

Hereditary Cancer



P +1888.354.8168 | F +1855.856.0655 | info@fulgentoncology.com

Highlighted fields are required information*

PATIENT INFORMATION

LAST NAME*

FIRST NAME*

SEX ASSIGNED AT BIRTH*

DATE OF BIRTH (MM/DD/YYYY)*

Male

Female

Unknown

MRN

ETHNICITY

ADDRESS

CITY

STATE

POSTAL CODE

COUNTRY

PHONE

EMAIL

SAMPLE TYPE

Blood

Buccal/Saliva

Other:

Extracted DNA & DNA Source
(Blood, Buccal, Tissue, Fibroblast):

SAMPLE DRAW DATE (MM/DD/YYYY)*

PATIENT ACKNOWLEDGEMENT

I have read the Informed Consent document and I give permission to Fulgent Genetics and its entities to perform genetic testing as described. I also give permission for my specimen and clinical information to be used in de-identified studies at Fulgent and for publication, if appropriate. My name or other personal identifying information will not be used in or linked to the results of any studies and publications. More information is available at [www.fulgentgenetics.com/policies/privacy-policy](#).

☐

Check this box if you are a New York state resident and give permission for Fulgent to retain any remaining sample longer than 60 days after sample collection.

☐

Opt out of research

X

Patient Signature (Required for billing purposes)*

Date (MM/DD/YYYY)

ORDERING PROVIDER

INSTITUTION/PRACTICE NAME

INSTITUTION PHONE / FAX

INSTITUTION EMAIL

ORDERING PROVIDER(S)

NPI (USA)/MINC (Canada)

PROVIDER TITLE (MD, DO, GC)

PROVIDER ADDRESS

CITY

STATE

POSTAL CODE

COUNTRY

PROVIDER PHONE

FAX/EMAIL REPORT TO

STATEMENT OF INFORMED CONSENT

By signing below, I, the ordering Medical Provider, confirm that testing is medically necessary and that test results may impact medical management for the patient.

I attest that the patient has received and read the Fulgent Informed Consent document, or has had it read to them, and that I have fully informed the patient about the purpose, capabilities, and limitations of the ordered test. The patient has voluntarily given his or her full consent for the ordered test and a signed copy of this consent is available on file. Any Fulgent Informed Consent that the patient agrees to at a later date will supersede and replace this Informed Consent.

X

Ordering Provider Signature (Required)*

Date (MM/DD/YYYY)

PATIENT COMMUNICATION CONSENT

☐

By checking this box, you acknowledge and agree that:

a.

By executing this agreement, you are providing express written consent for Fulgent, CSI, and Inform Diagnostics, their affiliates and subsidiaries, and parties making contact on their behalf to call and text you using automatic telephone dialing systems and artificial or pre-recorded messages at the telephone number you have provided, about your out-of-pocket estimation, even if your telephone number is currently listed on any state, federal, local or corporate Do Not Call list.

b.

Your Consent to be contacted through the use of automatic telephone dialing systems and artificial or pre-recorded messages is not required in order to purchase any property, goods, or services. If you would like to speak with our Benefits Investigation team, you can reach them at **+1 888.FULGENT (+1 888.385.4368), opt 3**.

SELECT TEST PANEL Please select only one test panel

All genes included on the cancer-specific panels are included on the full comprehensive panel. If multiple panels are selected, we will combine them into a single Custom Cancer Panel.

PAN-CANCER

Full Comprehensive Panel (FT-TP00048)

Includes all subpanels listed

Full Focus Panel (FT-TP00105)

BREAST & OVARIAN CANCERS

BRCA1 & BRCA2 Focus Panel (FT-TP01125)

Breast Cancer STAT Panel (FT-TP01030) (TAT: 10 DAYS)

Breast Cancer Focus Panel (FT-TP00101)

Ovarian Cancer Focus Panel (FT-TP00106)

Breast & Ovarian Cancer Focus Panel (FT-TP00462)

Breast Cancer Comprehensive Panel (FT-TP00043)

Ovarian Cancer Comprehensive Panel (FT-TP00053)

Breast & Ovarian Comprehensive Panel (FT-TP00461)

ENDOCRINE CANCERS

Multiple Endocrine Neoplasia Comprehensive Panel (FT-TP00182)

Paraganglioma-Pheochromocytoma Comprehensive Panel (FT-TP00055)

Thyroid Cancer Comprehensive Panel (FT-TP00059)

COLORECTAL CANCERS

Lynch Syndrome Focus Panel (FT-TP01543)

Colorectal Focus Panel (FT-TP00102)

Adenomatous Polyposis Focus Panel (FT-TP01534)

Colorectal Comprehensive Panel (FT-TP00044)

Polyposis Comprehensive Panel (FT-TP01535)

For the most up to date panel information and genes included please visit [FulgentOncology.com](#).

OTHERS

Endometrial Cancer Comprehensive Panel (FT-TP00046)

Gastric Cancer Comprehensive Panel (FT-TP00049)

Hematologic Malignancy Comprehensive Panel (FT-TP00050)

Melanoma Comprehensive Panel (FT-TP00051)

Nervous System/Brain Comprehensive Panel (FT-TP00052)

Pancreatic Cancer Comprehensive Panel (FT-TP00054)

Prostate Cancer Focus Panel (FT-TP00107)

Prostate Cancer Comprehensive Panel (FT-TP00056)

Renal/Urinary Cancer Comprehensive Panel (FT-TP00057)

Sarcoma Comprehensive Panel (FT-TP00058)

SINGLE GENE OR KNOWN MUTATION & ADDITIONAL REQUESTS

Custom Cancer Panel (FT-TP00045)

Please list gene(s) below.

TEST OPTION

Exclude VUS

Add RISE

Additional specimen required, please submit with proband to reduce delays.

