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# Subject: Magnetic Resonance Imaging (MRI) of the Breast

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Position Statement	Billing/Coding	Reimbursement	Program Exceptions	<u>Definitions</u>	Related Guidelines
<u>Other</u>	<u>References</u>	<u>Updates</u>			

## **DESCRIPTION:**

Magnetic resonance imaging (MRI) of the breast is an imaging modality for the detection and characterization of breast disease, assessment of local extent of disease, evaluation of treatment response, and guidance for biopsy and localization. Breast MRI should be bilateral except for individuals with a history of mastectomy or when the MRI is being performed expressly to further evaluate or follow findings in one breast. MRI findings should be correlated with clinical history, physical examination results, and the results of mammography and prior breast imaging. MRI of the breast is performed using MR scanners and intravenous magnetic resonance contrast agents. MRI examinations should be performed with a dedicated breast MRI coil unless obesity or other individual consideration requires modification of the imaging procedure.

**Summary and Analysis of Evidence:** An UpToDate review on "MRI of the breast and emerging technologies" (Slanetz) states that "Magnetic resonance imaging (MRI) of the breast is established for certain indications. The high sensitivity of MRI for breast cancer has led to the increasing use of MRI for breast cancer detection, assessment, and treatment monitoring. Breast MRI has been used for the assessment of silicone implant integrity, diagnosis of occult primary breast cancers, determination of disease extent in newly diagnosed breast cancer patients (when indicated), documentation of response to neoadjuvant chemotherapy, diagnosis of recurrence, clarification of inconclusive clinical or mammographic findings, and screening of high-risk patients." Magnetic resonance imaging (MRI) of the breast is a useful tool for the detection and characterization of breast cancer, assessment of local disease extent, evaluation of neoadjuvant treatment response, and guidance for biopsy and localization. MRI findings should be correlated with the clinical history, physical examination findings, and results of any recent breast imaging (ACR, 2023).

## **POSITION STATEMENT:**

Magnetic resonance imaging (MRI) of the breast **meets the definition of medical necessity** for the following indications:

#### Screening examination to detect breast cancer in any of the following:

- Breast cancer screening in members with high risk of breast cancer as defined by breast cancer risk assessment/clinical risk estimators (models), with calculated lifetime risk of 20% or greater of developing breast cancer
- Members with lifetime risk of 20% or greater of developing breast cancer based on history of lobular neoplasia (LCIS [lobular carcinoma in situ]/ALH [atypical lobular hyperplasia]) or ADH [atypical ductal hyperplasia]
- Members with history of extensive chest irradiation between ages (10 and 30)
- Members with known BRCA mutation
- Members not yet tested for BRCA gene, but with known BRCA mutation in first degree relative (parents, siblings, children)
- Personal history of germline mutations known to predispose to a high risk of breast cancer.

#### Evaluation of identified lesion, mass or abnormality in breast for any of the following

- Evaluation of suspected breast cancer when other imaging examinations, such as ultrasound and mammography, and physical examination are inconclusive for the presence of breast cancer, and biopsy could not be performed
- Inconclusive or conflicting findings on a diagnostic mammogram or ultrasound when the finding is not a palpable or a discrete mass
- Evaluation of suspicious mass, lesion, distortion, or abnormality of the breast in member with history of breast cancer when other imaging is inconclusive
- Nipple inversion (new) when mammographic and sonographic findings are inconclusive and a biopsy cannot be performed
- Member diagnosed with biopsy-proven lobular neoplasia (i.e., LCIS/ALH (lobular carcinoma in situ/atypical lobular hyperplasia) or ADH (atypical ductal hyperplasia)
- Spontaneous unilateral serous or bloody nipple discharge when conventional imaging is normal and there is no palpable mass
- Paget's disease of the nipple: to detect underlying ductal carcinoma when conventional imaging is normal and there is no palpable mass
- For a phylloides tumor diagnosed by biopsy, breast MRI may help determine extent of disease and resectability in selected cases
- Follow-up of a probably benign (BI-RADS 3) lesion seen only on prior MRI.

#### History of known breast cancer

MRI breast surveillance (annual) for any of the following:

- Members with history of breast cancer and dense breast tissue on mammography
- Members with personal history of breast cancer diagnosed before age 50

• Members with genetic or other risk factors placing them at high risk for a new cancer or recurrence.

#### Staging, treatment, and surveillance of members with a known history of breast cancer

- Initial staging when conventional imaging is indeterminate in defining the extent of cancer or presence of multifocal, multicentric, contralateral cancer or there is a discrepancy in estimated tumor size between physical exam and imaging
- Invasive lobular carcinoma that is poorly or inadequately defined by mammography, ultrasound, or physical exam
- To identify primary cancer in a member with axillary nodal adenocarcinoma and unidentified primary tumor
- Prior to treatment: To serve as a baseline for comparison prior to a member starting planned neoadjuvant chemotherapy
- During or after treatment: To identify candidates for breast conserving therapy or evaluate response to treatment, including preoperative neoadjuvant therapy (within three (3) months).

