

PATIENT INFORMATION

First name	MI	Last name	Date of birth (MM/DD/YYYY)
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Sex	MRN (medical record number)	Ancestry	
<input type="radio"/> M <input type="radio"/> F	<input type="text"/>	<input type="radio"/> Asian <input type="radio"/> Black/African American <input type="radio"/> White/Caucasian <input type="radio"/> Ashkenazi Jewish <input type="radio"/> Hispanic <input type="radio"/> Native American <input type="radio"/> Pacific Islander <input type="radio"/> French Canadian <input type="radio"/> Sephardic Jewish <input type="radio"/> Mediterranean <input type="radio"/> Other: _____	
Email address (for report access after release by medical professional)		Mobile phone	
<input type="text"/>		<input type="text"/>	
Address			
<input type="text"/>			
City	State	ZIP code	Country
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>

ORGANIZATION INFORMATION

Organization name and address			
Organization name			Phone
Address			Fax
City	State	ZIP code	Country
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Primary clinical contact			
Name			NPI
Email address (for report access)			Phone
Ordering physician			
<input type="radio"/> Same as primary clinical contact			
Name			NPI
Email address (for report access)			Phone
Additional clinical or laboratory contacts (optional)			
Name	Email address (for report access)	Name	Email address (for report access)
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Name	Email address (for report access)	Name	Email address (for report access)
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>

<input type="radio"/> INSURANCE BILLING (Please attach a copy of the patient's card.)	
We do not accept insurance for certain tests or patients outside the US. Before completing this section, confirm your test is eligible at www.invitae.com/billing.	
Primary insurance company name	Primary member ID#
<input type="text"/>	<input type="text"/>
Secondary insurance company name	Secondary member ID#
<input type="text"/>	<input type="text"/>
<input type="radio"/> Patient has Medicare and was treated as a hospital inpatient (>24 hour stay) in the last 14 days.	Prior-authorization #
<input type="text"/>	<input type="text"/>
Letter of Medical Necessity (LMN)	
<input type="radio"/> I have attached an LMN and/or other documents for insurance billing purposes.	
<input type="radio"/> I agree to allow Invitae to transfer the information from this requisition to an LMN and/or other documentation using the ordering physician's name as the signature for insurance billing.	

<input type="radio"/> INSTITUTIONAL BILLING
Invitae will send an invoice to the organization address above. Please contact Invitae if this order should be billed to a different location.

<input type="radio"/> PATIENT PAY BILLING
Invitae will send an electronic invoice to the patient email listed above

<input type="radio"/> OTHER BILLING
Invitae partner code:
<input type="text"/>

SPECIMEN INFORMATION

 Label each tube with the patient's full name, date of birth, and specimen collection date. A requisition form **MUST** accompany each specimen. www.invitae.com/specimen-requirements
Collection date (MM/DD/YYYY)
 / /

If not provided, date will be 1 day prior to our receipt of specimen. For DNA, provide date retrieved from archive.

Specimen type
 Blood Saliva DNA - source: _____

DNA must be extracted in a CLIA or other suitably certified laboratory. We are unable to accept blood or saliva from patients with allogeneic bone marrow transplants or a blood transfusion <2 weeks prior to specimen collection.

Special cases
 History of/current hematologic malignancy
 Resubmission

Deceased date (MM/DD/YYYY)
 / /
Specimen ID (IB # found on tube) - optional:
Is this patient deceased? Yes No

REASON FOR TESTING
ICD-10 codes
Previous results
Primary indication:
ONCOLOGY
 Hereditary breast and ovarian cancer (HBOC) syndrome
 Lynch syndrome
 Polyposis (FAP)
 Other: _____

CARDIOLOGY
 Aortopathy
 Arrhythmia
 Cardiomyopathy
 Other: _____

OTHER
 Neurology
 Other: _____

Testing for a personal history of disease? Yes No If yes, describe below.

Age at diagnosis: _____

Family history? Yes No If yes, describe in detail below and attach pedigree.

TEST SELECTION

 This form supports three different ways of indicating test selection. Please select **one** of the three options below.

OPTION 1: SELECT AN INVITAE PANEL FROM OUR TEST CATALOG

Select your desired test(s) from the attached test catalog and discard any pages without a selection.

OPTION 2: INVITAE TEST CODE

 Locate the test ID for the test you would like to order at www.invitae.com/tests or in our test catalog, and indicate the ID here.