Sample submitted with proband?

Yes

No

Additional Comments:

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Highlighted fields are required information*

BILLING OPTIONS

Please select a billing option and complete the relevant fields below: ☐ Insurance ☐ Institution ☐ Self-Pay

INSURANCE/BILLING INFORMATION

Please attach front and back of all insurance cards, ABN, medical criteria form

By signing above, the patient or payor authorizes Fulgent and its entities to contact them directly (including via text), and authorizes Fulgent and its entities to release medical information concerning the test to the assigned insurance company.

ICD-10 VALID CODE*		REFERRAL/PRIOR AUTH		FULGENT BENEFITS ID #
PRIMARY INSURANCE ID	INSURANCE NAME	STATE	GROUP	INSURANCE PHONE #
INSURANCE PLAN	NAME OF INSURED	RELATION TO PATIENT		DATE OF BIRTH (MM/DD/YYYY)
SECONDARY INSURANCE ID	INSURANCE NAME	STATE	GROUP	INSURANCE PHONE #
INSURANCE PLAN	NAME OF INSURED	RELATION TO PATIENT		DATE OF BIRTH (MM/DD/YYYY)

INSTITUTIONAL BILLING

- ☐ Use institution information above for billing
☐ Use information below for billing

INSTITUTION/PRACTICE NAME		ATTENTION TO	
ADDRESS			
CITY	STATE	POSTAL CODE	COUNTRY
PHONE	EMAIL		

SELF-PAY

- ☐ Use patient information above for billing
☐ Use information below for billing

By signing above, the patient or payor authorizes Fulgent and its entities to contact them directly, and use the provided billing instructions to bill the indicated method.

PAYOR LAST NAME		PAYOR FIRST NAME	
ADDRESS			
CITY	STATE	POSTAL CODE	COUNTRY
PHONE	EMAIL		

PATIENT CLINICAL HISTORY **Check all that apply**

☐ No personal history of cancer

INDICATIONS FOR TESTING **Check all that apply**

☐ Diagnostic ☐ Family History ☐ Family Variant ☐ Presymptomatic ☐ Other: _____

CANCER/TUMOR TYPE	AGE OF ONSET	PATHOLOGY AND OTHER INFO
<input type="checkbox"/> Brain		
<input type="checkbox"/> Breast		ER: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) PR: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) HER2/neu: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?)
<input type="checkbox"/> 2nd Primary Breast		ER: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) PR: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) HER2/neu: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?)
<input type="checkbox"/> Colorectal		Location: _____
<input type="checkbox"/> Hematologic		
<input type="checkbox"/> GI Polyps		<input type="checkbox"/> Adenomatous: _____ <input type="checkbox"/> Other: Number of polyp(s): _____

CANCER/TUMOR TYPE	AGE OF ONSET	PATHOLOGY AND OTHER INFO
<input type="checkbox"/> Melanoma		
<input type="checkbox"/> Ovarian		ER: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) PR: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) HER2/neu: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?)
<input type="checkbox"/> Fallopian tube		
<input type="checkbox"/> Primary peritoneal		
<input type="checkbox"/> Pancreatic		ER: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) PR: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?) HER2/neu: <input type="radio"/> POS(+) <input type="radio"/> NEG(-) <input type="radio"/> UNK(?)
<input type="checkbox"/> Prostate		Gleason score: _____ Metastatic: <input type="radio"/> Yes <input type="radio"/> No
<input type="checkbox"/> Uterine		
<input type="checkbox"/> Other Cancer		

CLINICAL HISTORY/SUSPECTED DIAGNOSIS

Please attach copy of recent CBC, copy of doctor's notes/clinical history, pathology reports, and any relevant test results.*

PATIENT TESTING HISTORY **Please attach relevant reports**

Germline testing results:	Microsatellite Instability (MSI) results:	Immunohistochemistry (IHC) results:
Somatic testing/Tumor profile results:	Other, specify: _____	

FAMILY HISTORY **Attach pedigree and additional pages as needed**

FAMILY MEMBER NAME (1)	RELATION TO PATIENT	SEX ASSIGNED AT BIRTH <input type="radio"/> Male <input type="radio"/> Female <input type="radio"/> Unknown
DIAGNOSIS AND/OR SYMPTOMS	AGE OF ONSET	DATE OF BIRTH (MM/DD/YYYY)
FAMILY MEMBER NAME (2)	RELATION TO PATIENT	SEX ASSIGNED AT BIRTH <input type="radio"/> Male <input type="radio"/> Female <input type="radio"/> Unknown
DIAGNOSIS AND/OR SYMPTOMS	AGE OF ONSET	DATE OF BIRTH (MM/DD/YYYY)

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INSTRUCTIONS

- 1. Complete the patient and provider information section.
- 2. Read and sign the consent statement. If needed, Patient Informed Consent forms for genetic testing can be found on FulgentOncology.com.
 - Signature from the provider on Page 1 of the TRF is required for all testing.
 - Signature from the patient is only required for billing purposes.
- 3. Indicate the test name and any relevant test options on Page 1 of this form.
- 4. Please visit FulgentOncology.com for specimen requirements.
 - Extracted DNA must be collected from a CLIA-certified laboratory or a laboratory meeting equivalent requirements as determined by CAP and/or CMS.

REQUIRED FOR INSURANCE CHECKLIST

- ☐ Detailed medical record (pedigree if available)
- ☐ ICD-10 code(s)
- ☐ Physician, patient, and insured signatures
- ☐ Copy of insurance card(s) - front/back
- ☐ Insurer specific forms (eg. ABN)
- ☐ Insurance authorization, if available
- ☐ For Medicare, a Medicare criteria form is required

For the most updated information and limitations on our products and services, please visit **FulgentOncology.com**.