#### Silicone implants

- Confirmation of silicone gel-filled breast implant ruptures in asymptomatic members, after an abnormal or indeterminate finding on mammography or breast ultrasound
- Postoperative evaluation of silicone breast implant complications when other imaging is inconclusive.

#### **Pre-operative evaluation**

• Pre-operative evaluation for known breast cancer when surgery planned within thirty (30) days.

#### Post-operative/procedural evaluation

• A follow-up study may be needed to help evaluate a member's progress after treatment, procedure, intervention or surgery. Documentation required.

## **BILLING/CODING INFORMATION:**

#### **CPT Coding**

77046	Magnetic resonance imaging, breast, without contrast material; unilateral		
77047	Magnetic resonance imaging, breast, without contrast material; bilateral		
77048	Magnetic resonance imaging, breast, without and with contrast material(s),		
	including computer-aided detection (CAD real-time lesion detection,		
	characterization and pharmacokinetic analysis), when performed, unilateral		
77049	Magnetic resonance imaging, breast, without and with contrast material(s),		
	including computer-aided detection (CAD real-time lesion detection,		
	characterization and pharmacokinetic analysis), when performed, bilateral		

## LOINC Codes:

The following information may be required documentation to support medical necessity: physician history and physical, physician progress notes, plan of treatment, radiographic study reports (e.g., mammography, breast ultrasound, MRI breast) and reason for magnetic resonance imaging of the breast.

Documentation Table	LOINC	LOINC	LOINC Time Frame Modifier Codes
	Codes	Time Frame	Narrative
		Modifier Code	
Physician history and	28626-0	18805-2	Include all data of the selected type that
physical			represents observations made six months
			or fewer before starting date of service
			for the claim
Attending physician visit	18733-6	18805-2	Include all data of the selected type that
note or treatment notes			represents observations made six months
			or fewer before starting date of service
			for the claim.
Radiology Study Reports	18726-0	18805-2	Include all data of the selected type that
(e.g., mammography,			represents observations made six months
breast ultrasound, MRI			or fewer before starting date of service
breast)			for the claim.

# **REIMBURSEMENT INFORMATION:**

Refer to section entitled **POSITION STATEMENT**.

### Follow-up study

A follow-up study may be needed to help evaluate a member's progress after treatment, procedure, intervention or surgery. Documentation requires a medical reason that clearly indicates why additional imaging is needed for the type and area(s) requested.

# **PROGRAM EXCEPTIONS:**

Federal Employee Plan (FEP): Follow FEP guidelines.

**Medicare Advantage products:** No Local Coverage Determination (LCD) was found at the time of the last guideline reviewed date. The following National Coverage Determination (NCD) was reviewed on the last guideline reviewed date: Magnetic Resonance Imaging, (220.2) located at cms.gov.

If this Medical Coverage Guideline contains a step therapy requirement, in compliance with Florida law 627.42393, members or providers may request a step therapy protocol exemption to this requirement if based on medical necessity. The process for requesting a protocol exemption can be found at <u>Coverage</u> <u>Protocol Exemption Request</u>.

## **DEFINITIONS:**

No guideline specific definitions apply.

## **RELATED GUIDELINES:**

Genetic Testing for Hereditary Breast or Ovarian Cancer, 05-82000-30

## **OTHER:**

None.

### **REFERENCES:**

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- Henderson JT, Webber EM, Weyrich MS, et al. Screening for Breast Cancer: Evidence Report and Systematic Review for the US Preventive Services Task Force. JAMA. 2024 Jun 11;331(22):1931-1946. [Abstract]
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- 16. Juanpere S, Perez E, Huc O, et al. Imaging of breast implants-a pictorial review. Insights Imaging. 2011 Dec;2(6):653-670.
- 17. Khatcheressian JL, Hurley P, Bantug E et al. Breast cancer follow-up and management after primary treatment: American Society of Clinical Oncology clinical practice guideline update. Journal of Clinical Oncology 2013; 31(7): 961-965.
- 18. Killelea B, Sowden M. Nipple inversion. In: UpToDate, Post TW (Ed), UpToDate, Waltham, MA. (Accessed on January 24, 2022.)
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- 20. National Cancer Institute Breast Cancer Risk Assessment Tool. Assessed 01/24/22.
- 21. National Cancer Institute Breast Cancer Screening (PDQ<sup>®</sup>), 09/13/21.
- 22. National Comprehensive Cancer Network (NCCN). NCCN Clinical Practice Guidelines in Oncology. Breast Cancer Screening and Diagnosis. Version 3.2023.
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# **COMMITTEE APPROVAL:**

This Medical Coverage Guideline (MCG) was approved by the Florida Blue Medical Policy and Coverage Committee on 05/22/25.

