Genetic Testing: Molecular Profiling Assays for Cancer Management

These services may or may not be covered by your HealthPartners plan. Please see your plan documents for your specific coverage information. If there is a difference between this general information and your plan documents, your plan documents will be used to determine your coverage.

Administrative Process
Prior authorization is required for molecular profiling assays for cancer management.

The scope of this coverage policy includes, but is not limited to, these services:

- Section 1: Cytogenetic and cytogenomic studies
- Section 2: Microsatellite instability (MSI) and mismatch repair (MMR) testing
- Section 3: Single-gene testing and targeted multiple-gene testing
- Section 4: Expanded genetic cancer marker testing
- Section 5: Testing for chimerism

Molecular profiling assays for cancer management other than as described below are also subject to a review for medical necessity, based on current clinical literature and expert recommendations.

Coverage

Indications that are covered

Section 1

Cytogenetic and cytogenomic studies

Cytogenetic and cytogenomic studies for these conditions, including, but not limited to, fluorescence in situ hybridization (FISH) and karyotype, are covered when criteria 1-2 below are met:

1. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must order the test.

Section 2

Microsatellite instability (MSI) testing and mismatch repair (MMR) testing

Microsatellite instability (MSI) and mismatch repair (MMR) testing for these conditions are covered when criteria 1-2 below are met:

1. The test is expected to directly impact management of the condition being evaluated.
2. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must order the test.
• Lung cancer
• Leukemia
• Lymphoma
• Melanoma
• Thyroid carcinoma
• Other hematological malignancies, lymphoproliferative disorders, and myelodysplastic/myeloproliferative conditions

The test is expected to directly impact management of the condition being evaluated.

1. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must order the test.

Section 3

Single-gene testing and targeted multiple-gene testing

Single-gene testing and targeted multiple-gene testing for these conditions are covered when criteria 1-3 below are met:

• Breast cancer (except for ductal carcinoma in situ [DCIS])
• Colorectal cancer (BRAF, KRAS, NRAS)
• Esophageal cancer
• Ewing sarcoma
• Gastrointestinal stromal tumor (GIST) (BRAF, KIT, PDGFRα, SDH)
• Glioblastoma multiforme
• Non-small cell lung cancer (NSCLC) (ALK, EGFR, KRAS)
• Leukemia
• Lymphoma
• Melanoma (BRAF, KIT)
• Thyroid carcinoma
• Other hematological malignancies, lymphoproliferative disorders, and myelodysplastic/myeloproliferative conditions

1. The test is expected to directly impact management of the condition being evaluated.
2. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must order the test.
3. The test does not include additional genes or analytes (genetic or molecular substances) that are not included in the list above, next to the condition being evaluated.

Section 4

Expanded genetic cancer marker testing

Afirma™ Genomic Sequencing Classifier (Veracyte, Inc.)

Afirma Genomic Sequencing Classifier is covered when criteria 1-3 below are met:

1. The test is expected to directly impact management of a thyroid nodule.
2. The ordering physician or healthcare provider must certify that the thyroid nodule being tested meets both of these criteria:
   A. The nodule has been labeled “indeterminate” during laboratory examination (atypia or lesion of undetermined significance), so that cancer is still considered to be a possible diagnosis.
   B. The nodule has a diameter of at least 1.0 cm.
3. The member is age 18 or older.

EndoPredict® (Myriad Genetics Laboratories, Inc.)

EndoPredict is covered when criteria 1-3 below are met:

1. The test is expected to directly impact management of breast cancer.
2. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must:
   A. Order the test.
   B. Certify that the test results will be used when making a decision about adjuvant chemotherapy treatment (chemotherapy that is given after surgery).
3. Previous test results must show all of the following:
   A. The tumor is estrogen receptor (ER) positive.
The tumor is human epidermal growth factor receptor 2 (HER2) negative.

B. The tumor is human epidermal growth factor receptor 2 (HER2) negative.

C. Axillary lymph nodes (located in the armpits) are free of cancer.

MammaPrint® (Agendia, Inc.)
(Each individual tumor being tested must independently meet these criteria.)

MammaPrint is covered when criteria 1-3 below are met:

1. The test is expected to directly impact management of breast cancer.
2. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must:
   A. Order the test.
   B. Certify that the test results will be used when making a decision about adjuvant chemotherapy treatment (chemotherapy that is given after surgery).
3. Previous test results must show all of the following:
   A. The tumor is estrogen receptor (ER) positive.
   B. The tumor is human epidermal growth factor receptor 2 (HER2) negative.
   C. Axillary lymph nodes (located in the armpits) are free of cancer.
   D. The tumor has a diameter between 0.5-2.0 cm.

Oncotype DX Breast Recurrence Score® (Genomic Health, Inc.)
(Each individual tumor being tested must independently meet these criteria.)

Oncotype DX Breast Recurrence Score is covered when criteria 1-4 below are met:

1. The test is expected to directly impact management of breast cancer.
2. A board-certified physician or advanced-practice registered nurse in pathology, endocrinology, genetics, oncology, or hematology (who is not affiliated with the commercial testing laboratory) must:
   A. Order the test.
   B. Certify that the test results will be used when making a decision about treatment with hormone therapy and adjuvant chemotherapy treatment (chemotherapy that is given after surgery).
3. Previous test results must show all of the following:
   A. The tumor is estrogen receptor (ER) or progesterone receptor (PR) positive.
   B. There is no evidence that the cancer has spread to distant organs or lymph nodes (distant metastasis).
   C. There are no more than three axillary lymph nodes (located in the armpits) that have tested positive for cancer.
4. Previous test results must show one of the following:
   A. The tumor is human epidermal growth factor receptor 2 (HER2) negative, regardless of tumor size.
   B. The tumor is HER2 positive and has a diameter less than 1.0 cm.

