

Lymphadenopathy on Extent-of-Disease Breast MRI: Proposed Audit Metrics

Ethan O. Cohen, MD, Hilda H. Tso, DO, Kyungmin Shin, MD, Sarah R. Martaindale, MD, Ashley C. Bragg, MD, Kanchan A. Phalak, MD, Rachel E. Perry, MD, Jia Sun, PhD, Jessica W. T. Leung, MD

<https://doi.org/10.2214/AJR.25.34159>

Accepted: December 15, 2025

Article Type: Research Letter

Article Section: Breast Imaging

The complete title page, as provided by the authors, is available at the end of this article.

ACCEPTED
MANUSCRIPT

Recommended citation:

Cohen EO, Tso HH, Shin K, et al. Lymphadenopathy on Extent-of-Disease Breast MRI: Proposed Audit Metrics. *AJR* 2025 Dec 24 [published online]. Accepted manuscript. doi:10.2214/AJR.25.34159

A supplement to this article is available at:

https://www.ajronline.org/doi/suppl/10.2214/AJR.25.34159/suppl_file/25_34159_suppl.pdf

The publication of this Accepted Manuscript is provided to give early visibility to the contents of the article, which will undergo additional copy-editing, typesetting, and review before it is published in its final form. During the production process, errors may be discovered that could affect the content of the Accepted Manuscript. All legal disclaimers that apply to the journal pertain. The reader is cautioned to consult the definitive version of record before relying on the contents of this document.

Preoperative breast MRI performed for extent-of-disease (EOD) evaluation not only helps detect occult synchronous breast malignancy but also provides detailed assessment for potential lymph node (LN) metastases [1]. The BI-RADS v2025 Manual recommends auditing EOD MRI examinations for abnormal interpretation rate (AIR), PPV, and additional cancer detection rate (CDR) [2]. Such audits are not recommended to evaluate performance in detecting abnormal axillary and internal mammary (IM) LNs (hereafter, lymphadenopathy), which may represent false-positive results. However, lymphadenopathy on EOD MRI is clinically relevant, for example upstaging disease and/or guiding treatment decisions. This study aimed to use BI-RADS-based audit metrics to evaluate the performance of EOD MRI in detecting lymphadenopathy.

The institutional review board approved this HIPAA-compliant retrospective study, waiving written informed consent. The reports of all breast MRI examinations from January 1, 2021 to December 31, 2022 were reviewed to identify EOD MRI examinations performed in patients with newly diagnosed breast cancer; such patients formed the final study sample. All patients were included in another study evaluating feasibility of EOD MRI audits [3]. The institution performs EOD MRI only after diagnostic mammography and, in patients with invasive breast cancer, staging whole-breast ultrasound that includes evaluation of ipsilateral regional (i.e., axillary levels 1-3 and IM) nodal basins. Axillary or IM lymphadenopathy detected on ultrasound undergoes image-guided biopsy before EOD MRI.

The reports of the EOD MRI examinations were reviewed to identify those reporting lymphadenopathy. The presence of one or more reported abnormal LNs at a given nodal basin was considered to represent a single finding of lymphadenopathy for that basin. The EMR was reviewed to identify 1-year lymphadenopathy outcomes. The following metrics were calculated to parallel the BI-RADS v2025 EOD MRI audit [2]: AIR for lymphadenopathy (AIR_{LN}), PPV2 for lymphadenopathy (PPV_{2LN} , percentage of lymphadenopathy classified as metastatic by image-guided biopsy or imaging follow-up), PPV3 for lymphadenopathy (PPV_{3LN} , percentage of biopsied lymphadenopathy yielding metastatic results), and additional CDR for lymphadenopathy (additional CDR_{LN} , metastatic

lymphadenopathy diagnosed beyond pre-examination known disease per 1000 examinations). The Supplemental Methods provide additional details.

Of 7196 breast MRI examinations performed during the study period, 1533 were EOD examinations. These examinations were performed in 1533 unique patients (mean age, 55.2 years), forming the final study sample. Of the EOD MRI examinations, 53 (3.5%) revealed lymphadenopathy. Table S1 summarizes patient and tumor characteristics for all cases and those with lymphadenopathy; Table 1 summarizes lymphadenopathy characteristics and outcomes; Figure 1 summarizes patient selection and outcomes stratified by overall BI-RADS categories on EOD MRI. Forty-six cases of lymphadenopathy were characterized by biopsy or further imaging; seven were lost to follow-up. AIR_{LN} was 3.46% (53/1533 [95% CI: 2.55-4.37%]), PPV_{2LN} was 13.21% (7/53 [95% CI: 4.10-22.32%]), PPV_{3LN} was 26.09% (6/23 [95% CI: 8.14-44.04%]), and additional CDR_{LN} was 4.57 per 1000 ($1000 \times 7/1533$ [95% CI: 1.20-7.94]).

