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## 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 |

Additional hereditary cancer risk tests

| 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 |
| 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) |
| 91461 | Lynch Syndrome Panel (5 genes) | EPCAM, MLH1, MSH2, MSH6, PMS2 |
| 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, <br> CDKN2A - 93939, CHEK2 - 93940, EPCAM/MSH2 - 91471, FH - 38805, FLCN - 38806, <br> HOXB13 - 38807, MEN1 - 93942, MITF - 38808, MLH1 - 91460, MSH6 - 91458, <br> MUTYH - 93944, NF1 - 93941, PALB2 - 92571, PMS2 - 91457, PTEN - 92566, RET - 93796, <br> SMARCA4 - 38809, STK11 - 92565, TP53 - 92560, VHL - 93943 |

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


[^0]:    Components of panels may be ordered separately