Section 5

Testing for chimerism

Testing for chimerism, including, but not limited to, comparative analysis with short tandem repeat (STR) markers and testing single nucleotide polymorphisms (SNP), is covered when criteria 1-2 below are met:

1. The test is expected to directly impact management of a malignant disease (cancer).
2. A board-certified physician or advanced-practice registered nurse in pathology, genetics, oncology, hematology, or immunology (who is not affiliated with the commercial testing laboratory) must:
   A. Order the test.
   B. Recommend the test for one of the following indications:
      • The member has a personal history of hematopoietic cell transplantation (HCT).
      • The member has a personal history of donor lymphocyte infusion (DLI).
      • The member has a personal history of immunomodulatory cytokine therapy or other cellular therapy.
      • The member is expected to receive HCT, DLI, immunomodulatory cytokine therapy, or other cellular therapy in the immediate future.

Indications that are not covered

1. The following services are considered experimental/investigational because reliable evidence does not permit conclusions concerning safety, effectiveness, or effect on health outcomes:
   A. Expanded genetic cancer marker testing, except as described in Section 4 (also, see Definitions)
   B. Genomic microarray testing for hematological malignancies
   C. MicroRNA testing
   D. Whole exome sequencing
   E. Whole genome sequencing
F. Proteomic pattern testing (see Definitions), including, but not limited to:
   - BDX-XL2® (Biodesix, Inc.)
   - CxBladder™ Detect (Pacific Edge Diagnostics USA, Ltd.)
   - CxBladder™ Monitor (Pacific Edge Diagnostics USA, Ltd.)
   - VeriStrat® (Biodesix, Inc.)
G. Testing of liquid biopsies (see Definitions), including, but not limited to:
   - GeneStrat® (Biodesix, Inc.)
   - Guardant360® (Guardant Health, Inc.)
   - Testing for circulating free DNA (cfDNA)
   - Testing for circulating tumor DNA (ctDNA)
   - SelectMDx® (MDxHealth SA)
H. Topographic genotyping (see Definitions), including, but not limited to:
   - BarreGEN® (Interpace Diagnostics Group, LLC)
   - RespriDx™ (Interpace Diagnostics Group, LLC)
   - PancraGEN® (Interpace Diagnostics Group, LLC)
I. Testing for the following indications:
   - Anal carcinoma
   - Basal cell carcinoma
   - Bladder cancer
   - Bone cancer (except for Ewing sarcoma)
   - Cancer of unknown primary site
   - Cervical cancer
   - Ductal carcinoma in situ (DCIS) of the breast
   - Hepatocellular carcinoma
   - Lung cancer (except for non-small cell lung cancer [NSCLC])
   - Multiple myeloma
   - Ovarian cancer
   - Penile cancer
   - Prostate cancer
   - Renal cancer
   - Soft tissue sarcomas (except for gastrointestinal stromal tumor [GIST])
   - Squamous cell carcinoma of the skin
   - Testicular cancer
   - Tracheal cancer
2. The following services are considered not medically necessary:
   A. Comparative analysis using short tandem repeat (STR) markers that is billed separately, as it is considered to be included as part of the genetic testing, except as described in Section 5 of Indications that are Covered. One example of comparative analysis using STR markers that is not covered is the know error® DNA Specimen Provenance Assay (DSPA) (Strand Diagnostics, LLC.).
   B. Direct-to-consumer genetic testing
   C. Genetic testing when the cause of the member's clinical condition, features, characteristics, or symptoms can be better explained by non-genetic factors, such as a known exposure to a toxic substance.
   D. Genetic testing that was not ordered by a licensed healthcare provider or physician (see Definitions) who has established a direct patient care relationship with the member to be tested.
   E. Genetic testing that is provided solely to satisfy data collection and analysis needs and that will not be used in direct clinical management.
   F. Predictive genetic testing for asymptomatic members under 18 years of age for conditions generally accepted as having an onset in adulthood.
   G. Repeat testing of a unique analyte (genetic or molecular substance) in a histologically-distinct tumor, whether at the same biopsy site or a different site, using the same or a similar genetic test, molecular profiling assay, or gene expression classifier.
   H. Multiple-gene panels that include genes not associated with ruling out potential causes for the clinical condition, features, characteristics, or symptoms being evaluated or that include genes not associated with the treatment or management of the condition being evaluated.
I. Epi proColor® (Epigenomics Inc.)
J. UroVysion™ Bladder Cancer Kit (Abbott Molecular, Inc.)
3. Genetic testing is considered not medically necessary when test results will not provide a diagnosis or unifying diagnosis or directly impact the treatment or management of the condition being evaluated.
**Definitions**

**Chimerism** is the presence of two genetically distinct cell lines in one person, such as cells derived from a transplant donor and a transplant recipient.

**Expanded genetic cancer marker tests** evaluate multiple genetic substances and other signs that may indicate the presence of cancer or give information about a cancer. Tests in this category could evaluate samples of blood, urine, stool, tumor tissue, or other sources. These tests could be offered before, during, or after cancer treatment.

**Healthcare provider** is any licensed non-physician (excluding naturopathic providers).

**Liquid biopsy** is a sample taken from blood, urine, or another bodily fluid to test for genetic substances and other signs that may indicate the presence of a tumor or give information about a tumor.

**Physician** is a licensed medical doctor or doctor of osteopathy.

**Proteomic pattern testing** examines the way proteins look and work in the body. The way these proteins look and work may indicate the presence of cancer or give information about a cancer.

**Topographic genotyping** (also known as molecular or integrated anatomic pathology) combines genetic testing with pathology services, such as sample preparation, genetic testing, and interpretation of the results by a pathologist.

**Codes**

If available, codes are listed below for informational purposes only, and do not guarantee member coverage or provider reimbursement. The list may not be all-inclusive.