Universally adopted and validated criteria for LN evaluation on breast MRI are lacking, contributing to interreader variability and potentially suboptimal diagnostic performance. One study reported varying reader accuracy in detecting LN metastases on breast MRI using the Node Reporting and Data System [1]. Another study reported lower radiologist performance in detecting lymphadenopathy than in detecting the primary lesion's quadrant or multiplicity for an abbreviated MRI protocol [6]. Such challenges support the utility of including lymphadenopathy-related metrics within quality improvement efforts.

The observed PPV_{3LN} was within the suggested range for PPV3 for screening mammography audits [2]. However, low percentages of EOD MRI examinations showed lymphadenopathy or yielded pathologically confirmed LN metastases. We believe that such results are acceptable given the management impact of precise nodal staging. Benchmarks are unavailable for lymphadenopathy-focused performance reviews as have been established for screening mammography audits [2]. Thus, thresholds would need to be determined if EOD MRI audits were to include lymphadenopathy-related metrics.

Study limitations include the paucity of lymphadenopathy on EOD MRI, review of only MRI reports, cases of lymphadenopathy lost to follow-up, and institutional workflow incorporating regional nodal staging with ultrasound before EOD MRI, limiting generalizability.

In conclusion, on EOD MRI examinations, AIR for lymphadenopathy was 3.46%, PPV of biopsied lymphadenopathy was 26.09%, and 4.57 cases of additional metastatic lymphadenopathy were diagnosed per 1000 examinations. Inclusion of lymphadenopathy-related metrics could aid quality improvement efforts and warrants consideration for future revisions to EOD MRI audit recommendations.

ACCEPTED
MANUSCRIPT

REFERENCES

1. Pediconi F, Maroncelli R, Pasculli M, et al. Performance of MRI for standardized lymph nodes assessment in breast cancer: are we ready for Node-RADS? *Eur Radiol* 2024; 34:7734-7745
2. Newell M, Destounis S, Leung J, DeMartini W, Lee C, Eby P. *ACR BI-RADS v2025 Manual*. Reston, VA: American College of Radiology, 2025
3. Cohen EO, Tso HH, Shin K, et al. Feasibility of Auditing Preoperative Breast MRI for Extent-of-Disease Evaluation Using the BI-RADS v2025 Manual. *Radiology* 2025; 317:e243803
4. Kinkel K, Helbich TH, Esserman LJ, et al. Dynamic high-spatial-resolution MR imaging of suspicious breast lesions: diagnostic criteria and interobserver variability. *AJR Am J Roentgenol* 2000; 175:35-43
5. Stoutjesdijk MJ, Fütterer JJ, Boetes C, van Die LE, Jager G, Barentsz JO. Variability in the description of morphologic and contrast enhancement characteristics of breast lesions on magnetic resonance imaging. *Invest Radiol* 2005; 40:355-362
6. Kadioglu ME, Metin Y, Metin NO, Tasci F, Ozdemir O, Kupeli A. The efficacy of abbreviated breast MRI protocols using 1.5 T MRI in the preoperative staging of newly diagnosed breast cancers. *Clin Imaging* 2023; 101:44-49

Table 1. Lymphadenopathy characteristics and outcomes among 53 EOD MRI examinations showing abnormal lymph nodes

Characteristic	Value
Location	
Ipsilateral axillary level 1	30
Ipsilateral axillary level 2	1
Ipsilateral axillary level 3	2
Ipsilateral axillary levels 1 and 2	1
Ipsilateral axillary levels 1 and 3	1
Ipsilateral internal mammary	8
Contralateral axillary level 1	7
Contralateral internal mammary	3
Outcome	
Benign by biopsy	17
Benign by 1-year ultrasound follow-up	22
Metastatic by biopsy ^a	6
Metastatic by FDG PET/CT ^b	1
Lost to follow-up	7
Treatment impact from detection of metastatic lymph nodes by EOD MRI	
Additional chest wall radiation	3
ALND rather than SLNB	1
No change	2
Not applicable (patient death before treatment)	1

Note—ALND = axillary LN dissection, EOD = extent-of-disease, LN = lymph node, SLNB = sentinel LN biopsy.

^aFour cases of ipsilateral axillary lymphadenopathy and two cases of ipsilateral internal mammary lymphadenopathy