Test code

Add-on code (optional)

OR

Invitae supports customization of your test. To create a custom panel, log in to your Invitae portal account or contact Client Services. Then indicate the ID associated with that panel here.

Custom panel ID

OPTION 3: FAMILY FOLLOW-UP TEST

 Learn more about Invitae's family follow-up test options at www.invitae.com/family.

Invitae proband RQ# _____

Relationship to proband _____

Gene(s) _____

Variant(s) _____

 Invitae's family follow-up testing analyzes the variant(s) indicated above. If you would like this report to include any variants of uncertain significance and be eligible for re-requisition, please include billing information on this requisition form and check here:

 To request a complimentary specimen collection kit visit www.invitae.com/request-a-kit
SHIPPING INSTRUCTIONS

Please ship specimen overnight in insulated containers:

Attn: Invitae Client Services, 1400 16th Street, San Francisco, CA 94103, USA

By signing this form, the medical professional acknowledges that the individual/family member authorized to make decisions for the individual (collectively, the "Patient") has been supplied information regarding and consented to undergo genetic testing, substantially as set forth in Invitae's Informed Consent for Genetic Testing (www.invitae.com/patient-consent), has been informed that Invitae may notify them of clinical updates related to genetic test results (in consultation with the ordering medical professional as indicated), and for orders originating outside the US, has been informed that the Patient's personal information and specimen will be transferred to and processed in the US. The Patient has further been informed and authorizes Invitae Corporation ("Invitae") and its designees to release information concerning testing to their insurer, if applicable, in order to process and/or appeal claims on behalf of the Patient. If a letter of medical necessity (LMN) has not been provided, the medical professional agrees to transfer the information from this requisition to a LMN and/or other documentation using the medical professional's name as the signature for insurance billing. For amounts received directly, the Patient has agreed to remit payment to Invitae for testing services rendered. I acknowledge that I offered pre-test genetic counseling to the Patient, if required by their insurer. In addition to the above, I attest that I am the ordering physician, or I am authorized by the ordering physician to order this test, or I am authorized under applicable law to order this test.

Medical professional signature
Date

HEREDITARY CANCER TEST CATALOG

All tests on this form fall into a single clinical area. If your order contains tests from multiple clinical areas, you will need to send in two sample tubes and your order will represent two billable events. Your test results will be delivered as two reports. Please contact Client Services with any questions. For Invitae's full test menu, please visit www.invitae.com.

FREQUENTLY-ORDERED PANELS

Test code	Test name	# gene(s)	Gene list
Cross-Cancer			
<input type="radio"/> 01101	Invitae Multi-Cancer Panel	83	ALK, APC, ATM, AXIN2, BAP1, BARD1, BLM, BMPR1A, BRCA1, BRCA2, BRIP1, CASR, CDC73, CDH1, CDK4, CDKN1B, CDKN1C, CDKN2A, CEBPA, CHEK2, CTNNA1, DICER1, DIS3L2, EGFR, EPCAM, FH, FLCN, GATA2, GPC3, GREM1, HOXB13, HRAS, KIT, MAX, MEN1, MET, MITF, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NF2, NTHL1, PALB2, PDGFRA, PHOX2B, PMS2, POLD1, POLE, POT1, PRKAR1A, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RB1, RECQL4, RET, RUNX1, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, STK11, SUFU, TERC, TERT, TMEM127, TP53, TSC1, TSC2, VHL, WRN, WT1
<input type="radio"/> 01102	Invitae Common Hereditary Cancers Panel	47	APC, ATM, AXIN2, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, CTNNA1, DICER1, EPCAM, GREM1, HOXB13, KIT, MEN1, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PDGFRA, PMS2, POLD1, POLE, PTEN, RAD50, RAD51C, RAD51D, SDHA, SDHB, SDHC, SDHD, SMAD4, SMARCA4, STK11, TP53, TSC1, TSC2, VHL

STAT Turnaround Time

SPECIAL INSTRUCTIONS: These panels have a guaranteed turnaround time of 5–12 calendar days from when the specimen is received. Genes cannot be removed and they cannot be ordered with any other non-STAT panels or genes. The option to re-requisition additional genes is available. Only blood and saliva are accepted (DNA is not accepted).

