

To submit an order via email, please send the completed test requisition form to [info@ambrygen.com](mailto:info@ambrygen.com)

<b>1. SPECIMEN INFORMATION</b>				
Collection Date				
<input type="checkbox"/> Send saliva kit to patient				
<b>PLEASE SUBMIT THE FOLLOWING WITH THE TRF:</b>				
1. Clinic Notes    2. Pedigree    3. Insurance Card and Authorization Documents				
<b>2. PATIENT INFORMATION</b>				
Name (Last, First, MI)			Sex at Birth <input type="checkbox"/> F <input type="checkbox"/> M	Date of Birth (MM/DD/YY)
Ethnicity: <input type="checkbox"/> African American <input type="checkbox"/> Asian <input type="checkbox"/> White <input type="checkbox"/> Hispanic <input type="checkbox"/> Portuguese <input type="checkbox"/> Native American <input type="checkbox"/> Other:				MRN
Address		City		State
Phone		Email		Zip
Preferred Billing <input type="checkbox"/> Insurance <input type="checkbox"/> Self-pay <input type="checkbox"/> Institutional				
<b>3. ORDERING PROVIDER INFORMATION</b>				
Organization Name, Number		Address		City, State
Ordering Provider Name (Last, First), Ambry Number <input type="checkbox"/>		<input type="checkbox"/>		<input type="checkbox"/>
<input type="checkbox"/>		<input type="checkbox"/>		<input type="checkbox"/>
Genetic Counselor/Other Healthcare Professional Name (Last, First), Ambry Number <input type="checkbox"/>		<input type="checkbox"/>		<input type="checkbox"/>
<b>4. PATIENT CLINICAL HISTORY</b> <small>Attach clinic notes and/or pedigree</small>				
Personal History of Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No	Age of Dx	Diagnosis Notes (cancer type, etc.)		Metastatic <input type="checkbox"/> Yes <input type="checkbox"/> No
ICD-10 Code(s)				
Family History of Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No	Family History Details (include relative, cancer type, and age of diagnosis)			
Prior Genetic Testing, IHC, or MSI <input type="checkbox"/> Yes <input type="checkbox"/> No	Patient Testing Details		Family Members Testing Details	
<b>5. TEST ORDERS</b>				
<b>REQUIRED: Select a Primary Test Order</b>				
<b>For Patients Meeting BRCA1/2 Testing Criteria</b>				
<input type="checkbox"/> BRCA1/2 test				
<b>For Patients Meeting Colorectal Cancer Syndrome Testing Criteria (Lynch)</b>				
Lynch Syndrome test: <input type="checkbox"/> MLH1, MSH2, MSH6, PMS2, EPCAM				
<b>For Patients Meeting Colorectal Cancer Syndrome Testing Criteria (polyposis)</b>				
Polyposis test: <input type="checkbox"/> APC/MUYTH				
<input type="checkbox"/> Other: _____				
<input type="checkbox"/> None of the above (patient does not meet any genetic testing criteria)				
Will the course of treatment change depending upon the results of the test? <input type="checkbox"/> Yes <input type="checkbox"/> No				
STAT TEST: <input type="checkbox"/> Date results needed (if known): _____				
<b>Was genetic counseling completed?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown   Date Genetic Counseling was Performed: _____				
Patient Signature (I agree to terms below):				Date:
Medical Professional Signature (I agree to terms below):				Date:
<b>TERMS AND CONDITIONS</b>				
<b>Patient Acknowledgement:</b> 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 Ambyr Genetics Corporation (Ambyr), authorize Ambyr 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 Ambyr money received from my health insurance company.				
<b>We will start testing immediately and we will attempt to contact the patient if the estimated out-of-pocket costs are &gt; USD \$100.</b>				
<b>For patient payment by credit card:</b> I hereby authorize Ambyr Genetics Corporation to bill my credit card as indicated above. In order to expedite consideration for eligibility for Ambyr'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 Ambyr Genetics Corporation to verify the above information for the sole purpose of assessing financial need, including the right to seek supporting documentation.				
<b>For NY Residents:</b> <input type="checkbox"/> I am a New York resident and I give Ambyr Genetics permission to store my sample for longer than 60 days. NOTE: If left blank, consent is interpreted as "NO".				
<b>Medical Professional:</b> 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 Ambyr Genetics to facilitate the provision of pre-test genetic counseling services by a third-party service, as required by the patient's insurance provider. Furthermore, all information on this TRF is true to the best of my knowledge. My signature applies to the attached letter of medical necessity.				
Blood/saliva from patients with a history of allogeneic bone marrow or stem cell transplant cannot be used for genetic testing. Blood/saliva from patients with active hematological disease is not recommended. An alternative specimen may be needed. Please see <a href="http://ambrygen.com/specimen-requirements">ambrygen.com/specimen-requirements</a> for details.				