^bSingle case of ipsilateral internal mammary lymphadenopathy

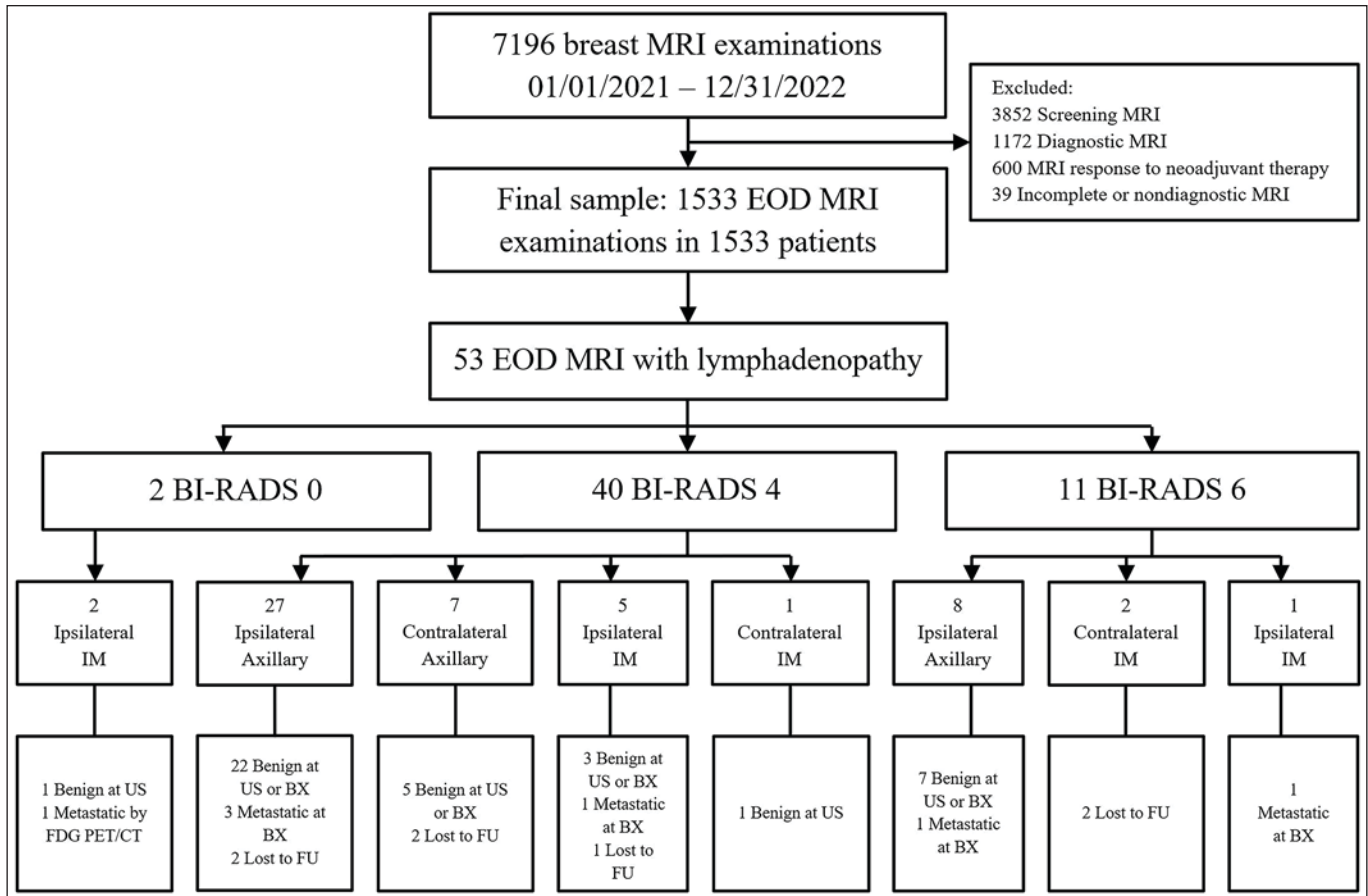


Fig. 1—Flowchart demonstrating patient selection process and outcomes for lymphadenopathy reported on EOD MRI stratified by examination’s overall BI-RADS assessment category (assigned at discretion of interpreting radiologist). DCIS = ductal carcinoma in situ, EOD = extent-of-disease, IDC = invasive ductal carcinoma, ILC = invasive lobular carcinoma, IM = internal mammary, IMC = invasive mammary carcinoma; US = ultrasound; FU = follow-up.

TITLE: Lymphadenopathy on Extent-of-Disease Breast MRI: Proposed Audit Metrics

AUTHORS (in order):

Ethan O. Cohen, MD¹ (not a trainee)

Hilda H. Tso, DO¹

Kyungmin Shin, MD¹

Sarah R. Martaindale, MD¹

Ashley C. Bragg, MD¹

Kanchan A. Phalak, MD¹

Rachel E. Perry, MD¹

Jia Sun PhD²

Jessica W.T. Leung, MD¹

1. Department of Breast Imaging, Division of Diagnostic Imaging
The University of Texas M.D. Anderson Cancer Center
1515 Holcombe, Unit 1350
Houston, TX USA 77030
2. Department of Biostatistics
The University of Texas MD Anderson Cancer Center
Unit 1411
PO Box 301402
Houston, TX USA 77030

Corresponding Author:

Ethan O. Cohen, MD

Phone: 214.514.6694

Email: ECohen@MDAnderson.org

X/TWITTER:

Cohen: @ECohenMD

Tso: @DrHildaTso

Shin: None

Martaindale: None

Bragg: None

Phalak: @KanchanPMD

Perry: None

Sun: None

Leung: @DrJessicaLeung

ARTICLE TYPE: Research Letter

FUNDING: None

DISCLOSURES: No disclosures relevant to the subject matter of this article.

ACKNOWLEDGEMENTS: The authors would like to acknowledge support by the NIH/NCI under award number P30 CA016672.