<input type="radio"/> 50001	Invitae Breast Cancer STAT Panel	7	BRCA1, BRCA2, CDH1, PALB2, PTEN, STK11, TP53
<input type="radio"/> 50001.1	Add-on ATM gene	1	ATM
<input type="radio"/> 50001.2	Add-on CHEK2 gene	1	CHEK2
<input type="radio"/> 50002	Invitae BRCA1 and BRCA2 STAT Panel	2	BRCA1, BRCA2

CANCER PANELS BY ORGAN SYSTEM

Test code	Test name	# gene(s)	Gene list
Breast Cancer			
<input type="radio"/> 01206	Invitae Breast Cancer Guidelines-Based Panel	11	ATM, BRCA1, BRCA2, CDH1, CHEK2, NBN, NF1, PALB2, PTEN, STK11, TP53
<input type="radio"/> 01206.1	Add-on gene with emerging data	1	BARD1
<input type="radio"/> 01202	Invitae Breast Cancer Panel	14	ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, NBN, NF1, PALB2, PTEN, RAD50, STK11, TP53
<input type="radio"/> 01202.1	Add-on preliminary-evidence genes	13	ABRAXAS1, AKT1, FANCC, FANCM, MRE11, MUTYH, PIK3CA, RAD51C, RAD51D, RINT1, SDHB, SDHD, XRCC2
Breast and Gynecologic Cancers			
<input type="radio"/> 01204	Invitae Breast and Gyn Cancers Guidelines-Based Panel	19	ATM, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, EPCAM, MLH1, MSH2, MSH6, NBN, NF1, PALB2, PMS2, PTEN, RAD51C, RAD51D, STK11, TP53
<input type="radio"/> 01204.1	Add-on gene with emerging data	1	BARD1
<input type="radio"/> 01201	Invitae Breast and Gyn Cancers Panel	23	ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, DICER1, EPCAM, MLH1, MSH2, MSH6, NBN, NF1, PALB2, PMS2, PTEN, RAD50, RAD51C, RAD51D, SMARCA4, STK11, TP53
<input type="radio"/> 01201.1	Add-on preliminary-evidence genes	13	ABRAXAS1, AKT1, CDC73, FANCC, FANCM, MRE11, MUTYH, PIK3CA, POLD1, RINT1, SDHB, SDHD, XRCC2

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HEREDITARY CANCER TEST CATALOG

CANCER PANELS BY ORGAN SYSTEM (continued)

