Cancer Test Requisition Form (Comprehensive)- Page 1 of 3

Ambry Genetics
A Konica Minolta Company
Aliso Viejo, CA 92656 USA
Toll Free: 866.262.7943 | Fax: 949.900.5501 | ambrygen.com

COMPLETE ENTIRE FORM AND SUBMIT CLINIC NOTES/PEDIGREE TO AVOID DELAYS

PLEASE SUBMIT THE FOLLOWING WITH THE TRF:
1. Clinic Notes 2. Pedigree 3. Insurance Card

PATIENT INFORMATION

Name (Last, First, MI)	Date of Birth (MM/DD/YY)	Date of Death (If applicable)	Phone Number/Email

Address City State Zip	Biological Sex ☐F ☐M	Ethnicity: ☐African American ☐Asian ☐Caucasian ☐Hispanic ☐Jewish (Ashkenazi) ☐Portuguese ☐Other:

SPECIMEN INFORMATION* (For phlebotomy service, select all services you are requesting)

Type(s) ☐Blood (EDTA preferred) ☐Saliva ☐DNA ☐Other: ☐Personal history of allogenic bone marrow or peripheral stem cell transplant

Collection Date	Specimen ID	Medical Record #

**Blood or saliva from patients with active/recent hematological disease will undergo additional review and may not be accepted in some cases. For these, cultured fibroblasts or fresh/fresh frozen normal tissue are preferred. See ambrygen.com/specimen-requirements for details.*

Phlebotomy Services Request: ☐Phlebotomy draw ☐Insurance preverification first ☐Send kit to patient**

*** As the patient's clinician, I am unaware of any potential for complication or difficulty in drawing blood for the listed patient(s). I understand that the phlebotomist has full authority to refuse to draw any patient if the safety of the phlebotomist and/or patient(s) are in question.*

ORDERING LICENSED PROVIDER/SENDING FACILITY (Each listed person will receive a copy of the report)

Facility Name (Facility Code)	Address	City	State /Country	Zip	Phone

Ordering Licensed Provider Name (Last, First)(Code)	NPI#	Phone	Fax/Email

Additional Results Recipients

Genetic Counselor or Other Medical Provider Name (Last, First) (Code)	Phone/Fax/Email

Genetic Counselor or Other Medical Provider Name (Last, First) (Code)	Phone/Fax/Email

CONFIRMATION OF INFORMED CONSENT, PRE-TEST GENETIC COUNSELING, AND MEDICAL NECESSITY FOR GENETIC TESTING

The undersigned person (or representative thereof) ensures he/she is a licensed medical professional authorized to order genetic testing and confirms that the patient has given appropriate consent. I confirm that testing is medically necessary and that test results may impact medical management for the patient. I agree to allow Ambry Genetics to facilitate the provision of pre-test genetic counseling services by a third party service, Informed DNA (unless otherwise noted), as required by the patient's insurance provider (unless this box is checked ☐). Furthermore, all information on this TRF is true to the best of my knowledge. My signature applies to the attached letter of medical necessity.

Signature Required for Processing Medical Professional Signature: Date:

☐ INSURANCE BILLING (Include copy of both sides of insurance card)

Patient Relation to Policy Holder? ☐Self ☐Spouse ☐Child	Name and DOB of Policy Holder (if not self)	
Insurance Company	Policy #	HMO Auth #

Out Of Pocket:
We will start testing immediately, unless you check the box below. We will attempt to contact you if your estimated out-of-pocket costs are > USD $100.
☐ Do not start testing until I approve payment terms regarding estimated out-of-pocket costs
Patient agrees to contact regarding out-of-pocket amount by:
☐ Email ☐ Phone (includes texts) - confirm mobile # _____

Special Billing Notes:

☐ INSTITUTIONAL BILLING

Facility Name ☐ *Send invoice to facility address above*

Address

Contact Name

Phone Number	E-mail/Fax

☐ PATIENT PAYMENT

☐ Check (Payable to Ambry Genetics)
☐ Credit Card (Call 949-900-5795)

Patient Acknowledgement: I acknowledge that the information provided by me is true and correct. For direct insurance billing: I authorize my insurance benefits to be paid directly to Ambry Genetics Corporation (Ambry), authorize Ambry to release medical information concerning my testing to my insurer, to be my designated representative for purposes of appealing any denial of benefits as needed and to request additional medical records for this purpose. I understand that I am financially responsible for any amounts not covered by my insurer and responsible for sending Ambry money received from my health insurance company.

For patient payment by credit card: I hereby authorize Ambry Genetics Corporation to bill my credit card as indicated above. In order to expedite consideration for eligibility for **Ambry's Patient Assistance Program**, please provide the total annual gross household income: $_____ and the number of family members in the household supported by the listed income: _____. I authorize Ambry Genetics Corporation to verify the above information for the sole purpose of assessing financial need, including the right to seek supporting documentation.