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>0005U</td>
<td>Oncology (prostate) gene expression profile by real-time RT-PCR of 3 genes (ERG, PCA3, and SPDEF), urine, algorithm reported as risk score</td>
</tr>
<tr>
<td>0006</td>
<td>Oncology (hepatic), mRNA expression levels of 161 genes, utilizing fresh hepatocellular carcinoma tumor tissue, with alpha-fetoprotein level, algorithm reported as a risk classifier</td>
</tr>
<tr>
<td>0007</td>
<td>Oncology (gastrointestinal neuroendocrine tumors), real-time PCR expression analysis of 51 genes, utilizing whole peripheral blood, algorithm reported as a nomogram of tumor disease index</td>
</tr>
<tr>
<td>0009U</td>
<td>Oncology (breast cancer), ERBB2 (HER2) copy number by FISH, tumor cells from formalin-fixed paraffin-embedded tissue isolated using image-based dielectrophoresis (DEP) sorting, reported as ERBB2 gene amplified or non-amplified</td>
</tr>
<tr>
<td>0011</td>
<td>Oncology, prostate cancer, mRNA expression assay of 12 genes (10 content and 2 housekeeping), RT-PCR test utilizing blood plasma and/or urine, algorithms to predict high-grade prostate cancer risk</td>
</tr>
<tr>
<td>0012</td>
<td>Oncology (urothelial), mRNA, gene expression profiling by real-time quantitative PCR of five genes (MDK, HOXA13, CDC2 [CDK1], IGFBP5, and XCR2), utilizing urine, algorithm reported as a risk score for having urothelial carcinoma</td>
</tr>
<tr>
<td>0013</td>
<td>Oncology (urothelial), mRNA, gene expression profiling by real-time quantitative PCR of five genes (MDK, HOXA13, CDC2 [CDK1], IGFBP5, and CXCR2), utilizing urine, algorithm reported as a risk score for having recurrent urothelial carcinoma</td>
</tr>
<tr>
<td>0013U</td>
<td>Oncology (solid organ neoplasia), gene rearrangement detection by whole genome next-generation sequencing, DNA, fresh or frozen tissue or cells, report of specific gene rearrangement(s)</td>
</tr>
<tr>
<td>0014U</td>
<td>Hematology (hematolymphoid neoplasia), gene rearrangement detection by whole genome next-generation sequencing, DNA, whole blood or bone marrow, report of specific gene rearrangement(s)</td>
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</tbody>
</table>
| 0016U | Oncology (hematolymphoid neoplasia), RNA, BCR/ABL1 major and minor breakpoint fusion transcripts, quantitative PCR amplification, blood or bone marrow, report of fusion not detected or
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>0017U</td>
<td>Oncology (hematolymphoid neoplasia), JAK2 mutation, DNA, PCR amplification of exons 12-14 and sequence analysis, blood or bone marrow, report of JAK2 mutation not detected or detected</td>
</tr>
<tr>
<td>0018U</td>
<td>Oncology (thyroid), microRNA profiling by RT-PCR of 10 microRNA sequences, utilizing fine needle aspirate, algorithm reported as a positive or negative result for moderate to high risk of malignancy</td>
</tr>
<tr>
<td>0019U</td>
<td>Oncology, RNA, gene expression by whole transcriptome sequencing, formalin-fixed paraffin embedded tissue or fresh frozen tissue, predictive algorithm reported as potential targets for therapeutic agents</td>
</tr>
<tr>
<td>0022U</td>
<td>Targeted genomic sequence analysis panel, non-small cell lung neoplasia, DNA and RNA analysis, 23 genes, interrogation for sequence variants and rearrangements, reported as presence/absence of variants and associated therapy(ies) to consider</td>
</tr>
<tr>
<td>0023U</td>
<td>Oncology (acute myelogenous leukemia), DNA, genotyping of internal tandem duplication, p.D835, p.I836, using mononuclear cells, reported as detection or nondetection of FLT3 mutation and indication for or against the use of midostaurin</td>
</tr>
<tr>
<td>0026U</td>
<td>Oncology (thyroid), DNA and mRNA of 112 genes, next-generation sequencing, fine needle aspirate of thyroid nodule, algorithmic analysis reported as a categorical result (“Positive, high probability of malignancy” or “Negative, low probability of malignancy”)</td>
</tr>
<tr>
<td>0027U</td>
<td>JAK2 (Janus kinase 2) (eg, myeloproliferative disorder) gene analysis, targeted sequence analysis exons 12-15</td>
</tr>
<tr>
<td>0036U</td>
<td>Exome (ie, somatic mutations), paired formalin-fixed paraffin-embedded tumor tissue and normal specimen, sequence analyses</td>
</tr>
<tr>
<td>0037U</td>
<td>Targeted genomic sequence analysis, solid organ neoplasm, DNA analysis of 324 genes, interrogation for sequence variants, gene copy number amplifications, gene rearrangements, microsatellite instability and tumor mutational burden</td>
</tr>
<tr>
<td>0040U</td>
<td>BCR/ABL1 (t(9;22)) (eg, chronic myelogenous leukemia) translocation analysis, major breakpoint, quantitative</td>
</tr>
<tr>
<td>0045U</td>
<td>Oncology (breast ductal carcinoma in situ), mRNA, gene expression profiling by real-time RT-PCR of 12 genes (7 content and 5 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as recurrence score</td>
</tr>
<tr>
<td>0046U</td>
<td>FLT3 (fms-related tyrosine kinase 3) (eg, acute myeloid leukemia) internal tandem duplication (ITD) variants, quantitative</td>
</tr>
<tr>
<td>0047U</td>
<td>Oncology (prostate), mRNA, gene expression profiling by real-time RT-PCR of 17 genes (12 content and 5 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a risk score</td>
</tr>
<tr>
<td>0048U</td>
<td>Oncology (solid organ neoplasm), DNA, targeted sequencing of protein-coding exons of 468 cancer-associated genes, including interrogation for somatic mutations and microsatellite instability, matched with normal specimens, utilizing formalin-fixed paraffin-embedded tumor tissue, report of clinically significant mutation(s)</td>
</tr>
<tr>
<td>0049U</td>
<td>NPM1 (nucleophosmin) (eg, acute myeloid leukemia) gene analysis, quantitative</td>
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<tr>
<td>0050U</td>
<td>Targeted genomic sequence analysis panel, acute myelogenous leukemia, DNA analysis, 194 genes, interrogation for sequence variants, copy number variants or rearrangements</td>
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<tr>
<td>0053U</td>
<td>Oncology (prostate cancer), FISH analysis of 4 genes (ASAP1, HDAC9, CHD1 and PTEN), needle biopsy specimen, algorithm reported as probability of higher tumor grade</td>
</tr>
<tr>
<td>0056U</td>
<td>Hematology (acute myelogenous leukemia), DNA, whole genome next-generation sequencing to detect gene rearrangement(s), blood or bone marrow, report of specific gene rearrangement(s)</td>
</tr>
<tr>
<td>0057U</td>
<td>Oncology (solid organ neoplasia), mRNA, gene expression profiling by massively parallel sequencing for analysis of 51 genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a normalized percentile rank</td>
</tr>
<tr>