Test code	Test name	# gene(s)	Gene list
Colorectal Cancer			
<input type="radio"/> 01252	Invitae Colorectal Cancer Guidelines-Based Panel	19	APC, AXIN2, BMPR1A, CHEK2, EPCAM, GREM1, MLH1, MSH2, MSH3, MSH6, MUTYH, NTHL1, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53
<input type="radio"/> 01252.1	Add-on gene with emerging data	1	RPS20
<input type="radio"/> 01251	Invitae Colorectal Cancer Panel	20	APC, AXIN2, BMPR1A, CDH1, CHEK2, EPCAM, GREM1, MLH1, MSH2, MSH3, MSH6, MUTYH, NTHL1, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53
<input type="radio"/> 01251.1	Add-on preliminary-evidence genes	9	ATM, BLM, BUB1B, CEP57, ENG, FLCN, GALNT12, MLH3, RPS20
Additional Organ Systems			
<input type="radio"/> 01271	Invitae Gastric Cancer Panel	19	APC, BMPR1A, CDH1, CTNNA1, EPCAM, KIT, MLH1, MSH2, MSH6, NF1, PDGFRA, PMS2, SDHA, SDHB, SDHC, SDHD, SMAD4, STK11, TP53
<input type="radio"/> 01302	Invitae Hereditary Paraganglioma-Pheochromocytoma Panel	10	MAX, NF1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127, VHL
<input type="radio"/> 01302.1	Add-on preliminary-evidence genes	4	EGLN1, FH, KIF1B, MEN1
<input type="radio"/> 01303	Invitae Hyperparathyroidism Panel	5	CASR, CDC73, CDKN1B, MEN1, RET
<input type="radio"/> 01561	Invitae Melanoma Panel	9	BAP1, BRCA2, CDK4, CDKN2A, MITF, POT1, PTEN, RB1, TP53
<input type="radio"/> 01561.1	Add-on preliminary-evidence genes	3	BRCA1, MC1R, TERT
<input type="radio"/> 01411	Invitae Myelodysplastic Syndrome/Leukemia Panel	16	ATM, BLM, CEBPA, EPCAM, GATA2, HRAS, MLH1, MSH2, MSH6, NBN, NF1, PMS2, RUNX1, TERC, TERT, TP53
<input type="radio"/> 01411.1	Add-on preliminary-evidence genes	5	BRCA1, BRCA2, BRIP1, CHEK2, PALB2
<input type="radio"/> 01411.2	Add-on dyskeratosis congenita genes	7	CTC1, DKC1, NHP2, NOP10, TERC, TERT, TINF2
<input type="radio"/> 01411.3	Add-on Fanconi anemia genes	17	BRCA2, BRIP1, ERCC4, FANCA, FANCB, FANCC, FANCD2, FANCE, FANCF, FANCG, FANCI, FANCL, FANCM, PALB2, RAD51C, SLX4, XRCC2
<input type="radio"/> 01461	Invitae Nervous System/Brain Cancer Panel	25	ALK, APC, DICER1, EPCAM, HRAS, MEN1, MLH1, MSH2, MSH6, NF1, NF2, PHOX2B, PMS2, PRKAR1A, PTCH1, PTEN, RB1, SMARCA4, SMARCB1, SMARCE1, SUFU, TP53, TSC1, TSC2, VHL
<input type="radio"/> 01461.1	Add-on preliminary-evidence genes	7	BAP1, BARD1, EZH2, GPC3, KIF1B, POT1, PTCH2
<input type="radio"/> 01461.2	Add-on hereditary paraganglioma pheochromocytoma genes	8	MAX, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127
<input type="radio"/> 01261	Invitae Pancreatic Cancer Panel	20	APC, ATM, BMPR1A, BRCA1, BRCA2, CDKN2A, EPCAM, MEN1, MLH1, MSH2, MSH6, NF1, PALB2, PMS2, SMAD4, STK11, TP53, TSC1, TSC2, VHL
<input type="radio"/> 01261.1	Add-on preliminary-evidence genes	3	CDK4, FANCC, PALLD
<input type="radio"/> 01261.2	Add-on chronic pancreatitis genes	5	CASR, CFTR, CTRC, PRSS1, SPINK1
<input type="radio"/> 01362	Invitae Prostate Cancer Panel	12	ATM, BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PMS2, TP53
<input type="radio"/> 01362.1	Add-on preliminary-evidence genes	5	BRIP1, FANCA, PALB2, RAD51C, RAD51D
<input type="radio"/> 01361	Invitae Renal/Urinary Tract Cancers Panel	24	BAP1, CDC73, CDKN1C, DICER1, DIS3L2, EPCAM, FH, FLCN, GPC3, MET, MLH1, MSH2, MSH6, PMS2, PTEN, SDHB, SDHC, SMARCA4, SMARCB1, TP53, TSC1, TSC2, VHL, WT1
<input type="radio"/> 01361.1	Add-on preliminary-evidence genes	6	BUB1B, CEP57, MITF, PALB2, SDHA, SDHD

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HEREDITARY CANCER TEST CATALOG

CANCER PANELS BY ORGAN SYSTEM (continued)