## Supplemental Information

### Hereditary Cancer Multi-Gene Tests

Test Name	Test Code	Genes
Adenomatous polyposis	8726	APC, MUTYH
BrainTumorNext® (29 genes)	8847	AIP, ALK, APC, CDKN1B, CDKN2A, DICER1, EPCAM, LZTR1, MEN1, MLH1, MSH2, MSH6, NBN, NF1, NF2, PHOX2B, PMS2, POT1, PRKAR1A, PTCH1, PTEN, SMARCA4, SMARCB1, SMARCE1, SUFU, TP53, TSC1, TSC2, VHL
BRCANext™^ (18 genes)	8855	ATM, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, EPCAM, MLH1, MSH2, MSH6, NBN, NF1, PALB2, PMS2, PTEN, RAD51C, RAD51D, TP53
BRCANext-Expanded™^ (23 genes)	8860	ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, DICER1, EPCAM, MLH1, MSH2, MSH6, NBN, NF1, PALB2, PMS2, PTEN, RAD51C, RAD51D, RECQL, SMARCA4, STK11, TP53
BRCAPlus® (8 genes)	8836	ATM, BRCA1, BRCA2, CDH1, CHEK2, PALB2, PTEN, TP53
CancerNext®^ (36 genes)	8824	APC, ATM, AXIN2, BARD1, BRCA1, BRCA2, BRIP1, BMPR1A, CDH1, CDK4, CDKN2A, CHEK2, DICER1, EPCAM, GREM1, HOXB13, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, RECQL, SMAD4, SMARCA4, STK11, TP53
CancerNext-Expanded®^ (77 genes)	8874	AIP, ALK, APC, ATM, AXIN2, BAP1, BARD1, BLM, BRCA1, BRCA2, BRIP1, BMPR1A, CDC73, CDH1, CDK4, CDKN1B, CDKN2A, CHEK2, CTNNA1, DICER1, EGFR, EGLN1, EPCAM, FANCC, FH, FLCN, GALNT12, GREM1, HOXB13, KIF1B, KIT, LZTR1, MAX, MEN1, MET, MIF, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NF2, NTHL1, PALB2, PDGFRA, PHOX2B, POT1, PMS2, POLD1, POLE, PRKAR1A, PTCH1, PTEN, RAD51C, RAD51D, RB1, RECQL, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2
ColoNext® (20 genes)	8822	APC, AXIN2, BMPR1A, CDH1, CHEK2, EPCAM, GREM1, MLH1, MSH2, MSH3, MSH6, MUTYH, NTHL1, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53
CustomNext-Cancer®^ (up to 91 genes) Required: complete CustomNext-Cancer supplemental form. <a href="http://ambrygen.com/forms">ambrygen.com/forms</a>	9510	AIP, ALK, APC, ATM, AXIN2, BAP1, BARD1, BLM, BRCA1, BRCA2, BRIP1, BMPR1A, CASR, CDC73, CDH1, CDK4, CDKN1B, CDKN2A, CFTR, CHEK2, CPA1, CTNNA1, CTSC, DICER1, EGFR, EGLN1, EPCAM, FAM175A(ABRAXAS1)†, FANCC, FH, FLCN, GALNT12, GREM1, HOXB13, KIF1B, KIT, LZTR1, MAX, MEN1, MET, MIF, MLH1, MLH3†, MRE11A †, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NF2, NTHL1, PALB2, PALLD†, PDGFRA, PHOX2B, POT1, PMS2, POLD1, POLE, PRKAR1A, PRSS1, PTCH1, PTEN, RAD50 †, RAD51C, RAD51D, RB1, RECQL, RET, RINT1†, RPS20†, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, SPINK1, STK11, SUFU, TERT†, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2
HBOC	8838	BRCA1, BRCA2
Lynch syndrome/HNPCC	8517	MLH1, MSH2, MSH6, PMS2 + EPCAM del/dup
MelanomaNext® (9 genes)	8849	BAP1, BRCA2, CDK4, CDKN2A, MIF, POT1, PTEN, RB1, 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, CTSC
PGLNext® (14 genes)	5504	EGLN1, FH, KIF1B, 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® (20 genes)	5900	BAP1, CHEK2, EPCAM, FH, FLCN, MET, MIF, MLH1, MSH2, MSH6, PMS2, PTEN, SDHA, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL

Genes Eligible for +RNAinsight®: APC, ATM, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, MLH1, MSH2, MSH6, MUTYH, NF1, PALB2, PMS2 EX1-10, PTEN, RAD51C, RAD51D, TP53

^ 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 (ATM, BARD1 [if tested], BLM [if tested], BRCA1, BRCA2, BRIP1, CDH1, CHEK2, FANCC [if tested], NBN, NF1, PALB2, PTEN, 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 (ATM, BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PALB2, PMS2, RAD51D, TP53)

Note: No additional ordering forms are required for processing

† Limited evidence gene