# **GUIDELINE UPDATE INFORMATION:**

10/15/04	New Medical Coverage Guideline.
10/15/05	Annual review. Added detection of local tumor recurrence in individuals with breast
	cancer who have radiographically dense breasts or old scar tissue from previous breast
	surgery that compromises the ability of combined mammography and ultrasonography
	to when services are covered. Updated references.
08/15/06	Scheduled review. Added 0159T (July CPT update). Updated references.
01/01/07	HCPCS update. Deleted 76093 and 76094. Added 77058 and 77059.
07/01/07	Scheduled review; revised position statements; reformatted guideline; updated
	references.
11/15/07	Added Li-Fraumeni syndrome or Cowden syndrome or Bannayan-Riley-Ruvalcaba
	syndrome or who have a first-degree relative with one of these syndromes to position
	statement. Updated definitions and references.
01/21/08	Updated Program Exceptions.
07/15/08	Revised position statement to indicate that MRI of breast may be considered medically
	necessary for the following indications for screening for breast cancer in the following
	patients: known BRCA1 or BRCA2 mutation in patient or relative; high risk (lifetime risk
	about 20 to 25 percent or greater) of developing breast cancer as identified by models
	that are largely defined by family history (breast, ovarian) annual screening 10 years
	after therapeutic chest irradiation; or Li-Fraumeni syndrome or Cowden syndrome or
	Bannayan-Riley-Ruvalcaba syndrome or who have a first-degree relative with one of
	these syndromes. Added the following indications to the position statement (meets the
	definition of medical necessity): for evaluation of the contralateral breast in patients
	with a new diagnosis of breast cancer who have normal clinical and mammographic
	findings in the contralateral breast for preoperative tumor mapping of the involved
	(ipsilateral) breast to evaluate the presence of multicentric disease in patients with
	clinically localized breast cancer who are candidates for breast-conservation therapy
	Deleted "in order to avoid biopsy" for the diagnosis of a suspicious breast lesion. Added
	"i.e., indeterminate breast lesion" for the diagnosis of a suspicious breast lesion. Added
	ipsilateral to the definitions section. Updated references.
09/15/08	Added criteria for detection of tumor recurrence.
05/21/09	Removed Federal Employee Plan (FEP) from BCBSF Radiology Management program
	exception statement. Added FEP program exception statement: FEP is excluded from the
	National Imaging Associates (NIA) review; follow FEP guidelines.
07/01/09	Updated BCBSF Radiology Management program exception; added BlueSelect.
10/15/09	Revised position statement (experimental or investigational), to clarify diagnosis of
	suspicious breast lesion. Added position statement for evaluation of residual tumor with
	positive margins after lumpectomy. Updated description section. Updated references.
01/01/10	Revised BCBSF Radiology Management program exception section.

10/15/10	Annual review: Revised position statement regarding mass, distortion, etc; added "or
	abnormality in breast". Added position statement for evaluation of a documented
	abnormality of the breast prior to obtaining an MRI-guided biopsy when there is
	documentation that other methods (e.g., palpation, ultrasound) are not able to localize
	the lesion for biopsy. Revised position statement regarding the evaluation of residual
	tumor in individuals found to have positive margins after lumpectomy, and deleted
	"when lumpectomy was performed as the definitive surgery". Updated references.
10/01/11	Revision; formatting changes.
10/15/11	Annual review; maintain position statements. Updated references.
06/15/12	Note added for MRI as first-line screening modality in women with increased risk for
	breast cancer.
01/15/13	Annual review; no change in position statements. Repositioned the statement for MRI a
	first-line screening modality in women with increased risk for breast (moved to the
	position statement section). Updated references.
01/01/14	Review. Updated program exception.
03/15/15	Annual review; revised position statement; updated description and references.
11/15/16	Revision; revised position statement. Added additional information related to breast MRI. Updated
00/15/10	references.
06/15/18	Revision; revised position statement. Updated references.
01/01/19	Annual HCPCs code update. Deleted 7/058, 7/059 and 01591. Added 7/046, 7/047,
02/45/20	77048 and 77049.
02/15/20	Review/revision. Revised criteria for silicone implants. Revised and added criteria for
	screening to detect breast cancer. Revised and add criteria for identified lesion, mass or
	abnormality in breast. Added indication and criteria for staging, treatment, and
	surveillance of members with a known history of breast cancer. Revised criteria for
05/45/00	evaluation of identified lesion, mass or abnormality in breast. Updated references.
05/15/22	Review/revision. Revised and expanded criteria for: screening examination to detect
	breast cancer, identified lesion, mass or abnormality in breast, staging, treatment, and
	surveillance for known history of breast cancer. Expanded criteria for history of known
07/04/00	breast cancer. Updated references.
07/01/22	Revision to Program Exceptions section.
06/10/23	Review: position statements and references updated.
08/21/23	Update to Program Exceptions section.
06/15/24	Review; no change in position statement. Updated references.
06/15/25	Review; deleted no history of breast cancer. Updated references.