Test code	Test name	# gene(s)	Gene list
Additional Organ Systems (continued)			
<input type="radio"/> 01511	Invitae Sarcoma Panel	26	APC, BLM, CDKN1C, DICER1, EPCAM, FH, HRAS, KIT, MLH1, MSH2, MSH6, NBN, NF1, PDGFRA, PMS2, PRKAR1A, PTCH1, RB1, RECQL4, SDHA, SDHB, SDHC, SDHD, SUFU, TP53, WRN
<input type="radio"/> 01511.1	Add-on preliminary-evidence genes	5	CDKN2A, POT1, PTCH2, TSC1, TSC2
<input type="radio"/> 01511.2	Add-on Diamond-Blackfan anemia genes	10	GATA1, RPL11, RPL26, RPL35A, RPL5, RPS10, RPS19, RPS24, RPS26, RPS7
<input type="radio"/> 01301	Invitae Thyroid Cancer Panel	7	APC, CHEK2, DICER1, PRKAR1A, PTEN, RET, TP53
<input type="radio"/> 01301.1	Add-on preliminary-evidence genes	4	MEN1, SDHB, SDHD, WRN
Pediatric Oncology			
<input type="radio"/> 01104	Invitae Pediatric Solid Tumors Panel	48	ALK, APC, AXIN2, BAP1, BLM, BMPR1A, CDC73, CDKN1C, DICER1, DIS3L2, EPCAM, FH, GPC3, HRAS, MAX, MEN1, MLH1, MSH2, MSH6, NBN, NF1, NF2, PHOX2B, PMS2, PRKAR1A, PTCH1, PTEN, RB1, RECQL4, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, SMARCB1, SMARCE1, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, WRN, WT1
<input type="radio"/> 01105	Invitae Pediatric Hematologic Malignancies Panel	16	ATM, BLM, CEBPA, EPCAM, GATA2, HRAS, MLH1, MSH2, MSH6, NBN, NF1, PMS2, RUNX1, TERC, TERT, TP53
<input type="radio"/> 01106	Invitae Pediatric Nervous System/Brain Tumors Panel	24	ALK, APC, DICER1, EPCAM, HRAS, MEN1, MLH1, MSH2, MSH6, NF1, NF2, PHOX2B, PMS2, PRKAR1A, PTCH1, PTEN, RB1, SMARCB1, SMARCE1, SUFU, TP53, TSC1, TSC2, VHL
<input type="radio"/> 01106.1	Hereditary paraganglioma pheochromocytoma genes	8	MAX, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127
Individual Hereditary Cancer Conditions			
<input type="radio"/> 01724	Invitae Ataxia-Telangiectasia Test	1	ATM
<input type="radio"/> 01728	Invitae BAP1 Hereditary Cancer Predisposition Syndrome Test	1	BAP1
<input type="radio"/> 01722	Invitae Basal Cell Nevus Syndrome Panel	2	PTCH1, SUFU
<input type="radio"/> 01722.1	Add-on preliminary-evidence gene	1	PTCH2
<input type="radio"/> 01720	Invitae Birt-Hogg-Dubé Syndrome Test	1	FLCN
<input type="radio"/> 01730	Invitae Bloom Syndrome Test	1	BLM
<input type="radio"/> 01731	Invitae Carney Complex Test	1	PRKAR1A
<input type="radio"/> 01732	Invitae CASR-Related Conditions Test	1	CASR
<input type="radio"/> 01729	Invitae CDC73-Related Conditions Test	1	CDC73
<input type="radio"/> 01745	Invitae Chronic Pancreatitis Panel	5	CASR, CFTR, CTRC, PRSS1, SPINK1
<input type="radio"/> 01703	Invitae Constitutional Mismatch Repair-Deficiency Panel	5	EPCAM, MLH1, MSH2, MSH6, PMS2
<input type="radio"/> 04164	Invitae Costello Syndrome Test	1	HRAS
<input type="radio"/> 01719	Invitae DICER1 Syndrome Test	1	DICER1
<input type="radio"/> 01744	Invitae Familial Acute Myeloid Leukemia with Mutated CEBPA Test	1	CEBPA
<input type="radio"/> 01709	Invitae Familial Adenomatous Polyposis Test	1	APC
<input type="radio"/> 01712	Invitae Familial Gastrointestinal Stromal Tumor Syndrome Panel	7	KIT, NF1, PDGFRA, SDHA, SDHB, SDHC, SDHD

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HEREDITARY CANCER TEST CATALOG

CANCER PANELS BY ORGAN SYSTEM (continued)

