input type="checkbox"/> Breast Cancer STAT (7-10 day TAT) ATM, BRCA1, BRCA2, CDH1, CHEK2, PALB2, PTEN, STK11, TP53	ALK, APC, ATM, DICER1, EPCAM, HRAS, LZTR1, MEN1, MLH1, MSH2, MSH6, NF1, NF2, PHOX2B, PMS2, PRKAR1A, PTCH1, PTEN, POT1, SMARCA4, SMARCB1, SMARCE1, SUFU, TP53, TSC1, TSC2, VHL
<input type="checkbox"/> Breast Comprehensive ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, MLH1, MRE11, MSH2, NBN, NF1, PALB2, PTEN, RAD50, RAD51C, RAD51D, STK11, TP53, XRCC2	<input type="checkbox"/> Ovarian Comprehensive BARD1, BRCA1, BRCA2, BRIP1, CDH1, EPCAM, MLH1, MRE11, MSH2, MSH6, NBN, PALB2, PMS2, RAD51C, RAD51D, SMARCA4, STK11, TP53
<input type="checkbox"/> Breast and Ovarian Comprehensive APC, BMRP1A, CDH1, CTNNA1, EPCAM, KIT, MLH1, MSH2, MSH6, NF1, PDGFRA, PMS2, SMAD4, STK11, TP53	<input type="checkbox"/> Pancreatic Comprehensive APC, ATM, BMRP1A, BRCA1, BRCA2, CDK4, CDKN2A, EPCAM, FANCC, MEN1, MLH1, MSH2, MSH6, NF1, PALB2, PMS2, SMAD4, STK11, TP53, TSC1, TSC2, VHL
<input type="checkbox"/> Endometrial Comprehensive BRCA1, BRCA2, EPCAM, MLH1, MSH2, MSH6, PMS2, POLD1, PTEN, TP53	<input type="checkbox"/> Paraganglioma-Pheochromocytoma Comprehensive FH, MAX, NF1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127, VHL
<input type="checkbox"/> Gastric Comprehensive APC, BMRP1A, CDH1, CTNNA1, EPCAM, KIT, MLH1, MSH2, MSH6, NF1, PDGFRA, PMS2, SMAD4, STK11, TP53	<input type="checkbox"/> Prostate Comprehensive ATM, BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PMS2, TP53
<input type="checkbox"/> Hematologic Malignancy Comprehensive ATM, BLM, CEBPA, EPCAM, GATA2, HRAS, MLH1, MSH2, MSH6, NBN, NF1, PMS2, RUNX1, TERC, TERT, TP53	<input type="checkbox"/> Renal/Urinary Comprehensive BAP1, CDC73, CDKN1C, DICER1, DIS3L2, EPCAM, FH, FLCN, GPC3, MET, MTF, MLH1, MSH2, MSH6, PMS2, PTEN, SDHA, SDHB, SDHC, SDHD, SMARCA4, SMARCB1, TP53, TSC1, TSC2, VHL, WT1
<input type="checkbox"/> Melanoma Comprehensive BAP1, BRCA2, CDK4, CDKN2A, CHEK2, MC1R, MITF, POT1, PTEN, RB1, SLC45A2, TP53, TYR	<input type="checkbox"/> Sarcoma Comprehensive APC, BLM, CDKN1C, DICER1, EPCAM, FH, HRAS, KIT, MLH1, MSH2, MSH6, NBN, NF1, PDGFRA, PMS2, PRKAR1A, PTCH1, RB1, RECQL4, SDHA, SDHB, SDHC, SDHD, SUFU, TP53, WRN
<input type="checkbox"/> Thyroid Comprehensive APC, CHEK2, DICER1, PRKAR1A, PTEN, RET, TP53	<input type="checkbox"/> Custom Select genes to create a custom panel.

CGx Personal/Family History Questionnaire Please complete Questionnaire

PATIENT'S PERSONAL HISTORY (Hx)

Cancer/Tumor	Personal Hx	Age at Dx
Breast <input type="checkbox"/> Triple negative (ER-, PR-, HER2-) <input type="checkbox"/> Multiple primaries	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Ovarian <input type="checkbox"/> Check if non-epithelial	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Prostate (Gleason score >7)	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Pancreatic	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Colon/Rectal	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Stomach	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Melanoma	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Other Cancer(s):		

OTHER PERSONAL INFORMATION

<input type="checkbox"/> Bone marrow transplant	<input type="checkbox"/> Previous genetic testing for hereditary cancer
	<input type="checkbox"/> Current diagnosis of a hematologic cancer

OTHER PERSONAL INFORMATION

Relationship	Maternal	Paternal	Cancer Site(s)	Age at Dx
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		

ICD-10 DIAGNOSIS CODES: Additional documentation supporting Medical Necessity may be attached.

