

CLIENT INFORMATION	
ORDERING PHYSICIAN	NPI #
TREATING PHYSICIAN	NPI #
PHYSICIAN/AUTHORIZED SIGNATURE	
Client#	
Client Name	
Address	
Phone Number	Fax Number
PATIENT INFORMATION	
Name (LAST, FIRST, MIDDLE):	
Date of Birth:	Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female
Address:	
City, State, Zip:	
Phone Number:	MRN / PT ID#:
BILLING INFORMATION (face sheet & front and back of insurance card must be attached)	
Bill: <input type="checkbox"/> My Account <input type="checkbox"/> Insurance <input type="checkbox"/> Medicare <input type="checkbox"/> Medicaid <input type="checkbox"/> Patient <input type="checkbox"/> Workers Comp <input type="checkbox"/> See attached	
Patient Hospital Status: <input type="checkbox"/> InPatient <input type="checkbox"/> OutPatient <input type="checkbox"/> Non-Patient	
Insured Information: Name	
Relationship to Patient (circle one)	Self Spouse Child Other:
Primary Insurance Co:	Authorization #
Billing Address	Insured #
Billing City, State, Zip	Group #
SPECIMEN INFORMATION	
Collection Date:	Time: <input type="checkbox"/> AM <input type="checkbox"/> PM
Specimen Type: <input type="checkbox"/> Peripheral Blood	Specimen ID #:
PATIENT CLINICAL CANCER HISTORY	
<input type="checkbox"/> No personal history of cancer	
<input type="checkbox"/> Breast: Invasive or DCIS, age at Dx _____ (Check all that apply) <input type="checkbox"/> Bilateral <input type="checkbox"/> Premenopausal <input type="checkbox"/> Triple Negative (ER-, PR-, HER-)	
<input type="checkbox"/> Ovarian: Age at Dx _____	<input type="checkbox"/> Endometrial: Age at Dx _____
<input type="checkbox"/> Prostate: Age at Dx _____	If Prostate, Gleason score: _____
<input type="checkbox"/> Pancreatic: Age at Dx _____	<input type="checkbox"/> Renal: Age at Dx _____
<input type="checkbox"/> Endocrine: Age at Dx _____	Type: <input type="checkbox"/> Thyroid <input type="checkbox"/> Pheochromocytoma <input type="checkbox"/> Paranglioma
<input type="checkbox"/> Colorectal: Age at Dx _____	
MSI Testing Done: <input type="checkbox"/> Yes <input type="checkbox"/> No MSI Result: <input type="checkbox"/> High <input type="checkbox"/> Stable <input type="checkbox"/> Low	
IHC Testing Done: <input type="checkbox"/> Yes <input type="checkbox"/> No IHC Result: <input type="checkbox"/> Present <input type="checkbox"/> Absent IHC of _____	
MLH1 Methylation status: <input type="checkbox"/> Not done <input type="checkbox"/> Methylated tumor only <input type="checkbox"/> Methylated tumor and normal tissue <input type="checkbox"/> Unmethylated	
BRAF V600E: <input type="checkbox"/> Not Done <input type="checkbox"/> Present <input type="checkbox"/> Absent	
<input type="checkbox"/> Colon Polyps: Age of first polyp _____ If Adenomatous, total # _____	
<input type="checkbox"/> Other Cancers: Type _____ Age(s) of Dx: _____	
<input type="checkbox"/> History of Bone Marrow Transplant: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, please contact us at 800-345-4363 prior to sample collection.	
<input type="checkbox"/> History of blood transfusion, date of last transfusion _____	
CLINICAL INDICATION (attach previous test results if applicable)	
Narrative Diagnosis:	
All Diagnoses should be provided by the ordering physician or an authorized designee. Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Service (Highest Specificity Required)	
ICD-CM	ICD-CM
Genetic Counseling Provided: <input type="checkbox"/> Yes <input type="checkbox"/> No Counselor name: _____	
Institution: _____ Phone #: _____	
Note: Some insurance companies require genetic counseling before approving BRCAAssure and VistoSeq testing. If you are unsure, please contact us at 855-488-8750.	