<td>0069U</td>
<td>Oncology (colorectal), microRNA, RT-PCR expression profiling of miR-31-3p, formalin-fixed paraffin-embedded tissue, algorithm reported as an expression score</td>
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<tr>
<td>0079U</td>
<td>Comparative DNA analysis using multiple selected single-nucleotide polymorphisms (SNPs), urine and buccal DNA, for specimen identity verification</td>
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<tr>
<td>0089U</td>
<td>Oncology (melanoma), gene expression profiling by RTqPCR, PRAME and LINC00518, superficial collection using adhesive patches</td>
</tr>
<tr>
<td>0090U</td>
<td>Oncology (cutaneous melanoma), mRNA gene expression profiling by RT-PCR of 23 genes (14 content and 9 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a categorical result (ie, benign, indeterminate, malignant)</td>
</tr>
<tr>
<td>0111U</td>
<td>Oncology (colon cancer), targeted KRAS (codons 12, 13, and 61) and NRAS (codons 12, 13, and 61) gene analysis utilizing formalin-fixed paraffin-embedded tissue</td>
</tr>
<tr>
<td>0113U</td>
<td>Oncology (prostate), measurement of PCA3 and TMPRSS2-ERG in urine and PSA in serum following prostatic massage, by RNA amplification and fluorescence-based detection, algorithm reported as risk score</td>
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<td>Code</td>
<td>Description</td>
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<tr>
<td>0118U</td>
<td>Transplantation medicine, quantification of donor-derived cell-free DNA using whole genome next-generation sequencing, plasma, reported as percentage of donor-derived cell-free DNA in the total cell-free DNA</td>
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<tr>
<td>0120U</td>
<td>Oncology (B-cell lymphoma classification), mRNA, gene expression profiling by fluorescent probe hybridization of 58 genes (45 content and 13 housekeeping genes), formalin-fixed paraffin-embedded tissue, algorithm reported as likelihood for primary mediastinal B-cell lymphoma (PMBCL) and diffuse large B-cell lymphoma (DLBCL) with cell of origin subtyping in the latter</td>
</tr>
<tr>
<td>0136U</td>
<td>ATM (ataxia telangiectasia mutated) (eg, ataxia telangiectasia) mRNA sequence analysis (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>0137U</td>
<td>PALB2 (partner and localizer of BRCA2) (eg, breast and pancreatic cancer) mRNA sequence analysis (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>0138U</td>
<td>BRCA1 (BRCA1, DNA repair associated), BRCA2 (BRCA2, DNA repair associated) (eg, hereditary breast and ovarian cancer) mRNA sequence analysis (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>0153U</td>
<td>Oncology (breast), mRNA, gene expression profiling by next-generation sequencing of 101 genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a triple negative breast cancer clinical subtype(s) with information on immune cell involvement</td>
</tr>
<tr>
<td>0154U</td>
<td>Oncology (urothelial cancer), RNA, analysis by real-time RT-PCR of the FGFR3 (fibroblast growth factor receptor 3) gene analysis (ie, p.R248C [c.742C&gt;T], p.S249C [c.746C&gt;G], p.G370C [c.1108G&gt;T], p.Y373C [c.1118A&gt;G], FGFR3-TACC3v1, and FGFR3-TACC3v3) utilizing formalin-fixed paraffin-embedded urothelial cancer tumor tissue, reported as FGFR gene alteration status</td>
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<tr>
<td>0171U</td>
<td>Targeted genomic sequence analysis panel, profile myeloid leukemia, myelodysplastic syndrome, and myeloproliferative neoplasms, DNA analysis, 23 genes, interrogation for sequence variants, rearrangements and minimal residual disease, reported as presence/absence</td>
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<tr>
<td>81120</td>
<td>IDH1 (isocitrate dehydrogenase 1 [NADP+], soluble) (eg, glioma), common variants (eg, R132H, R132C)</td>
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<tr>
<td>81121</td>
<td>IDH2 (isocitrate dehydrogenase 2 [NADP+], mitochondrial) (eg, glioma), common variants (eg, R140W, R172M)</td>
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<td>81170</td>
<td>ABL1 (ABL proto-oncogene 1, non-receptor tyrosine kinase) (eg, acquired imatinib tyrosine kinase inhibitor resistance), gene analysis, variants in the kinase domain</td>
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<td>81175</td>
<td>ASXL1 (additional sex combs like 1, transcriptional regulator) (eg, myelodysplastic syndrome, myeloproliferative neoplasms, chronic myelomonocytic leukemia), gene analysis; full gene sequence</td>
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<tr>
<td>81176</td>
<td>ASXL1 (additional sex combs like 1, transcriptional regulator) (eg, myelodysplastic syndrome, myeloproliferative neoplasms, chronic myelomonocytic leukemia), gene analysis; targeted sequence analysis (eg, exon 12)</td>
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<tr>
<td>81201</td>
<td>APC (adenomatous polyposis coli) (eg, familial adenomatosis polyposis [FAP], attenuated FAP) gene analysis; full gene sequence</td>
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<tr>
<td>81203</td>
<td>APC (adenomatous polyposis coli) (eg, familial adenomatosis polyposis [FAP], attenuated FAP) gene analysis; duplication/deletions variants</td>
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<tr>
<td>81206</td>
<td>BCR/ABL1 (t(9;22)) (eg, chronic myelogenous leukemia) translocation analysis; major breakpoint, qualitative or quantitative</td>
</tr>
<tr>
<td>81207</td>
<td>BCR/ABL1 (t(9;22)) (eg, chronic myelogenous leukemia) translocation analysis; minor breakpoint, qualitative or quantitative</td>
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<tr>
<td>81208</td>
<td>BCR/ABL1 (t(9;22)) (eg, chronic myelogenous leukemia) translocation analysis; other breakpoint, qualitative or quantitative</td>
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<tr>
<td>81210</td>
<td>BRAF (v-raf murine sarcoma viral oncogene homolog B1) (eg, colon cancer), gene analysis, V600E variant</td>
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<tr>
<td>81218</td>
<td>CEBPA (CCAAT/enhancer binding protein [C/EBP], alpha) (eg, acute myeloid leukemia), gene analysis, full gene sequence</td>
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<tr>
<td>81219</td>
<td>CALR (calreticulin) (eg, myeloproliferative disorders), gene analysis, common variants in exon 9</td>
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<tr>
<td>81228</td>
<td>Cyto genomic constitutional (genome-wide) microarray analysis; interrogation of genomic regions for copy number variants (eg, bacterial artificial chromosome [BAC] or oligo-based comparative genomic hybridization [CGH] microarray analysis)</td>
