

Hereditary Cancer Complete Genetic Testing Menu

Comprehensive and Guideline-Based Hereditary Cancer panels

Test code	Test name	Genes
38600	Comprehensive Hereditary Cancer Panel (66 genes)	APC, ATM, AXIN2, BAP1, BARD1, BLM, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN1B, CDKN2A (p16, p14), CHEK2, DICER1, EGFR, EPCAM, FANCA, FANCC, FANCM, FH, FLCN, GALNT12, GREM1, HOXB13, MAX, MEN1, MET, MITF, MLH1, MRE11 (MRE11A), MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, POT1, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RECQL, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2
38611	Guideline-Based Hereditary Cancer Panel (32 genes)	APC, ATM, AXIN2, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A (p16,p14), CHEK2, EPCAM, GREM1, HOXB13, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, SMAD4, STK11, TP53

Cancer-specific panels

Test code	Test name	Genes
38621	Hereditary Breast Cancer Panel (16 genes)	ATM, BRCA1, BRCA2, CDH1, CHEK2, EPCAM, MLH1, MSH2, MSH6, NBN, NF1, PALB2, PMS2, PTEN, STK11, TP53
38631	Hereditary Colorectal Cancer Panel (19 genes)	APC, AXIN2, BMPR1A, CDH1, EPCAM, GREM1, MLH1, MSH2, MSH3, MSH6, MUTYH, NTHL1, PMS2, POLD1, POLE, PTEN, SMAD4, STK11, TP53
38641	Hereditary Endocrine Cancer Panel (12 genes)	FH, MAX, MEN1, NF1, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, TMEM127, VHL
92587	BRCA Panel Plus (7 genes)	BRCA1, BRCA2, CDH1, PALB2, PTEN, STK11, TP53

Hereditary cancer syndrome tests

Test code	Test name	Genes
Breast cancer syndrome		
91863	BRCA Panel	BRCA1, BRCA2
91864	BRCA Ashkenazi Jewish Screen	Common founder variants BRCA1 c.68_69delAG, BRCA1 c.5266dupC, BRCA2 c.5946delT
92140	BRCA Ashkenazi Jewish Screen w/Reflex to BRCA Panel (BRCA1, BRCA2)	Ashkenazi Jewish screen; if negative reflex to BRCA Panel (BRCA1, BRCA2)
91866	BRCA1 and BRCA2 Deletion and Duplication	This test detects large deletion/duplication mutations in the BRCA1 and BRCA2 genes which are not detectable by DNA sequencing
Lynch syndrome		
91461	Lynch Syndrome Panel (5 genes)	EPCAM, MLH1, MSH2, MSH6, PMS2
Other cancer risk		
38651	Nevoid Basal Cell Carcinoma (NBCCS) (Gorlin) Syndrome Panel (PTCH1, SUFU)	PTCH1, SUFU
38661	Tuberous Sclerosis Complex Panel (TSC1, TSC2)	TSC1, TSC2
94053	Juvenile Polyposis Panel (BMPR1A and SMAD4)	BMPR1A, SMAD4
93945	Hereditary Cancer Single Site(s)	APC, ATM, AXIN2, BAP1, BARD1, BLM, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN1B, CDKN2A (p16, p14), CHEK2, DICER1, EGFR, EPCAM, FANCA, FANCC, FANCM, FH, FLCN, GALNT12, GREM1, HOXB13, MAX, MEN1, MET, MITF, MLH1, MRE11 (MRE11A), MSH2, MSH3, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, POT1, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RECQL, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SMAD4, SMARCA4, STK11, SUFU, TMEM127, TP53, TSC1, TSC2, VHL, XRCC2
See genes column	Single-gene tests	APC – 93797, ATM – 38802, BAP1 – 38803, BLM – 38804, CDH1 – 92568, CDKN2A – 93939, CHEK2 – 93940, EPCAM/MSH2 – 91471, FH – 38805, FLCN – 38806, HOXB13 – 38807, MEN1 – 93942, MITF – 38808, MLH1 – 91460, MSH6 – 91458, MUTYH – 93944, NF1 – 93941, PALB2 – 92571, PMS2 – 91457, PTEN – 92566, RET – 93796, SMARCA4 – 38809, STK11 – 92565, TP53 – 92560, VHL – 93943

Components of panels may be ordered separately.

Please contact Quest Genomics Client Services at 1.866.GENE.INFO (1.866.436.3463) or visit QuestHereditaryCancer.com with questions