FOR NY RESIDENTS:
☐ I am a New York resident and I give Ambry Genetics permission to store my sample for longer than 60 days. **NOTE:** If left blank, consent is interpreted as "NO".

Signature Required For Insurance/Self-Pay Patients and NY Sample Storage Consent: Date:

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INDICATIONS FOR TESTING (Check all that apply)

ICD-10 code(s): _____

Will patient management be changed depending on the test results? ☐ Yes ☐ No ☐ STAT TEST: Date results needed (if known):_____

PATIENT CLINICAL HISTORY

☐ No personal history of cancer

Cancer/Tumor	Age at Dx	Pathology and Other Info
Brain tumor		
Breast		Type: ER☐(+) ☐(-) ☐unk PR☐(+) ☐(-) ☐unk HER2/neu☐(+) ☐(-) ☐unk Metastatic:☐Y ☐N
2nd primary breast		Type: ER☐(+) ☐(-) ☐unk PR☐(+) ☐(-) ☐unk HER2/neu☐(+) ☐(-) ☐unk Metastatic:☐Y ☐N
Colorectal		Location:
Melanoma		
Ovarian		☐ Fallopian tube ☐ Primary peritoneal
Pancreatic		
Prostate		Gleason Score: Metastatic: ☐Y ☐N
Uterine		
Hematologic		Type: ☐Allogenic bone marrow or peripheral stem cell transplant*
Other Cancer		Type:
GI polyps		☐Adenomatous Polyp #:☐1 ☐2-5 ☐6-9 ☐10-19 ☐20-99 ☐100+ ☐Other type: Polyp #:☐1 ☐2-5 ☐6-9 ☐10-19 ☐20-99 ☐100+

Other clinical history:

**Blood or saliva from patients with active/recent hematological disease will undergo additional review and may not be accepted in some cases. For these, cultured fibroblasts or fresh/fresh frozen normal tissue are preferred. See ambrygen.com/specimen-requirements for details.*

PATIENT TESTING HISTORY (Please include copies of any previous test results)

☐ No previous molecular and/or genetic testing

☐ Germline genetic testing Test(s) performed:_____

Result (s):_____

☐ Somatic test/tumor profile Test(s) performed:_____

Result(s): _____

☐ Microsatellite instability analysis:

☐ Stable (MSS) ☐ Unstable/high (MSI-H) ☐ Unstable/low (MSI-L)

☐ IHC, if multiple primaries, tumor used:_____

☐ Proteins present: _____ ☐ Proteins absent: _____

FAMILY HISTORY*

**Completing this section is not mandatory for ordering if a pedigree and/or clinical note with family history is supplied, but is recommended and helps with results interpretation and claims filing.*

☐ None (maternal) ☐ None (paternal) ☐ Maternal history unknown ☐ Paternal history unknown

Relation to patient	Maternal	Paternal	Cancer/Polyp Type	Dx age	Relation to patient	Maternal	Paternal	Cancer/Polyp Type	Dx age
	☐	☐				☐	☐		
	☐	☐				☐	☐		
	☐	☐				☐	☐		
	☐	☐				☐	☐		
	☐	☐				☐	☐		
	☐	☐				☐	☐		

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For multiple tests, testing will be run concurrently (initiated at the same time) unless otherwise specified. For reflexive testing (second test starts pending first test outcome), indicate the order of reflexive tests in the notes section or next to the test check box. For reflex test orders, any positive findings (pathogenic/likely pathogenic) in the first test will be reported, and the second test will be cancelled; all other findings will automatically reflex (including VUS).

MULTI-GENE ORDERS

Select the indication for testing:

☐ Hereditary polyposis[1]

☐ Lynch syndrome/HNPCC[2]

☐ Hereditary breast and ovarian cancer[3]

☐ Testing is clinically indicated for other gene(s):

☐ None of the above

To complete your multi-gene order, please select a test option to the right. (See supplemental information on page 4 for details).

[1] APC/MUTYH [2] MLH1, MSH2, MSH6, PMS2, EPCAM [3] BRCA1/2
** Required: completed CustomNext-Cancer supplemental form.* ambrygen.com/forms

☐	BrainTumorNext	8847	27 gene brain tumor test
☐	BRCAplus	8836	8 gene breast cancer test
☐	BreastNext	8820	17 gene breast cancer test
☐	CancerNext	8824	34 gene cancer test
☐	CancerNext-*Expanded*	8874	67 gene cancer test
☐	ColoNext	8822	17 gene colorectal cancer test
☐	CustomNext-*Cancer*	9510	up to 81 gene custom test*
☐	GYNplus	8835	13 gene ovarian/uterine cancer test
☐	MelanomaNext	8849	8 gene melanoma test
☐	OvaNext	8830	25 gene ovarian/breast/uterine cancer test
☐	PancNext	8042	13 gene pancreatic cancer test
☐	Pancreatitis panel	8022	6 gene pancreatitis test
☐	PGLNext	5504	12 gene PGL/PCC test
☐	ProstateNext	8845	14 gene prostate cancer test
☐	RenalNext	5900	19 gene renal cancer test

