| Disease | Drug | Company | Therapeutic Role | Type of Drug | Diagnostic Test Specified | Approval Type | Medicare Payment Category |
|-----------------------------------|--|----------------------------------|--|---|--|------------------|---------------------------------|
| Breast Cancer (BC) | Margenza (margetuximab- | Macrogenics | Third-line for metastatic, | HER2-neu receptor | None | Full | Part B |
| | cmkb) Phesgo (pertuzumab, trastuzumab, hyaluronidase- zzxf) | Genentech | HER2+ disease Neoadjuvant for HER2+ early BC, adjuvant for high risk HER2+ early BC, and first-line for HER2+ metastatic BC. | antagonist Conjugate | HER2 protein overexpression or HER2 gene amplification | Full | Part B |
| | Trodelvy® (sacituzumab govitecan-hziy) | Immunomedics | Third-line for triple negative BC | Trop-2-directed antibody and topoisomerase inhibitor conjugate | None | Accelerated | Part B |
| | Tukysa® (tucatinib) | Seattle Genetics | Second-line for unresectable or metastatic HER2+ BC | Kinase inhibitor | None | Full | Part D |
| Cholangiosarcoma | Pemazyre® (pemigatinib) | Incyte | Second-line for unresectable/metastatic disease with a FGFR2 fusion or rearrangement | Kinase inhibitor | Presence of an FGFR2 fusion or rearrangement | Accelerated | Part D |
| Diffuse Large B-Cell Lymphoma | Monjuvi®(tafasitamab-cxix) | MorphoSys/Incyte | Relapsed/refractory patients who are not eligible for ASCT | CD19-directed cytolytic antibody | None | Accelerated | Part B |
| Epitheliod sarcoma | Tazverik® (tazemetostat) | Epizyme, Inc. | Patients with metastatic/locally advanced disease ineligible for complete resection | Methyltransferase inhibitor | None | Accelerated | Part D |
| Follicular Lymphoma (FL) | | | Third-line for relapsed/refractory EZH2+ positive disease and for FL patients w/o satisfactory treatment options. | | Tumors are positive for an EZH2 mutation | Accelerated | |
| Gastrointestinal Stromal Tumor | Ayvakit (avapritinib) | Blueprint Medicines Corp. | First-line for unresectable/metastatic PDGFRA mutation-positive disease | Kinase inhibitor | Positive for PDGFRA exon 18 mutation, including PDGFRA D842V mutations | Full | Part D |
| | Qinlock® (ripretinib) | Deciphera Pharmaceuticals | 4th-line therapy | Kinase inhibitor | None | Full | Part D |
| Lung Cancer, Non-Small Cell | Gavreto® (pralsetinib) | Genentech/Blueprint Medicines | Metastatic RET fusion-positive disease | Kinase inhibitor | Presence of a <i>RET</i> gene fusion | Accelerated | Part D |
| | Retevmo® (selpercatinib) | Eli Lilly | Metastatic RET fusion-positive disease | Kinase inhibitor | Presence of a RET gene fusion (FDA-approved test for the detection of RET gene fusion not currently available) | Accelerated | Part D |
| | Tabrecta (capmatinib) | Novartis | Metastatic disease positive for MET exon 14 skipping | Kinase inhibitor | Presence of a mutation that leads to MET exon 14 skipping. | Accelerated | Part D |
| Lung Cancer, Small-Cell | Zepzelca (lurbinectedin) | Jazz Pharmaceuticals | Metastatic small cell disease progressing on/after platinumbased chemo | Alkylating agent | None | Accelerated | Part B |
| Lymphoma, Mantle Cell | Tecartus (brexucabtagene autoleucel) | KITE/Gilead | Relapsed/refractory disease | Genetically modified autologous t-cells | None | Accelerated | Part B |
| Multiple Myeloma | Blenrep (belantamab mafodotin-blmf) | GlaxoSmithKline | therapy for relapsed/refractory | Antibody-drug Conjugate | None | Accelerated | Part B |
| | Sarclisa® (isatuximab-irfc) | Sanofi | 3rd-line therapy after lenalidomide and proteasome inhibitor | CD38-directed cytolitic antibody | None | Full | Part B |
| Myelodysplastic Syndrome | Inqovi® (decitabine, cedaruzidine) | Taiho Oncology | Myelodysplastic Syndrome (subtypes) | Conjugate of nucleoside metabolic and cytadine deaminase inhibitors | None | Full | Part D |
| Neuroblastoma | Danyelza® (namitamab-gqgk) | Y-mAbs Therapeutics | Relapsed/refractory high risk disease in bone marrow which is responsive to prior therapy | GD2-binding antibody | None | Accelerated | Part B |
| Prostate Cancer | Orgovyx (relugolix) | Myovant Sciences | Advanced disease | GnRH receptor antagonist | None | Full | Part D |
| Medullary Thyroid Cancer | Gavreto® (pralsetinib) | Genentech/Blueprint Medicines | Advanced or metastatic RET- mutant medullary disease | Kinase inhibitor | Presence of RET gene mutation (FDA-approved | Accelerated | Part D |
| | Retevmo® (selpercatinib) | Eli Lilly | | | test not currently available). | | |
| Thyroid Cancer | Gavreto® (pralsetinib) | Genentech/Blueprint Medicines | Advanced or metastatic RET fusion-positive patients who | Kinase inhibitor | Presence of a RET gene fusion (FDA-approved test not currently | Accelerated | Part D |
| | Retevmo® (selpercatinib) | Eli Lilly | are radio-iodone sensitive | | available) | | |