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<tr>
<td>81229</td>
<td>Cyto genomic constitutional (genome-wide) microarray analysis; interrogation of genomic regions for copy number and single nucleotide polymorphism (SNP) variants for chromosomal abnormalities</td>
</tr>
<tr>
<td>81233</td>
<td>BTK (Bruton's tyrosine kinase) (eg, chronic lymphocytic leukemia) gene analysis, common variants (eg, in urothelial cancer tissue, reported as FGFR gene alteration status</td>
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<td>0154U</td>
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<td>Code</td>
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<tr>
<td>81235</td>
<td>EGFR (epidermal growth factor receptor) gene analysis, common variants (eg, exon 19 LREA deletion, L858R, T790M, G719A, G719S, L861Q)</td>
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<tr>
<td>81236</td>
<td>EZH2 (enhancer of zeste 2 polycomb repressive complex 2 subunit) (eg, myelodysplastic syndrome, myeloproliferative neoplasms) gene analysis, full gene sequence</td>
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<tr>
<td>81237</td>
<td>EZH2 (enhancer of zeste 2 polycomb repressive complex 2 subunit) (eg, diffuse large B-cell lymphoma) gene analysis, common variant(s) (eg, codon 646)</td>
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<tr>
<td>81245</td>
<td>FLT3 (fms-related tyrosine kinase 3) (eg, acute myeloid leukemia), gene analysis; internal tandem duplication (ITD) variants (ie, exons 14, 15)</td>
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<tr>
<td>81246</td>
<td>FLT3 (fms-related tyrosine kinase 3) (eg, acute myeloid leukemia), gene analysis; tyrosine kinase domain (TKD) variants (eg, D835, I836)</td>
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<tr>
<td>81261</td>
<td>IGH@ (Immunoglobulin heavy chain locus) (eg, leukemias and lymphomas, B-cell), gene rearrangement analysis to detect abnormal clonal population(s); amplified methodology (eg, polymerase chain reaction)</td>
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<tr>
<td>81262</td>
<td>IGH@ (Immunoglobulin heavy chain locus) (eg, leukemias and lymphomas, B-cell), gene rearrangement analysis to detect abnormal clonal population(s); direct probe methodology (eg, Southern blot)</td>
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<tr>
<td>81263</td>
<td>IGK@ (Immunoglobulin kappa light chain locus) (eg, leukemia and lymphoma, B-cell), variable region somatic mutation analysis</td>
</tr>
<tr>
<td>81264</td>
<td>IGK@ (Immunoglobulin kappa light chain locus) (eg, leukemia and lymphoma, B-cell), gene rearrangement analysis, evaluation to detect abnormal clonal population(s)</td>
</tr>
<tr>
<td>81265</td>
<td>Comparative analysis using Short Tandem Repeat (STR) markers; patient and comparative specimen (eg, pre-transplant recipient and donor germline testing, post-transplant non-hematopoietic recipient germline [eg, buccal swab or other germline tissue sample] and donor testing, twin zygosity testing, or maternal cell contamination of fetal cells)</td>
</tr>
<tr>
<td>81266</td>
<td>Comparative analysis using Short Tandem Repeat (STR) markers; each additional specimen (eg, additional cord blood donor, additional fetal samples from different cultures, or additional zygosity in multiple birth pregnancies) (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>81267</td>
<td>Chimerism (engraftment) analysis, post transplantation specimen (eg, hematopoietic stem cell), includes comparison to previously performed baseline analyses; without cell selection</td>
</tr>
<tr>
<td>81268</td>
<td>Chimerism (engraftment) analysis, post transplantation specimen (eg, hematopoietic stem cell), includes comparison to previously performed baseline analyses; with cell selection (eg, CD3, CD33), each cell type</td>
</tr>
<tr>
<td>81270</td>
<td>JAK2 (Janus kinase 2) (eg, myeloproliferative disorder) gene analysis, p.Val617Phe (V617F) variant</td>
</tr>
<tr>
<td>81272</td>
<td>KIT (v-kit Hardy-Zuckerman 4 feline sarcoma viral oncogene homolog) (eg, gastrointestinal stromal tumor [GIST], acute myeloid leukemia, melanoma), gene analysis, targeted sequence analysis (eg, exons 8, 11, 13, 17, 18)</td>
</tr>
<tr>
<td>81273</td>
<td>KIT (v-kit Hardy-Zuckerman 4 feline sarcoma viral oncogene homolog) (eg, mastocytosis). gene analysis, D816 variant(s)</td>
</tr>
<tr>
<td>81275</td>
<td>KRAS (v-Ki-ras2 Kirsten rat sarcoma viral oncogene) (eg, carcinoma) gene analysis, variants in codons 12 and 13</td>
</tr>
<tr>
<td>81276</td>
<td>KRAS (Kirsten rat sarcoma viral oncogene homolog) (eg, carcinoma) gene analysis; additional variant(s) (eg, codon 61, codon 146)</td>
</tr>
<tr>
<td>81277</td>
<td>Cytogenomic neoplasia (genome-wide) microarray analysis, interrogation of genomic regions for copy number and loss-of-heterozygosity variants for chromosomal abnormalities</td>
</tr>
<tr>
<td>81287</td>
<td>MGMT (O-6-methylguanine-DNA methyltransferase) (eg, glioblastoma multiforme), methylation analysis</td>
</tr>
<tr>
<td>81288</td>
<td>MLH1 (mutL homolog 1, colon cancer, nonpolyposis type 2) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; promoter methylation analysis</td>
</tr>
<tr>
<td>81292</td>
<td>MLH1 (mutL homolog 1, colon cancer, nonpolyposis type 2) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; full sequence analysis</td>
</tr>
<tr>
<td>81294</td>
<td>MLH1 (mutL homolog 1, colon cancer, nonpolyposis type 2) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; duplication/deletion variants</td>
</tr>
<tr>
<td>81295</td>
<td>MSH2 (mutS homolog 2, colon cancer, nonpolyposis type 1) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; duplication/deletion variants</td>
</tr>
<tr>
<td>81296</td>
<td>MSH2 (mutS homolog 2, colon cancer, nonpolyposis type 1) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; full sequence analysis</td>
</tr>
<tr>
<td>81298</td>
<td>MSH6 (mutS homolog 6 [E. coli]) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; duplication/deletion variants</td>
</tr>
<tr>
<td>81300</td>
<td>MSH6 (mutS homolog 6 [E. coli]) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; duplication/deletion variants</td>
</tr>
<tr>
<td>81301</td>
<td>Microsatellite instability analysis (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) of markers for mismatch repair deficiency (eg, BAT25, BAT26), includes comparison of neoplastic and normal tissue, if performed</td>
</tr>
</tbody>
</table>
81305 MYD88 (myeloid differentiation primary response 88) (eg, Waldenstrom's macroglobulinemia, lymphoplasmacytic leukemia) gene analysis, p.Leu265Pro (L265P) variant

