

Date faxed: _____

This form MUST be faxed directly to BCBS FEP at 1-855-895-3504 on (or before) the sample collection date. Please include a copy of this form when sending your order/sample to Invitae. The Invitae billing team will follow up with BCBS FEP to ensure coverage. Thank you!

BCBS FEP Auth Department: Please fax your determination letter directly to Invitae at fax# 510-417-3932.

Prior Authorization Request Form Genetic Testing for Hereditary Breast and/or Ovarian Cancer

Notice: The Federal Employee Program has a 15 Day turn-around time on all Prior Authorization Requests According to the Blue Cross Blue Shield Service Benefit Plan Failure to complete this form in its entirety may result in delayed processing or an adverse determination for insufficient information.

Patient Information

Patient's Name:

Blue Cross Blue Shield ID Number:

Birth Date:

Patient's Phone Number:

Billing Provider Information

Ordering Physician/Provider Information

Name: Invitae Corporation
Address: 1400 16th Street
San Francisco, CA 94103

Please check this box if the ordering and billing provider are the same

Provider's Name:
Provider's Address:

Tax ID/NPI Number: NPI: 1316206220

Tax ID/NPI Number:

Office Contact: Patrick Hamm

Office Contact:

Phone: 510-417-3932

Phone:

Fax: 510-417-3932

Fax:

Please enter all codes requested; "by report" codes must have a description of why the code is being used.

ICD-10 CODE(S):

CPT/HCPC CODE(S):

PATIENT CLINICAL INFORMATION

Please provide the following documentation: Anticipated Date(s) of Service: _____

- History and physical and/or consultation notes including:
 - Ethnicity/Ancestry – Ashkenazi Jewish Heritage
 - Personal or family history of cancer (if applicable) including:
 - Family relationship(s): (maternal or paternal), (family member e.g., sibling, aunt, grandparent), (living or deceased) (if applicable)
 - Breast Cancer, Bilateral Breast Cancer, or Ovarian Cancer
 - Site(s) of cancer*
 - Age at diagnosis
 - BRCA1/BRCA2 mutation history (if applicable)
 - Genetic counseling/professional results (if applicable)
- Laboratory or Pathology reports (e.g., BRCA results for BART testing requests, or hormone receptor assay) (if applicable)
- Additionally, for BART Testing (please answer the following):

Did patient have BRCA testing prior to 2006 when BART testing was not available? Yes No

BRCA1 and BRCA 2 mutation testing results (if available/if applicable) Positive Negative

View our Medical Policy on line at <http://www.fepblue.org/medical-policies.jsp>

Fax Number: 1-855-895-3504

Phone Number: 1-800-633-4581

This facsimile transmission may contain protected and privileged, highly confidential medical, Personal and Health Information (PHI) and/or legal information. The information is intended only for the use of the individual or entity named above.

If you are not the intended recipient of this material, you may not use, publish, discuss, disseminate or otherwise distribute it. If you are not the intended recipient, or if you have received this transmission in error, please notify the sender immediately and **confidentially** destroy the information that faxed in error.

Thank you for your help in maintaining appropriate confidentiality.

Revised: Effective:

BCBS Federal Employee Program Statement of Medical Necessity Form	Genetic Testing for Hereditary Breast and/or Ovarian Cancer Syndrome (BRCA1/BRCA2)
Patient Information	Physician Information
Name: _____	Physician Name: _____
Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Tax ID/NPI Number: _____
BCBS Federal Employee Program ID Number: _____	Specialty: <input type="checkbox"/> PCP <input type="checkbox"/> Specialist <input type="checkbox"/> Genetic Counselor (Please Identify): _____
Birth Date (mm/dd/yy): _____	Rendering Provider (Laboratory): Invitae Corporation
Date of Service (mm/dd/yy): _____	Rendering Provider ID Number: NPI: 1316206220
Has the patient undergone prior BRCA testing? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Coding Information	
BRCA1 and BRCA2 Gene Analysis CPT Codes (check all that apply): <input type="checkbox"/> 81162 (BRCA1 & 2; Full sequence & full duplication/deletion variants) <input checked="" type="checkbox"/> 81211 (BRCA1 & 2; Full sequence & common duplication/deletion variants)	
ICD-10 Codes (list all applicable codes): _____	
Medical Necessity Criteria	
BRCA1 and BRCA2 Mutation Testing: Genetic testing for BRCA1 and BRCA2 mutations in adults (at least 18 years of age or older) may be considered medically necessary when any of the following criteria are met (please check all applicable boxes):	
<input type="checkbox"/> Individual (male or female) from a family with a known deleterious BRCA1/BRCA2 mutation; List mutation (Identify gene): _____	
<input type="checkbox"/> Personal history of breast cancer (including invasive and ductal carcinoma <i>in situ</i>) and one or more of the following circumstances: <input type="checkbox"/> Diagnosed at age 45 or younger <input type="checkbox"/> Two breast primaries when first breast cancer diagnosis occurred on or before 50 years of age (includes bilateral [contralateral] disease or cases where there are two or more clearly separate [ipsilateral] primary tumors either synchronously or asynchronously) <input type="checkbox"/> Diagnosed on or before 50 years of age with one or more close blood relative with breast cancer at any age or with a limited family history <input type="checkbox"/> Diagnosed on or before 60 years of age with triple negative breast cancer <input type="checkbox"/> Diagnosed at any age with one or more close blood relatives with breast cancer diagnosed on or before 50 years of age <input type="checkbox"/> Diagnosed at any age with two or more close blood relatives with breast cancer at any age <input type="checkbox"/> Diagnosed at any age with one or more close blood relatives with epithelial ovarian cancer, fallopian tube, or primary peritoneal cancer <input type="checkbox"/> Diagnosed at any age with two or more close blood relatives with pancreatic cancer or aggressive prostate cancer (Gleason score ≥ 7) at any age <input type="checkbox"/> Close male blood relative with breast cancer <input type="checkbox"/> Adult individual of ethnicity associated with deleterious founder mutations (e.g., Ashkenazi Jewish descent); <i>Please indicate maternal or paternal family member/s:</i> _____	
<input type="checkbox"/> Personal history of epithelial ovarian, fallopian tube or primary peritoneal cancer	
<input type="checkbox"/> Personal history of male breast cancer at any age	
<input type="checkbox"/> Personal history of pancreatic cancer or aggressive prostate cancer (Gleason score > 7) at any age with two or more close blood relatives with breast and/or ovarian cancer (including fallopian tube or primary peritoneal cancer), and/or pancreatic or aggressive prostate cancer (Gleason score > 7) at any age	
<input type="checkbox"/> An adult without a personal history, but with a family history only of a:	
<input type="checkbox"/> First- or second-degree blood relative meeting any of the above criteria (Please indicate maternal or paternal family member & age) <i>Family Member:</i> _____ <i>Age:</i> _____ (years) <i>Tumor type:</i> _____	
<input type="checkbox"/> A third-degree blood relative with both of the following: · Breast cancer (including invasive and ductal carcinoma <i>in situ</i>) and/or ovarian or fallopian tube or primary peritoneal cancer; and · At least two close blood relatives with breast cancer (at least one with breast cancer at age 50 or younger) and/or ovarian (including) fallopian tube or primary peritoneal cancer <i>Family Member:</i> _____ <i>Age:</i> _____ (years) <i>Tumor type:</i> _____ <i>Family Member:</i> _____ <i>Age:</i> _____ (years) <i>Tumor type:</i> _____ <i>Family Member:</i> _____ <i>Age:</i> _____ (years) <i>Tumor type:</i> _____	
I confirm that this test is medically necessary in accordance with (BCBS FEP) medical policy and that the information provided is accurate and factual based on the patient's medical records and history. I confirm that this test is medically necessary for the risk and assessment, diagnosis or detection of a disease, illness, impairment, symptom, syndrome or disorder. The results will determine my patient's medical management and treatment decisions. I confirm that I have been trained to provide genetic counseling, and that I have conducted a full personal and family history which includes a first, second and third degree pedigree. I confirm that I have provided genetic testing information and pre-test counseling to the patient and they have consented to genetic testing. I have scheduled post-test counseling to review the test results and determine future medical management and treatment plans.	
Ordering Provider Signature: _____ Date (mm/dd/yy): _____	
Please print name: _____ NPI number: _____	
Date of genetic counseling (mm/dd/yy): _____	