OPTIONAL ADD-ON

Add AmbryScore: ☐ Breast (Supplemental Ordering Form REQUIRED) ☐ Prostate

SINGLE SYNDROME ORDERS

Check to order	Test Name	Test Code	Description
Hereditary Breast and/or Ovarian Cancer			
☐	*ATM*	9014	Ataxia-telangiectasia
☐	*BRCA1/2*	8838	Hereditary breast and ovarian cancer
☐	*BRCA1/2* Ashkenazi Jewish 3-site mutation panel	5892	
☐	*BRCA1/2* Ashkenazi Jewish 3-site mutation panel with reflex to *BRCA1/2* analysis if negative	5894	
☐	*CHEK2*	9016	
☐	*DICER1*	5260	
☐	*PALB2*	2366	
☐	*PTEN*	2106	*PTEN*-related disorders (including Cowden syndrome)
☐	*TP53*	2866	Li-Fraumeni syndrome
Gastrointestinal Cancer			
☐	*APC*	3040	Familial adenomatous polyposis
☐	*APC* and *MUTYH*	8726	Adenomatous polyposis
☐	*BMPR1A* and *SMAD4*	8604	Juvenile polyposis syndrome
☐	*CDH1*	4726	Hereditary diffuse gastric cancer
☐	*EPCAM* del/dup	8519	Lynch syndrome
☐	Lynch syndrome	8517	*MLH1, MSH2, MSH6, PMS2 + EPCAM* del/dup
☐	Lynch syndrome (sequential)	8515	Step 1: *MLH1, MSH2,* and *MSH6 + EPCAM* del/dup; Step 2: *PMS2*
☐	*MLH1*	8508	Lynch syndrome
☐	*MSH2 + EPCAM* del/dup	8510	Includes *MSH2* inversion
☐	*MSH2* inversion	2226	Lynch syndrome
☐	*MSH6*	8512	Lynch syndrome
☐	*MUTYH*	4661	*MUTYH*-associated polyposis
☐	*PMS2*	4646	Lynch syndrome
☐	*STK11*	2766	Peutz-Jeghers syndrome

Check to order	Test Name	Test Code	Description
Genitourinary Cancer			
☐	*BAP1*	9044	
☐	*FH*	6301	Hereditary leiomyomatosis and renal cell cancer
☐	*FLCN*	5921	Birt-Hogg-Dubé syndrome
☐	*VHL*	2606	Von-Hippel Lindau disease
☐	*TSC1* and *TSC2*	5904	Tuberous sclerosis complex
Endocrine Tumors			
☐	*MEN1*	2646	Multiple endocrine neoplasia type 1
☐	*RET* gene sequence	2680	Multiple endocrine neoplasia type 2
Skin Cancer/Melanoma			
☐	*CDKN2A* and *CDK4* concurrent	4708	Familial atypical multiple mole melanoma (FAMMM)
☐	*PTCH1*	5684	Gorlin syndrome
Other Hereditary Cancer Testing			
☐	*NF1*	5704	Neurofibromatosis type 1
☐	*NF2*	9024	Neurofibromatosis type 2
☐	*RB1*	5426	Hereditary retinoblastoma
☐	*SMARCB1*	7180	Schwannomatosis
Other Orders			
☐	Please visit ambrygen.com for a list of available tests. Test Code(s): _____ Gene/Test Name(s): _____		

SPECIFIC SITE ANALYSIS (Please include a copy of relative's report)

