

Policy Number: PA.018.MPC Last Review Date: 08/18/2022 Effective Date: 09/01/2022

PA.018.MPC Gene Expression Testing for Breast Cancer

Maryland Physicians Care considers **gene expression testing** for breast cancer medically necessary for the following indications:

Prosigna (Prosigna® Breast Cancer Prognostic Gene Signature Assay) is considered medically necessary <u>only for post-menopausal women who meet all of the</u> <u>following criteria</u>:

- 1. Diagnosed with Stage I or II breast cancer within the previous six months; and
- 2. Lymph node-negative or Stage II with 1-3 positive nodes; and
- 3. Estrogen receptor (ER) positive tumor; and
- 4. Her2 negative tumor; and
- 5. The woman would be a candidate for adjuvant chemotherapy and
- 6. The result of the test will guide the decision whether or not to use chemotherapy; and
- 7. The woman would choose to receive chemotherapy if offered.

MammaPrint® is considered medically necessary <u>only for women who meet all of</u> <u>the following criteria</u>:

- 1. Diagnosed with stage I or II breast cancer within the previous six months; and
- 2. Lymph node-negative; and
- 3. Tumor smaller than 5.0 centimeters; and
- 4. Estrogen receptor (ER) positive or negative tumor; and
- 5. Her2 negative tumor; and
- 6. The woman would be a candidate for adjuvant chemotherapy and
- 7. The result of the test will guide the decision whether or not to use chemotherapy; and
- 8. The woman would choose to receive chemotherapy if offered.

Oncotype DX (Oncotype DX® Breast Recurrence Score) is considered medically necessary <u>only for women who meet all the following criteria</u>:

- 1. Diagnosed with Stage I or II breast cancer within the previous six months; and
- 2. Lymph node-negative; and
- 3. Estrogen receptor (ER) positive tumor; and
- 4. Her2 negative tumor; and
- 5. The woman would be a candidate for adjuvant chemotherapy and



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- 6. The result of the test will guide the decision whether or not to use chemotherapy; and
- 7. The woman would choose to receive chemotherapy if offered.

Breast Cancer Index (BCI) (bioTheranostics) is considered medically necessary only for women who meet all of the following criteria and have Medicare coverage:

- 1. Member with non-relapsed, ER+ breast cancer,
- 2. Was lymph node negative,
- 3. Is completing five years of endocrine therapy,
- 4. Patient must be eligible for consideration of extended endocrine therapy based on published clinical trial data or practice guidelines,
- 5. Physician or patient is concerned about continuing anti-hormonal therapy because of documented meaningful toxicity or possible significant patient specific side effects,
- 6. The test results will be discussed with the patient (including the limitations of the testing method, the risks and benefits of either continuing or stopping the therapy based on the test, and current cancer management guidelines).

Endopredict (EndoPredict® for Breast Cancer Prognosis) is considered medically necessary only for women with T1-3, N0-1 breast cancer when the following criteria are met:

- 1. Tumor size greater than 0.5cm (5mm) in greatest dimension (T1b-T3); and
- 2. Patient is post-menopausal, and
- 3. Pathology (excisional or biopsy) reveals invasive carcinoma of the breast that is ER-positive, Her2-negative, and
- 4. Patient is either lymph node-negative or has 1-3 positive lymph nodes, and
- 5. Patient has no evidence of distant metastasis, and
- 6. Test result will be used to determine treatment choice between endocrine therapy alone vs. endocrine therapy plus chemotherapy.

Note: The EndoPredict test should not be ordered if a physician does not intend to act upon the test result.



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Limitations

- 1. <u>A maximum of one genomic assay per breast tumor is considered</u> <u>medically necessary and therefore second or subsequent genomic assays</u> <u>on the same tumor are not covered</u>.
- 2. Gene expression testing for breast cancer will <u>only be covered if the test is</u> <u>ordered by an oncologist.</u>
- 3. Since gene expression assays analyze ribonucleic acid (RNA) and RNA is unstable:
 - Assay samples are to be obtained at the time of tumor excision;
 - Time from excision to fixation should be less than 1 hour (see College of American Pathologists/American Society of Clinical Oncology (CAP/ASCO) protocol; reference below);
 - Formalin fixed paraffin blocks that are greater than six months old should not be tested.
- Maryland Physicians Care considers Mammaprint, Oncotype DX, and Prosigna to be <u>experimental/investigational for any other uses, reasons, or tissue</u> <u>type</u>.

Background

About one in eight women will develop breast cancer during their lifetime in the United States. Breast cancer is the second most commonly diagnosed cancer among women behind skin cancer. Breast cancer refers to a malignant tumor in the breast caused by an uncontrolled growth in cells and it is always caused by a genetic abnormality. Leading risk factors include increasing in age and being of the female sex.

If breast cancer is detected, various tests can be performed for additional insight on the rate of growth, likelihood of spreading or return and the potential effectiveness of treatment options. Mutigene tests are one set of tests utilized and they can include:

- Oncotype DX: multigene test that predicts whether Stage I or Stage II breast cancer that is estrogen receptor positive and node negative will spread to other parts of the body.
- MammaPrint: multigene test that predicts whether Stage I or Stage II breast cancer that is node negative will spread to other parts of the body.
- Prosigna: multigene test for post-menopausal women that predicts whether Stage I or Stage II breast cancer that is estrogen receptor positive and node negative will spread to other parts of the body



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If these tests indicate a high chance of spreading, chemotherapy may be given to lower the risk.

Oncotype DX Scores (risk of recurrence)

- 0 < 18: low risk
- 18-31: intermediate risk
- \geq 31: high risk

Prosigna Scores (risk of recurrence):

- 0-40: low risk
- 41-60:intermediate risk
- 61-100: high risk

Codes:

CPT/HCPCS Codes	
Code	Description
*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
	*Lab services does not require PA when performed by contracted lab in a freestanding facility and/or services allowed in a participating physician's office if the office is not in regulated space.
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
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
81523	Oncology (breast), mRNA, next-generation sequencing gene expression profiling of 70 content genes and 31 housekeeping genes, utilizing formalin-



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	fixed paraffin-embedded tissue, algorithm reported as index related to risk to distant metastasis
S3854	Gene expression profiling panel for use in the management of breast cancer treatment

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