

Medical Necessity Form: Genetic Testing for Inherited BRCA1 or BRCA2



BlueCross BlueShield of Delaware

BCBSD requires that prior authorization for BRCA1 and BRCA2 genetic testing be obtained prior to ordering the test. In order for BCBSD to gather relevant medical information for review, providers must complete and sign the form below. Completed forms should be faxed to BCBSD's Medical Management Department at **302.421.8864** or **800.670.4862**.

Patient Information		
Patient Name		
Patient's Date of Birth	BCBSD Member ID Number	Proposed Date of Service
Physician and Genetic Counselor Information		
Ordering Physician Name	Phone Number	Fax Number
Rendering Physician Name	Phone Number	Fax Number
Procedure Code	Diagnosis Code(s)	
Genetic Counselor Name	Phone Number	Date of Visit
Outcome: Genetic Testing Recommended <input type="checkbox"/> Y <input type="checkbox"/> N Patient Requested Test <input type="checkbox"/> Y <input type="checkbox"/> N		

Please check Y to those that apply to the patient (personal history) and/or the patient's family (family history, on either the mother or father's side). If Y is checked, please also list the relationship to the patient of the individual diagnosed (e.g., self, maternal aunt, sister, paternal cousin) and her/his age at diagnosis.

Hereditary Breast and Ovarian Cancer Syndrome			
<input type="checkbox"/> Y <input type="checkbox"/> N	Biologically related individual from a family with a known BRCA1 or BRCA2 mutation	Relationship	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of breast cancer at or before age 45	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of breast cancer at or before age 50	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of breast cancer at any age	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of breast cancer and an individual of ethnicity associated with higher mutation frequency (e.g., Ashkenazi Jewish, Icelandic, Swedish, Hungarian or other)	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of epithelial ovarian, fallopian tube, or primary peritoneal cancer at any age	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of male breast cancer	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Family history only —meeting any of the above criteria	Relationship	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Personal history of triple negative breast cancer under age 60	Relationship Self	Age at Diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N	Individual with at least a 10% risk of carrying a BRCA mutation based on validated quantitative risk assessment tools (BRCAPRO, Yale, Univ of PA, BOADICEA and Tyrer-Cuzick)	Please send a pedigree and a copy of the models >10%	

Please provide any additional information regarding the reason for testing:

I confirm that information given on this form is accurate as of this date.

Signature of Physician or Authorized Representative

Date