TEST REQUEST																														
<p style="text-align: right;">^ Please attach a copy of family member's test results</p> <p>VISTASEQ® HEREDITARY CANCER TESTS (See reverse side for gene list)</p> <p>Panels:</p> <p><input type="checkbox"/> Hereditary Cancer Panel (27 Gene Assay) (481220)</p> <p><input type="checkbox"/> Hereditary Cancer Panel Minus BRCA1/2 genes (25 Gene Assay) (481240)</p> <p><input type="checkbox"/> Breast Cancer Panel (19 Gene Assay) (481319)</p> <p><input type="checkbox"/> High/Moderate Risk Breast Cancer Panel (9 Gene Assay) (481452)</p> <p><input type="checkbox"/> GYN Cancer Panel (11 Gene Assay) (481330)</p> <p><input type="checkbox"/> Breast and GYN Cancer Panel (25 Gene Assay) (481341)</p> <p><input type="checkbox"/> High Risk Colorectal Cancer Panel (7 Gene Assay) (481352)</p> <p><input type="checkbox"/> Colorectal Cancer Panel (22 Gene Assay) (481363)</p> <p><input type="checkbox"/> Brain/CNS/PNS Cancer Panel (17 Gene Assay) (481386)</p> <p><input type="checkbox"/> Endocrine Cancer Panel (13 Gene Assay) (481374)</p> <p><input type="checkbox"/> Pancreatic Cancer Panel (14 Gene Assay) (481385)</p> <p><input type="checkbox"/> Renal Cell Cancer Panel (19 Gene Assay) (481407)</p> <p><input type="checkbox"/> Lynch Syndrome Panel (5 Gene Assay) (483543)</p> <p><input type="checkbox"/> Prostate Cancer Panel (10 Gene Assay) (483555)</p> <p>Single Genes:</p> <p><input type="checkbox"/> APC Comprehensive Analysis* (483484)</p> <p><input type="checkbox"/> MEN1 Comprehensive Analysis* (483460)</p> <p><input type="checkbox"/> RET Comprehensive Analysis* (483472)</p> <p>HEREDITARY BREAST AND OVARIAN CANCER (test components on back)</p> <p><input type="checkbox"/> BRCAAssure® Comprehensive Analysis* (485030)</p> <p><input type="checkbox"/> BRCAAssure® BRCA1 Targeted Analysis^ (485066)</p> <p><input type="checkbox"/> BRCAAssure® BRCA2 Targeted Analysis^ (485081)</p> <p><input type="checkbox"/> BRCAAssure® Ashkenazi Jewish Panel (485097)</p> <p>LYNCH SYNDROME GERMLINE TESTING</p> <p><input type="checkbox"/> VistoSeq® Lynch Syndrome Panel (483543)</p> <p><input type="checkbox"/> VistoSeq® MLH1 Comprehensive Analysis* (483496)</p> <p><input type="checkbox"/> VistoSeq® MSH2 Comprehensive Analysis* (483508)</p> <p><input type="checkbox"/> VistoSeq® MSH6 Comprehensive Analysis* (483520)</p> <p><input type="checkbox"/> VistoSeq® PMS2 Comprehensive Analysis* (483532)</p> <p><input type="checkbox"/> EPCAM Deletion/Duplication Analysis (511654)</p> <p>Individual Tests: _____</p> <p>See reverse side for additional individual gene test list and write in test name/test code in the space above</p> <p><input type="checkbox"/> Mutation Specific Sequencing (451382)</p> <p>Required: Gene(s): _____</p> <p>Mutation(s): _____</p> <tr style="background-color: #800040; color: white;"> <th>PREVIOUS GENETIC TESTING FOR CANCER</th> </tr> <tr> <td> <p>Has the patient had genetic testing for cancer? If yes, please document the gene(s) tested and results: _____</p> </td> </tr> <tr style="background-color: #800040; color: white;"> <th>FAMILY HISTORY</th> </tr> <tr> <td> <p style="text-align: right;">^ Please attach a copy of family member's test results</p> <p>Is there a family history of cancer? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown</p> <p>Other family members tested positive for a hereditary cancer syndrome? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Type ^ : _____</p> <p>Is the patient adopted? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown</p> <p>Does the patient have Ashkenazi Jewish Ancestry? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Please attach pedigree or complete table below. 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I understand a preauthorization approval from my health plan does not guarantee full payment.</p> <p>Labcorp will attempt to contact me if my estimated out-of-pocket payment is more than \$300. Testing may be canceled if Labcorp is unable to reach me. No matter my estimated cost, my actual out-of-pocket cost may be higher or lower than the estimate provided. It is my responsibility to contact my health plan regarding concerns over my coverage and benefits.</p> <p>Patient's Signature (required) _____</p> <p>Patient Phone Number: _____</p> <p><input type="checkbox"/> Prior-authorization form has been sent separately. Please contact Prior Authorization at 855-488-8750 if you have any questions about our service.</p> <tr style="background-color: #800040; color: white;"> <th>INFORMED CONSENT</th> </tr> <tr> <td> <p>I have obtained informed consent of the patient (or the patient's authorized representative) for the ordered genetic test(s) in accordance with applicable law. 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VistaSeq® Hereditary Cancer Panels - Gene List