Gene(s): _____ Mutation(s): _____

Relative Name: _____

Relationship to Relative:_____Accession # (if tested at Ambry): _____

Positive control sample: ☐ will be provided ☐ already at Ambry ☐ not available

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Supplemental Information

Hereditary Cancer Multi-Gene Tests

Test Name	Test Code	Genes
Adenomatous polyposis	8726	*APC, MUTYH*
BrainTumorNext (27 genes)	8847	*AIP, ALK, APC, CDKN1B, CDKN2A, DICER1, MEN1, MLH1, MSH2, MSH6, NBN, NF1, NF2, PHOX2B, PMS2, POT1, PRKAR1A, PTCH1, PTEN, SMARCA4, SMARCB1, SMARCE1, SUFU, TP53, TSC1, TSC2, VHL*
BRCAplus (8 genes)	8836	*ATM, BRCA1, BRCA2, CDH1, CHEK2, PALB2, PTEN, TP53*
BreastNext* (17 genes)	8820	*ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, MRE11A, MUTYH, NBN, NF1, PALB2, PTEN, RAD50, RAD51C, RAD51D, TP53*
CancerNext*,^ (34 genes)	8824	*APC, ATM, BARD1, BRCA1, BRCA2, BRIP1, BMPR1A, CDH1, CDK4, CDKN2A, CHEK2, DICER1, EPCAM, GREM1, HOXB13, MLH1, MRE11A, MSH2, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, POLD1, POLE, PTEN, RAD50, RAD51C, RAD51D, SMAD4, SMARCA4, STK11, TP53*
CancerNext-*Expanded*^,^ (67 genes)	8874	*AIP, ALK, APC, ATM, BAP1, BARD1, BLM, BRCA1, BRCA2, BRIP1, BMPR1A, CDH1, CDK4, CDKN1B, CDKN2A,CHEK2, DICER1, EPCAM, FANCC, FH, FLCN, GALNT12, GREM1, HOXB13, MAX, MEN1, MET, MITF, MLH1, MRE11A, MSH2, MSH6, MUTYH, NBN, NF1, NF2, PALB2, PHOX2B, POT1, PMS2, POLD1, POLE, PRKAR1A, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RB1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2*
ColoNext (17 genes)	8822	*APC, BMPR1A, CDH1, CHEK2, EPCAM, GREM1, MLH1, MSH2, MSH6, MUTYH, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53*
CustomNext-*Cancer*^,^ (up to 81 genes) Required: complete CustomNext-*Cancer* supplemental form. ambrygen.com/forms	9510	*AIP, ALK, APC, ATM, AXIN2, BAP1, BARD1, BLM, BRCA1, BRCA2, BRIP1, BMPR1A, CASR, CDC73, CDH1, CDK4, CDKN1B, CDKN2A, CFTR, CHEK2, CPA1, CTNNA1, CTRC, DICER1, EGFR, EPCAM, FANCC, FH, FLCN, GALNT12, GREM1, HOXB13, KIT, MAX, MEN1, MET, MITF, MLH1, MRE11A, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NF2, NTHL1, PALB2, PDGFRA, PHOX2B, POT1, PMS2, POLD1, POLE, PRKAR1A, PRSS1, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RB1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, SPINK1, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2*
GYNplus (13 genes)	8835	*BRCA1, BRCA2, BRIP1 , EPCAM, MLH1, MSH2, MSH6, PALB2, PMS2, PTEN, RAD51C, RAD51D, TP53*
HBOC	8838	*BRCA1, BRCA2*
Lynch syndrome/HNPCC	8517	*MLH1, MSH2, MSH6, PMS2* + *EPCAM* del/dup
MelanomaNext (8 genes)	8849	*BAP1, BRCA2, CDK4, CDKN2A, MITF, PTEN, RB1, TP53*
OvaNext* (25 genes)	8830	*ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, DICER1, EPCAM, MLH1, MRE11A, MSH2, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, PTEN, RAD50, RAD51C, RAD51D, SMARCA4, STK11, TP53*
PancNext (13 genes)	8042	*APC, ATM, BRCA1, BRCA2, CDKN2A, EPCAM, MLH1, MSH2, MSH6, PALB2, PMS2, STK11, TP53*
Pancreatitis panel (6 genes)	8022	*CASR, CFTR, CPA1, PRSS1, SPINK1, CTRC*
PGLNext (12 genes)	5504	*FH, MAX, MEN1, NF1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127, VHL*
ProstateNext^ (14 genes)	8845	*ATM, BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PALB2, PMS2, RAD51D, TP53*
RenalNext (19 genes)	5900	*BAP1, EPCAM, FH, FLCN, MET, MITF, MLH1, MSH2, MSH6, PMS2, PTEN, SDHA, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL*

* AmbryScore for Breast is available as an add on to this panel if all of the following eligibility criteria are met:
- Female biological sex
- 18-84 years old
- Non-Ashkenazi Jewish, N. European ancestry
- No personal history of cancer (excluding non-melanoma skin cancer)
- No personal history of atypical hyperplasia or lobular carcinoma in situ (LCIS)
- No personal or family history of a mutation in a breast cancer susceptibility gene[1]

[1] *ATM, BARD1, BLM* (if tested), *BRCA1, BRCA2, BRIP1, CDH1, CHEK2, FANCC* (if tested), *MRE11A, NBN, NF1, PALB2, PTEN, RAD50, RAD51C, RAD51D, STK11, TP53*

Note: AmbryScore supplemental ordering form is required for processing

^ AmbryScore for Prostate is available as an add on to this panel if all of the following eligibility criteria are met:
- Male biological sex
- 18-84 years old
- N. European ancestry
- No personal or family history of a mutation in a prostate cancer susceptibility gene[2]

[2] *ATM, BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PALB2, PMS2, RAD51D, TP53*

Note: No additional ordering forms are required for processing