Test code	Test name	# gene(s)	Gene list
Individual Hereditary Cancer Conditions (continued)			
○ 01734	Invitae Familial Platelet Disorder with Propensity to Myeloid Malignancy Test	1	RUNX1
○ 05311	Invitae Fanconi Anemia Panel	17	BRCA2, BRIP1, ERCC4, FANCA, FANCB, FANCC, FANCD2, FANCE, FANCF, FANCG, FANCI, FANCL, FANCM, PALB2, RAD51C, SLX4, XRCC2
○ 05317	Invitae GATA2 Deficiency Test	1	GATA2
○ 01701	Invitae Hereditary Breast and Ovarian Cancer Syndrome Panel	2	BRCA1, BRCA2
○ 01707	Invitae Hereditary Diffuse Gastric Cancer Syndrome Test	1	CDH1
○ 01727	Invitae Hereditary Leiomyomatosis and Renal Cell Cancer Test	1	FH
○ 01723	Invitae Hereditary Papillary Renal Cell Carcinoma Test	1	MET
○ 01711	Invitae Juvenile Polyposis Syndrome Panel	2	BMPR1A, SMAD4
○ 01705	Invitae Li-Fraumeni Syndrome Test	1	TP53
○ 01702	Invitae Lynch Syndrome Panel	5	EPCAM, MLH1, MSH2, MSH6, PMS2
○ 01713	Invitae Melanoma-Pancreatic Cancer Syndrome Panel	2	CDK4, CDKN2A
○ 01717	Invitae Multiple Endocrine Neoplasia Type 1 Test	1	MEN1
○ 01718	Invitae Multiple Endocrine Neoplasia Type 2 Test	1	RET
○ 01710	Invitae MUTYH-Associated Polyposis Syndrome Test	1	MUTYH
○ 01708	Invitae Neurofibromatosis Type 1 Test	1	NF1
○ 01708.1	Add-on Legius syndrome gene	1	SPRED1
○ 04167	Invitae Neurofibromatosis Type 2 Test	1	NF2
○ 04167.1	Add-on schwannomatosis gene	1	SMARCB1
○ 01725	Invitae Nijmegen Breakage Syndrome Test	1	NBN
○ 01726	Invitae Oligodontia-Colorectal Cancer Syndrome Test	1	AXIN2
○ 01736	Invitae Perlman Syndrome Test	1	DIS3L2
○ 01706	Invitae Peutz-Jeghers Syndrome Test	1	STK11
○ 01704	Invitae PTEN-Related Disorders Test	1	PTEN
○ 01737	Invitae RECQL4-Related Disorders Test	1	RECQL4
○ 01738	Invitae Retinoblastoma Test	1	RB1
○ 01714	Invitae Rhabdoid Tumor Predisposition Syndrome Panel	2	SMARCA4, SMARCB1
○ 04168	Invitae Schwannomatosis Test	1	SMARCB1
○ 04168.1	Add-on neurofibromatosis type 2 gene	1	NF2
○ 01739	Invitae Simpson-Golabi-Behmel Syndrome Test	1	GPC3
○ 01715	Invitae Small Cell Carcinoma of the Ovary Hypercalcemic Type Test	1	SMARCA4
○ 01721	Invitae Tuberous Sclerosis Complex Panel	2	TSC1, TSC2

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HEREDITARY CANCER TEST CATALOG

CANCER PANELS BY ORGAN SYSTEM (continued)

Test code	Test name	# gene(s)	Gene list
Individual Hereditary Cancer Conditions (continued)			
<input type="radio"/> 01716	Invitae von Hippel-Lindau Syndrome Test	1	VHL
<input type="radio"/> 01740	Invitae Weaver Syndrome Test	1	EZH2
<input type="radio"/> 01741	Invitae Werner Syndrome Test	1	WRN
<input type="radio"/> 01742	Invitae Wilms Tumor Panel	5	CDC73, CDKN1C, DIS3L2, GPC3, WT1
<input type="radio"/> 01743	Invitae WT1-Related Disorders Test	1	WT1