Gene Panel	Genes
Hereditary Cancer Panel	<i>APC, ATM, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, EPCAM, FAM175A, MLH1, MSH2, MSH6, MUTYH, NBN, PALB2, PMS2, PRKAR1A, PTEN, RAD51C, RAD51D, SMAD4, STK11, TP53</i>
Hereditary Cancer Panel Minus BRCA1/2 genes	<i>APC, ATM, BARD1, BMPR1A, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, EPCAM, FAM175A, MLH1, MSH2, MSH6, MUTYH, NBN, PALB2, PMS2, PRKAR1A, PTEN, RAD51C, RAD51D, SMAD4, STK11, TP53</i>
Breast Cancer Panel	<i>ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, FAM175A, MRE11A, MUTYH, NBN, NF1, PALB2, PTEN, RAD50, RAD51C, RAD51D, STK11, TP53</i>
High/Moderate Risk Breast Cancer Panel	<i>ATM, BRCA1, BRCA2, CDH1, CHEK2, PALB2, PTEN, STK11, TP53</i>
GYN Cancer Panel	<i>BRCA1, BRCA2, CHEK2, EPCAM, MLH1, MSH2, MSH6, MUTYH, PMS2, PTEN, TP53</i>
Breast and GYN Cancer Panel	<i>ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, EPCAM, FAM175A, MRE11A, FANCC, MLH1, MSH2, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, PTEN, RAD50, RAD51C, RAD51D, STK11, TP53</i>
High Risk Colorectal Cancer Panel	<i>APC, EPCAM, MLH1, MSH2, MSH6, MUTYH, PMS2</i>
Colorectal Cancer Panel	<i>APC, ATM, AXIN2, BLM, BMPR1A, BRCA1, BRCA2, CDH1, CDK2A, CHEK2, EPCAM, MLH1, MSH2, MSH6, MUTYH, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53</i>
Brain/CNS/PNS Cancer Panel	<i>ALK, APC, MEN1, MLH1, MSH2, MSH6, NBN, NF1, NF2, PHOX2B, PMS2, PTCH1, RB1, SMARCB1, SUFU, TP53, VHL</i>
Endocrine Cancer Panel	<i>CDC73, MAX, MEN1, NF1, PRKAR1A, PTEN, RET, SDHB, SDHC, SDHD, TMEM127, TP53, VHL</i>
Pancreatic Cancer Panel	<i>APC, ATM, BRCA1, BRCA2, CDKN2A, EPCAM, MLH1, MSH2, MSH6, PALB2, PMS2, STK11, TP53, VHL</i>
Renal Cell Cancer Panel	<i>EPCAM, FH, FLCN, GPC3, MET, MTF, MLH1, MSH2, MSH6, PMS2, PTEN, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL, WT1</i>
Lynch Syndrome Panel	<i>MLH1, MSH2, MSH6, PMS2, EPCAM</i>
Prostate Cancer Panel	<i>ATM, BRCA1, BRCA2, CHEK2, HOXB13, MLH1, MSH2, MSH6, PALB2, PMS2</i>

BRCAAssure® Test Components

Comprehensive Analysis	BRCA1 Target Analysis	BRCA2 Target Analysis	Ashkenazi Jewish Panel
Includes full gene sequencing and deletion/duplication analysis of BRCA1/2 genes	Includes sequencing of known familial mutation(s) on BRCA1 gene	Includes sequencing of known familial mutation(s) on BRCA2 gene	Includes screening for three known pathogenic variants: two in BRCA1 gene, and one in BRCA2 gene

Lynch Syndrome Germline Testing - Individual Test List (Test Names / Test Code)

Single Gene Tests

MLH1 Deletion/Duplication Analysis (511690)
 MSH2 Deletion/Duplication Analysis (511705)
 MSH6 Deletion/Duplication Analysis (511720)
 PMS2 Deletion/Duplication Analysis (511725)

Familial Tests with Known Mutations[▲]

MLH1 (511635)
 MSH2 (511750)
 MSH6 (511765)
 PMS2 (511776)
 EPCAM Deletion/Duplication Analysis (511654)

[▲] Please attach a copy of family member's test results

Patient, client, and billing information is requested for timely processing of this case. Medicare and other third party payors require that services be medically necessary for coverage, and generally do not cover routine screening tests

Symbols Legend

= Medicare deems investigational. Medicare does not pay for services it deems investigational.

Refer to Determining Necessity of ABN Completion.

Determining Necessity of Advance Beneficiary Notice of Non-coverage (ABN) Completion*

- 1. Diagnose.** Determine your patient's diagnosis.
- 2. Document.** Write the diagnosis code(s) on the front of this requisition.
- 3. Verify.** Determine if the laboratory test(s) ordered for the patient is subject to the Local Coverage Determination or National Coverage Determination. This information can be located in the policies published by your Medicare Administrative Contractor (MAC), CMS, or www.Labcorp.com/MedicareMedicalNecessity.
- 4. Review.** If the diagnosis code for your patient **does not** meet the medical necessity requirements set forth by Medicare or the test is being performed more frequently than Medicare allows, an ABN should be completed.

*An ABN should be completed for all tests that are considered investigational (experimental or for research use) by Medicare.

How to Complete an Advance Beneficiary Notice of Non-coverage (ABN)

Medicare is very specific in requiring that all of the information included on the ABN must be completed. Additionally, Labcorp requests that the specimen number or bar code label be included on the form. To be valid, an ABN must:

1. Be executed on the CMS approved ABN form (CMS-R-131).
2. Identify the Medicare Part B Beneficiary, using the name as it appears on the patient's red, white, and blue Medicare card.
3. Indicate the test(s)/procedure(s) which may be denied within the relevant reason column.
4. Include an estimated cost for the test(s)/procedures(s) subject to the ABN.
5. Have "Option 1", "Option 2", or "Option 3" designated by the beneficiary.
6. Be signed **and** dated by the beneficiary or his/her representative **prior to** the service being rendered.