

COMPLETE ENTIRE FORM TO AVOID DELAYS

Form containing sections: PATIENT INFORMATION, FAMILY HISTORY, PATIENT HISTORY, SPECIMEN INFORMATION, ORDERING PHYSICIAN/SENDING FACILITY, CONFIRMATION OF INFORMED CONSENT AND MEDICAL NECESSITY FOR GENETIC TESTING, INDICATIONS FOR TESTING, TEST REQUESTED - GI CANCER TEST OPTIONS, INSURANCE BILLING, INSTITUTIONAL BILLING, and Research & Recontact Consent.

**REQUIRED INSURANCE ORDERING CHECKLIST**

- Clinic Notes (Pedigree if available)
- ICD-10 Code(s)
- Clinician & Patient Signatures
- Insurer Specific Forms (i.e. ABN)
- Copy of Insurance Cards

**Hereditary Cancer Multi-Gene Tests**

Test Name	Test Code	Genes
<b>Adenomatous polyposis</b>	<b>8726</b>	<b>APC, MUTYH</b>
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 (6 genes)	8836	BRCA1, BRCA2, CDH1, PALB2, PTEN, TP53
BRCAPlus-Expanded (8 genes)	8837	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
<b>CancerNext (34 genes)</b>	<b>8824</b>	<b>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</b>
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, MIF, 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, TMMEM127, TP53, TSC1, TSC2, VHL, XRCC2
<b>ColoNext (17 genes)</b>	<b>8822</b>	<b>APC, BMPR1A, CDH1, CHEK2, EPCAM, GREM1, MLH1, MSH2, MSH6, MUTYH, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53</b>
CustomNext-Cancer (up to 67 genes) Required: complete CustomNext-Cancer supplemental form. <a href="http://ambrygen.com/forms">ambrygen.com/forms</a>	9510	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, MIF, 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, TMMEM127, 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
<b>Lynch syndrome/HNPCC</b>	<b>8517</b>	<b>MLH1, MSH2, MSH6, PMS2 + EPCAM del/dup</b>
MelanomaNext (8 genes)	8849	BAP1, BRCA2, CDK4, CDKN2A, MIF, 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
<b>PancNext (13 genes)</b>	<b>8042</b>	<b>APC, ATM, BRCA1, BRCA2, CDKN2A, EPCAM, MLH1, MSH2, MSH6, PALB2, PMS2, STK11, TP53</b>
<b>Pancreatitis panel (4 genes)</b>	<b>8022</b>	<b>CFTR, PRSS1, SPINK1, CTRC</b>
PGLNext (12 genes)	5504	FH, MAX, MEN1, NF1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMMEM127, 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, MIF, MLH1, MSH2, MSH6, PMS2, PTEN, SDHA, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL

**Updated Ordering Process (as of June 8, 2016)**

We have improved the ordering and reporting process for our hereditary cancer panels. This helps confirm that testing for one or more of the following genes is clinically indicated: *APC, BRCA1, BRCA2, EPCAM, MLH1, MSH2, MSH6, MUTYH, and PMS2*.

**If you are ordering a multi-gene test**, please first select a clinically indicated condition and complete your order by selecting a multi-gene order.

Please indicate if your patient meets clinical and/or insurance testing criteria, or if the testing is otherwise clinically indicated for one or more of the following conditions:

- Adenomatous polyposis (*APC/MUTYH*)
- Hereditary breast and ovarian cancer (*BRCA1/2*)
- Lynch syndrome/HNPCC (*MLH1, MSH2, MSH6, PMS2, EPCAM*)

If testing is not clinically indicated for your patient for any of the listed options, please either fill in the other gene(s) option or select "none of the above".

To complete your multi-gene order, please select one of the appropriate test options and/or select "other" and enter an appropriate test code(s)/test name(s).

**For single gene orders**, please enter the gene(s) and/or test name(s), as well as the relevant test code in the single gene orders section.

For additional details about our single gene testing options, please visit [ambrygen.com/hereditary-cancer-single-gene-tests](http://ambrygen.com/hereditary-cancer-single-gene-tests).

**Example**

For a ColoNext multi-gene order, when Lynch syndrome testing is clinically indicated for the patient:

**Multi-Gene Orders:** For multi-gene orders, first select which of the following conditions is clinically indicated based on the patient's personal and/or family history:

- Adenomatous polyposis (*APC/MUTYH*)
- Hereditary breast and ovarian cancer (*BRCA1/2*)
- Lynch syndrome/HNPCC (*MLH1, MSH2, MSH6, PMS2, EPCAM*)
- Testing is clinically indicated for other gene(s):
- None of the above

To complete your multi-gene order, please select a test option below (see supplemental pages for details):

- ColoNext (8822)     PancNext (8042)     Pancreatitis panel (8022)
- CancerNext (8824)     Other Test Code: \_\_\_\_\_ Test Name: \_\_\_\_\_

For single gene genetic testing of *CDH1*:

**Single Syndrome Orders**

Single gene analysis is available for listed panels. Visit [ambrygen.com/hereditary-cancer-single-gene-tests](http://ambrygen.com/hereditary-cancer-single-gene-tests) for details.

- Lynch syndrome (8517)\*

Test Code(s): 4726                      Gene/Test Name(s): *CDH1*

\* If patient has Medicare and only Lynch is being ordered, please order 8515.

**Single Site Analysis (SSA) (Include relative report)**

Gene(s): \_\_\_\_\_ Mutation(s): \_\_\_\_\_

Previously Tested Relative (name): \_\_\_\_\_

Relationship to Relative: \_\_\_\_\_

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