<input type="checkbox"/> C50.019 Malignant neoplasm of nipple and areola, unspecified female breast	<input type="checkbox"/> C18.2 Malignant neoplasm of ascending colon
<input type="checkbox"/> Z85.43 Personal history of malignant neoplasm of ovary	<input type="checkbox"/> Z85.51 Personal history of malignant neoplasm of bladder
<input type="checkbox"/> C56.3 Malignant neoplasm of bilateral ovaries	<input type="checkbox"/> C73 Malignant neoplasm of thyroid gland
<input type="checkbox"/> C56.9 Malignant neoplasm of unspecified ovary	<input type="checkbox"/> Z85.850 Personal history of malignant neoplasm of thyroid
<input type="checkbox"/> C61 Malignant neoplasm of prostate	<input type="checkbox"/> C95.00 Leukemia not having achieved remission
<input type="checkbox"/> Z85.07 Personal history of malignant neoplasm of pancreas	<input type="checkbox"/> Z85.6 Personal history of leukemia
<input type="checkbox"/> Z85.841 Personal history of malignant neoplasm of brain	<input type="checkbox"/> C34.90 Malignant neoplasm of unspecified part of the bronchus or lung
<input type="checkbox"/> C71.0 Malignant neoplasm of cerebrum, except lobes and ventricle	<input type="checkbox"/> C80.1 Neoplasm, unspecified
<input type="checkbox"/> C54.1 Malignant neoplasm of endometrium	<input type="checkbox"/> C16.9 Malignant neoplasm of stomach, unspecified
	<input type="checkbox"/> C43.30 Malignant melanoma of unspecified part of face
	<input type="checkbox"/> Other _____

Medical Necessity Required for insurance I, the provider, attest that I am the ordering physician or am authorized under applicable laws and regulations to order genetic testing for the patient. I attest that this test is medically necessary for the diagnosis or detection of a disease or disorder, and that the results will be used in medical management and care decisions for the patient. I further attest that any information entered on this Test Requisition Form, or otherwise provided by me on behalf of the patient, is true and correct to the best of my knowledge, and that the patient has consented to receive communications about his/her genetic test from Mainstream Diagnostic Laboratory.

Patient Informed Consent Patient must consent I, the patient, voluntarily consent to the collection and testing of my sample. I certify that the specimen is fresh and has not been adulterated in any manner. I authorize the laboratory to release the results of this testing to the ordering provider. I further authorize my insurance benefits to be paid directly to Mainstream Diagnostic Laboratory for services rendered. I acknowledge that the lab may be treated as an out-of-network provider. In the event I receive payment for laboratory services from my insurer, I will remit said payment to the lab within 14 days of receipt. I will either endorse the original check, or produce a personal check for the entire payment amount, and forward it to the lab. When selecting Self Pay above, I acknowledge financial responsibility for all lab charges associated with the processing of this test requisition. All rights to the samples will belong to the laboratory conducting the testing. There will be no compensation in the event of an invention resulting from research and development using this sample. I agree to allow my provided samples to be used for the purpose of (diagnosis/research) (development/quality control). I understand that if I agree, any information identifying me will be kept confidential so that it will not be possible to determine from whom the sample was drawn. Your signature on this form indicates that you understand to your satisfaction the information about Mainstream Diagnostic Laboratory and agree to have the test done. In no way does this waive your legal rights or release anyone from their legal and professional responsibilities. If you have further questions concerning matters related to this consent, you may wish to seek professional genetic counseling prior to signing this form. Consultation with a medical geneticist, genetic counselor, or your referring healthcare provider also may be warranted after the test has been completed.

Opt In for Research I give permission for my sample and clinical information to be used in de-identified studies at Mainstream Diagnostic Laboratory and for publication, if Mainstream Diagnostic Laboratory deems it appropriate. I understand that my name and/or other identifying information will NOT be used in or linked to the results of any studies and publications. More information is available at www.mainstreamlab.com.

Provider Name (Print)	Provider NPI #	Clinic Address	Clinic Phone/Fax
Provider Signature	Date	Patient Signature (or Legal Guardian)	Date
X		X	