81307 PALB2 (partner and localizer of BRCA2) (eg, breast and pancreatic cancer) gene analysis; full gene sequence

81308 PALB2 (partner and localizer of BRCA2) (eg, breast and pancreatic cancer) gene analysis; known familial variant

81309 PIK3CA (phosphatidylinositol-4, 5-biphosphate 3-kinase, catalytic subunit alpha) (eg, colorectal and breast cancer) gene analysis, targeted sequence analysis (eg, exons 7, 9, 20)

81310 NPM1 (nucleoplasmin) (eg, acute myeloid leukemia) gene analysis, exon 12 variants


81313 PCA3/KLK3 (prostate cancer antigen 3 [non-protein coding]/kallikrein-related peptidase 3 [prostate specific antigen]) ratio (eg, prostate cancer)

81314 PGDFRA (platelet-derived growth factor receptor, alpha polypeptide) (eg, gastrointestinal stromal tumor (GIST)) gene analysis, targeted sequence analysis (eg, exons 12, 18)

81315 PML/RARalpha, (t(15;17)), (promyelocytic leukemia/retinoic acid receptor alpha) (eg, promyelocytic leukemia) translocation analysis; common breakpoints (eg, intron 3 and intron 6), qualitative or quantitative

81316 PML/RARalpha, (t(15;17)), (promyelocytic leukemia/retinoic acid receptor alpha) (eg, promyelocytic leukemia) translocation analysis; single breakpoint (eg, intron 3, intron 6 or exon 6), qualitative or quantitative

81317 PMS2 (postmeiotic segregation increased 2 [S. cerevisiae]) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; full sequence analysis

81319 PMS2 (postmeiotic segregation increased 2 [S. cerevisiae]) (eg, hereditary non-polyposis colorectal cancer, Lynch syndrome) gene analysis; duplication/deletion variants

81320 PLCG2 (phospholipase C gamma 2) (eg, chronic lymphocytic leukemia) gene analysis, common variants (eg, R665W, S707F, L845F)

81321 PTEN (phosphatase and tensin homolog) (eg, Cowden syndrome, PTEN hamartoma tumor syndrome) gene analysis; full sequence analysis