HEREDITARY CANCER INDIVIDUAL GENES

<input type="radio"/>	ABRAXAS1	<input type="radio"/>	CDK4	<input type="radio"/>	EPCAM	<input type="radio"/>	GATA2	<input type="radio"/>	MSH3	<input type="radio"/>	POT1	<input type="radio"/>	RPS10	<input type="radio"/>	SPINK1
<input type="radio"/>	AKT1	<input type="radio"/>	CDKN1B	<input type="radio"/>	ERCC4	<input type="radio"/>	GPC3	<input type="radio"/>	MSH6	<input type="radio"/>	PRKAR1A	<input type="radio"/>	RPS19	<input type="radio"/>	SPRED1
<input type="radio"/>	ALK	<input type="radio"/>	CDKN1C	<input type="radio"/>	EZH2	<input type="radio"/>	GREM1	<input type="radio"/>	MUTYH	<input type="radio"/>	PRSS1	<input type="radio"/>	RPS20	<input type="radio"/>	STK11
<input type="radio"/>	APC	<input type="radio"/>	CDKN2A	<input type="radio"/>	FANCA	<input type="radio"/>	HOXB13	<input type="radio"/>	NBN	<input type="radio"/>	PTCH1	<input type="radio"/>	RPS24	<input type="radio"/>	SUFU
<input type="radio"/>	ATM	<input type="radio"/>	CEBPA	<input type="radio"/>	FANCB	<input type="radio"/>	HRAS	<input type="radio"/>	NF1	<input type="radio"/>	PTCH2	<input type="radio"/>	RPS26	<input type="radio"/>	TERC
<input type="radio"/>	AXIN2	<input type="radio"/>	CEP57	<input type="radio"/>	FANCC	<input type="radio"/>	KIF1B	<input type="radio"/>	NF2	<input type="radio"/>	PTEN	<input type="radio"/>	RPS7	<input type="radio"/>	TERT
<input type="radio"/>	BAP1	<input type="radio"/>	CFTR	<input type="radio"/>	FANCD2	<input type="radio"/>	KIT	<input type="radio"/>	NHP2	<input type="radio"/>	RAD50	<input type="radio"/>	RUNX1	<input type="radio"/>	TINF2
<input type="radio"/>	BARD1	<input type="radio"/>	CHEK2	<input type="radio"/>	FANCE	<input type="radio"/>	MAX	<input type="radio"/>	NOP10	<input type="radio"/>	RAD51C	<input type="radio"/>	SDHA	<input type="radio"/>	TMEM127
<input type="radio"/>	BLM	<input type="radio"/>	CTC1	<input type="radio"/>	FANCF	<input type="radio"/>	MC1R	<input type="radio"/>	NTHL1	<input type="radio"/>	RAD51D	<input type="radio"/>	SDHAF2	<input type="radio"/>	TP53
<input type="radio"/>	BMPR1A	<input type="radio"/>	CTNNA1	<input type="radio"/>	FANCG	<input type="radio"/>	MDM2	<input type="radio"/>	PALB2	<input type="radio"/>	RB1	<input type="radio"/>	SDHB	<input type="radio"/>	TSC1
<input type="radio"/>	BRCA1	<input type="radio"/>	CTRC	<input type="radio"/>	FANCI	<input type="radio"/>	MEN1	<input type="radio"/>	PALLD	<input type="radio"/>	RECQL4	<input type="radio"/>	SDHC	<input type="radio"/>	TSC2
<input type="radio"/>	BRCA2	<input type="radio"/>	DICER1	<input type="radio"/>	FANCL	<input type="radio"/>	MET	<input type="radio"/>	PDGFRA	<input type="radio"/>	RET	<input type="radio"/>	SDHD	<input type="radio"/>	VHL
<input type="radio"/>	BRIP1	<input type="radio"/>	DIS3L2	<input type="radio"/>	FANCM	<input type="radio"/>	MITF	<input type="radio"/>	PHOX2B	<input type="radio"/>	RINT1	<input type="radio"/>	SLX4	<input type="radio"/>	WRN
<input type="radio"/>	BUB1B	<input type="radio"/>	DKC1	<input type="radio"/>	FH	<input type="radio"/>	MLH1	<input type="radio"/>	PIK3CA	<input type="radio"/>	RPL11	<input type="radio"/>	SMAD4	<input type="radio"/>	WT1
<input type="radio"/>	CASR	<input type="radio"/>	EGFR	<input type="radio"/>	FLCN	<input type="radio"/>	MLH3	<input type="radio"/>	PMS2	<input type="radio"/>	RPL26	<input type="radio"/>	SMARCA4	<input type="radio"/>	XRCC2
<input type="radio"/>	CDC73	<input type="radio"/>	EGLN1	<input type="radio"/>	GALNT12	<input type="radio"/>	MRE11	<input type="radio"/>	POLD1	<input type="radio"/>	RPL35A	<input type="radio"/>	SMARCB1		
<input type="radio"/>	CDH1	<input type="radio"/>	ENG	<input type="radio"/>	GATA1	<input type="radio"/>	MSH2	<input type="radio"/>	POLE	<input type="radio"/>	RPL5	<input type="radio"/>	SMARCE1		

Invitae continually updates its panels based on the most recent evidence. Please note that if an order is placed using an older version of this form, Invitae reserves the right to upgrade any ordered panel(s) to the current version(s). To avoid confusion, please consider placing your order using our online test catalog.