81322 PTEN (phosphatase and tensin homolog) (eg, Cowden syndrome, PTEN hamartoma tumor syndrome) gene analysis; duplication/deletion variant

81327 SEPT9 (Septin9) (eg, colorectal cancer) methylation analysis

81334 RUNX1 (run related transcription factor 1) (eg, acute myeloid leukemia, familial platelet disorder with associated myeloid malignancy), gene analysis, targeted sequence analysis (eg, exons 3-8)

81340 TRB@ (T cell antigen receptor, beta) (eg, leukemia and lymphoma), gene rearrangement analysis to detect abnormal clonal population(s); using amplification methodology (eg, polymerase chain reaction)

81341 TRB@ (T cell antigen receptor, beta) (eg, leukemia and lymphoma), gene rearrangement analysis to detect abnormal clonal population(s); using direct probe methodology (eg, Southern blot)

81342 TRG@ (T cell antigen receptor, gamma) (eg, leukemia and lymphoma), gene rearrangement analysis, evaluation to detect abnormal clonal population(s)

81345 TERT (telomerase reverse transcriptase) (eg, thyroid carcinoma, glioblastoma multiforme) gene analysis, targeted sequence analysis (eg, promoter region)

81400 Molecular pathology procedure, Level 1 (eg, identification of single germline variant [eg, SNP] by techniques such as restriction enzyme digestion or melt curve analysis) ACADM (acyl-CoA dehydrogenase, C-4 to C-12 straight chain, MCAD) (eg, medium chain acyl dehydrogenase deficiency), K304E variant ACE (angiotensin converting enzyme) (eg, hereditary blood pressure regulation), insertion/deletion variant AGTR1 (angiotensin II receptor, type 1) (eg, essential hypertension), 1166A>C variant BCKDHA (branched chain keto acid dehydrogenase E1, alpha polypeptide) (eg, maple syrup urine disease, type 1A), Y438N variant CCR5 (chemokine C-C motif receptor 5) (eg, HIV resistance), 32-bp deletion mutation/794 825del32 deletion CLRN1 (clarin1) (clarin1) (eg, Usher syndrome, type 3), N48K variant F2 (coagulation factor 2) (eg, hereditary hypercoagulability), 1199G>A variant F5 (coagulation factor V) (eg, hereditary hypercoagulability), HR2 variant F7 (coagulation factor VII [serum prothrombin conversion accelerator]) (eg, hereditary hypercoagulability), R353Q variant F13B (coagulation factor X, prothrombin) (eg, hereditary hypercoagulability), V34L variant FGB (fibrinogen beta chain) (eg, hereditary ischemic heart disease), -455G>A variant FGFR1 (fibroblast growth factor receptor 1) (eg, Pfeiffer syndrome type 1, craniosynostosis), P252R variant FGFR3 (fibroblast growth factor receptor 3) (eg, Muenke syndrome), P250R variant FKTN (fukutin) (eg, Fukuyama congenital muscular dystrophy), retrotransposon insertion variant GNE (glucosamine [UDP-N-acetyl]2-epimerase/N-acetylmannosamine kinase) (eg, inclusion body myopathy 2 [IBM2], Nonaka myopathy), M712T variant IVD (isovaleryl-CoA dehydrogenase) (eg, isovaleric acidemia), A282V variant LCT (lactase-hydrolyzate) (eg, lactose intolerance), 13910 C>T variant NEB (neulin) (eg, nemaline myopathy 2), exon 55 deletion variant PCDH15 (protocadherin-related 15) (eg, Usher syndrome type 1F), R245X variant SERPINE1 (serpine
peptidase inhibitor clade E, member 1, plasminogen activator inhibitor -1, PAI-1 (eg, thrombophilia), 4G variant SHOC2 (soc-2 suppressor of clear homolog) (eg, Noonan-like syndrome with loose anagen hair), S2G variant SRY (sex determining region Y) (eg, 46,XX testicular disorder of sex development, gonadal dysgenesis), gene analysis TOR1A (torsin family 1, member A [torsin A]) (eg, early-onset primary dystonia [DYT1]), 907_909delGAG (904_906delGAG) variant

81401 MOLECULAR PATHOLOGY PROCEDURE LEVEL 2
81402 MOLECULAR PATHOLOGY PROCEDURE LEVEL 3
81403 MOLECULAR PATHOLOGY PROCEDURE LEVEL 4
81404 MOLECULAR PATHOLOGY PROCEDURE LEVEL 5
81405 MOLECULAR PATHOLOGY PROCEDURE LEVEL 6
81406 MOLECULAR PATHOLOGY PROCEDURE LEVEL 7
81407 MOLECULAR PATHOLOGY PROCEDURE LEVEL 8
81408 MOLECULAR PATHOLOGY PROCEDURE LEVEL 9

81445 Targeted genomic sequence analysis panel, solid organ neoplasm, DNA analysis, and RNA analysis when performed, 5-50 genes (eg, ALK, BRAF, CDKN2A, EGFR, ERBB2, KIT, KRAS, NRAS, MET, PDGFRA, PDGFRB, PGR, PIK3CA, PTEN, RET), interrogation for sequence variants and copy number variants or rearrangements, if performed

81450 Targeted genomic sequence analysis panel, hematolymphoid neoplasm or disorder, DNA analysis, and RNA analysis when performed, 5-50 genes (eg, BRAF, CEBPA, DNMT3A, EZH2, FLT3, IDH1, IDH2, JAK2, KRAS, KIT, MLL, NRAS, NPM1, NOTCH1), interrogation for sequence variants, and copy number variants or rearrangements, or isoform expression or mRNA expression levels, if performed

81455 Targeted genomic sequence analysis panel, solid organ or hematolymphoid neoplasm, DNA and RNA analysis when performed, 51 or greater genes (eg, ALK, BRAF, CDKN2A, CEPA, DNMT3A, EGFR, ERBB2, EZH2, FLT3, IDH1, IDH2, JAK2, KRAS, KIT, MLL, NRAS, NPM1, NOTCH1, PDGFRA, PDGFRB, PGR, PIK3CA, PTEN, RET), interrogation for sequence variants and copy number variants or rearrangements, if performed

81504 Oncology (tissue of origin), microarray gene expression profiling of > 2000 genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as tissue similarity scores

81518 Oncology (breast), mRNA, gene expression profiling by real-time RT-PCR of 11 genes (7 content and 4 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithms reported as percentage risk for metastatic recurrence and likelihood of benefit from extended endocrine therapy

81519 Oncology (breast), mRNA, gene expression profiling by real-time RT-PCR of 21 genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as recurrence score

81520 Oncology (breast), mRNA, gene expression profiling by hybrid capture of 58 genes (50 content and 8 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a recurrence risk score

81521 Oncology (breast), mRNA, microarray gene expression profiling of 70 content genes and 465 housekeeping genes, utilizing fresh frozen or formalin-fixed paraffin-embedded tissue, algorithm reported as index related to risk of distant metastasis

81522 Oncology (breast), mRNA, gene expression profiling by RT-PCR of 12 genes (8 content and 4 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as recurrence risk score

81525 Oncology (colon), mRNA, gene expression profiling by real-time RT-PCR of 12 genes (7 content and 5 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a recurrence score

81540 Oncology (tumor of unknown origin), mRNA, gene expression profiling by real-time RT-PCR of 92 genes (87 content and 5 housekeeping) to classify tumor into main cancer type and subtype, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a probability of a predicted main cancer type and subtype

81541 Oncology (prostate), mRNA gene expression profiling by real-time RT-PCR of 46 genes (31 content and 15 housekeeping), utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a disease-specific mortality risk score

81542 Oncology (prostate), mRNA, microarray gene expression profiling of 22 content genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as metastasis risk score

81545 Oncology (thyroid), gene expression analysis of 142 genes, utilizing fine needle aspirate, algorithm reported as a categorical result (eg, benign or suspicious)

81551 Oncology (prostate), promoter methylation profiling by real-time RT-PCR of 3 genes (GSTP1, APC, RASSF1), utilizing formalin-fixed, paraffin-embedded tissue, algorithm reported as a likelihood of prostate cancer detection on repeat biopsy

81552 Oncology (uveal melanoma), mRNA, gene expression profiling by real-time RT-PCR of 15 genes (12 content and 3 housekeeping), utilizing fine needle aspirate or formalin-fixed paraffin-embedded tissue,
algorithm reported as risk of metastasis

81599  Unlisted multianalyte assay with algorithmic analysis

88120  Cytopathology, in situ hybridization (eg, FISH), urinary tract specimen with morphometric analysis, 3-5 molecular probes, each specimen; manual

88121  Cytopathology, in situ hybridization (eg, FISH), urinary tract specimen with morphometric analysis, 3-5 molecular probes, each specimen; using computer-assisted technology

88182  Flow cytometry, cell cycle or DNA analysis

88245  Chromosome analysis for breakage syndrome; baseline Sister Chromatic Exchange (SCE), 20-25 cells

88248  Chromosome analysis for breakage syndromes; baseline breakage, score 50-100 cells, count 20 cells, 2 karyotypes (eg, for ataxia telangiectasia, Fanconi anemia, fragile X)

88249  Chromosome analysis for breakage syndromes; score 100 cells, clastogen stress (eg, diepoxybutane, mitomycin C, ionizing radiation, UV radiation

88261  Chromosome analysis; count 5 cells, 1 karyotype, with banding

88262  Chromosome analysis; count 15-20 cells, 2 karyotypes, with banding

88263  Chromosome analysis; count 45 cells for mosaicism, 2 karyotypes, with banding

88264  Chromosome analysis, analyze 20-25 cells

88271  Molecular cytogenetics; DNA probe, each (eg, FISH)

88272  Molecular cytogenetics; chromosomal in situ hybridization, analyze 3-5 cells (eg, for derivatives and markers)

88273  Molecular cytogenetics; chromosomal in situ hybridization, analyze 10-30 cells (eg, for microdeletions)

88274  Molecular cytogenetics; interphase in situ hybridization, analyze 25-99 cells

88275  Molecular cytogenetics; interphase in situ hybridization, analyze 100-300 cells

88280  Chromosome analysis; additional karyotypes, each study

88283  Chromosome analysis; additional specialized banding technique (eg, NOR, C-banding)

88285  Chromosome analysis; additional cells counted, each study

88289  Chromosome analysis; additional high resolution study

88299  Unlisted cytogenetic study

88358  Morphometric analysis; tumor (eg, DNA ploidy)

88364  In situ hybridization (eg, FISH), per specimen; each additional single probe stain procedure (List separately in addition to code for primary procedure)

88365  In situ hybridization (eg, FISH), per specimen; initial single probe stain procedure

88366  In situ hybridization (eg, FISH), per specimen; each multiplex probe stain procedure

88367  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), using computer-assisted technology, per specimen; initial single probe stain procedure

88368  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), manual, per specimen; initial single probe stain procedure

88369  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), manual, per specimen; each additional single probe stain procedure (List separately in addition to code for primary procedure)

88373  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), using computer-assisted technology, per specimen; each additional single probe stain procedure (List separately in addition to code for primary procedure)

88374  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), using computer-assisted technology, per specimen; each multiplex probe stain procedure

88377  Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), manual, per specimen; each multiplex probe stain procedure

G9840  RAS (KRAS and NRAS) gene mutation testing performed before initiation of anti-EGFR MoAb

S3854  Gene expression profiling panel for use in the management of breast cancer treatment

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References


