

| 6 | | | | | | | | | | | | _ | |
|--|--------------|-------------------|---|--|------------------|-------------|--------------|---------------------------|--|------------------------------|-----------------|-------------|-------------|
| ▶ OB Ultra | sound Med | lical History | | | | | | | | | | | |
| ▼ Oncology Member Medical History | | | | | | | | | | | | | |
| Oncology | Member H | listory | | | | | | | | | | | |
| Sequence # | Details | Date of Diagnosis | Cell Type & Location | Stage @ Diagnosis | Current Stag | e Treatment | History Date | S Treatment Type and Site | <u>Date of Diagnosis</u> for Recurrence or <u>Metastasis</u> | Site of Recurr Metastasis | ence or | Comme | <u>ents</u> |
| | | | (| | | | | | | | | | |
| Oncology Member History Details Find View All First 1 of 1 1 Last | | | | | | | | | | | | | |
| Sequence | # 1 Ye | ear of Diagr | nosis Month of | Diagnosis | Day of Diagnosis | Cell T | ype & Locat | ion | | | | | |
| Stag | ge @ Diagr | nosis | | | | | Current Sta | age | | | | | |
| Type of Treatment Dates Treatment Dates | | | | | | | | | | | | | |
| Recurrence or Metastasis Date of Diagnosis Comments | | | | | | | | | | | | | |
| ▼ CPT, ICD, Guideline | | | | | | | | | | | | | |
| CPT Info Member Rational Text | | | | | | | | | | | | | |
| Dup Service | ce Group Nam | ne Unit CPT | Description | * <u>Status</u> | Rationa | <u>le</u> | Repost Pro | vider Language | Member Language | Modifier | <u>BodyPart</u> | Description | |
| | | 1 770 | Magnetic resonance breast, without and material(s), includin Aided Detection (C/ lesion detection, chi and pharmacokineti when performed; bil | with contrast g Computer- D real-time aracterization, c analysis), | g 🔻 | a | Repost | | | | | | + = |

Q Repost

∨ 35

Withdrawn

SOCCPT Q SOCCPT

| ▼ Activity History | | | | | | | | | |
|--------------------|-------------|--|---------------|--------------|---------|------------|---------|----------------------|-------------------|
| | <u>Step</u> | Activity | <u>Status</u> | <u>Start</u> | | <u>End</u> | | Assigned Group | |
| | 10 | Initiate Request | 3 - Completed | 02/23/2024 | 11:34AM | 02/23/2024 | 11:34AM | Intake Workgroup | WEBUSER WEBUSER |
| | <u>20</u> | Give Verbal Recap, If Approved - Do Not Send To IVR | 3 - Completed | 02/23/2024 | 11:34AM | 02/23/2024 | 11:34AM | Intake Workgrou | WEBUSER WEBUSE |
| | <u>21</u> | Pending Additional Information | 4 - Canceled | 02/23/2024 | 12:17PM | | | Pending Addition Q | Q |
| | <u>30</u> | RN Review - Give Verbal Recap, If Approved - Do Not Send To IVR | 4 - Canceled | | | | | RN Workgroup Q | |
| | <u>40</u> | MD Review - Give Verbal Recap, If Approved - Do Not Send To IVR | 4 - Canceled | | | | | MD Priority Work | a |
| | <u>50</u> | NU Wrap Up - No Verbal Notification Required | 4 - Canceled | | | | | NU No Verbal No Q | Q |
| | <u>60</u> | Notify Member of Decision - SilverLink | 4 - Canceled | | | | | NU Member Noti | Q |
| | <u>70</u> | URGENT Initiate Request | 4 - Canceled | | | | | Intake Workgrou | Q |
| | 80 | URGENT Give Verbal Recap, If Approved - Do Not Send To IVR | 4 - Canceled | | | | | Intake Workgrou | |
| | 90 | URGENT RN Review - Give Verbal Recap, If Approved - Do Not Send To IVR | 2 - Started | | | | | RN Workgroup Q | Lyndon Delafuente |
| | 100 | URGENT MD Review - Give Verbal Recap, If Approved - Do Not Send To IVR | 1 - Queued | | | | | MD Main Workgr Q | |
| | <u>110</u> | URGENT NU Wrap Up | 1 - Queued | | | | | Eliza Notification Q | Q |
| | <u>120</u> | URGENT Notify Provider of Approval | 1 - Queued | 1 | | | | Eliza Notification Q | Lakita DENT NU |
| | <u>130</u> | Urgent Notify Member of Decision - SilverLink | 1 - Queued | | : | | | NU Member Noti | |

▼ Member History Information

Claims Summary

Medical Status: Pending Created: 02/23/2024 Member: Fornarelli, Rosamaria

SO Status: Open Auth Start: Physician: DUCKWORTH, APRIL M

Auth End: Facility: MIDSTATE RADIOLOGY ASSOCIATES LLC

| <u>Status</u> | <u>CPT</u> | | Rationale |
|---------------|------------|---|-----------|
| Pending | 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 | |
| Withdrawn | SOCCPT | SOCCPT | 35 |

| ICD Version | ICD ID | |
|-------------|---------|--|
| 10 | C50.912 | MALIGNANT NEOPLASM OF UNSPECIFIED SITE OF LEFT FEMALE BREAST |
| 10 | Z17.0 | ESTROGEN RECEPTOR POSITIVE STATUS [ER+] |
| 10 | Z51.11 | ENCOUNTER FOR ANTINEOPLASTIC CHEMOTHERAPY |

02/23/2024 12:17PM

ctammalwar

UPADNotes UPADNotes 02/23/2024 11:35AM webuser

77049 Is

This request is not in scope for real time claims lookups for conservative therapy. No claims were found for diagnostic ultrasound. This request is not in scope for real time claims lookups for X-rays.

The medical record for this patient is required to complete medical necessity review. This request will be pended until relevant medical records are uploaded at eviCore.com. (If the medical record is not currently attached to this case, DO NOT transfer to nurse) Cigna - ESNTLPLUS

Is this request for a customer who is currently enrolled in an approved oncology clinical trial OR to maintain compliance with an ongoing oncology clinical trial protocol?: No

Is this case Routine/Standard?: Yes

The medical record for this patient is required to complete medical necessity review.

signs and symptoms indicating the exam

br/>-Prior diagnostic studies with results (e.g. imaging studies or biopsies)

br/>-Prior management including conservative therapies

br/>-Medications with dose and duration

br/>>br/>-How would you like to proceed?:
Continue to documentation upload

Are you ready to upload documentation now?: No, I will upload at a later time. Please note that faxes received by eviCore may take up to 24 hours to process. Web uploaded documents have faster processing.

REQUEST CLINICAL INFO - UPADS Survey

02/23/2024 11:34AM webuser

The medical record for this patient is required to complete medical necessity review. This request will be pended until relevant medical records are uploaded at eviCore.com.

Str/>Medical records include:

Current signs and symptoms indicating the exam

Prior diagnostic studies with results (e.g. imaging studies or biopsies)

Prior management including conservative therapies

Medications with dose and duration

Web Portal Attestation Note

02/23/2024 11:34AM

webuser

The web user attested that this case was not urgent.

Hartford HealthCare Hospital of Central Connecticut Radiology Imaging Result

| Name: | DOB: | Sex: Female | Patient Class: Outpatient |
|--|------------------------|------------------|--|
| Procedures Performed: | Exam Date and Time: | Reason for Exam: | Dlagnosis: |
| US Breast diagnostic limited- Bilateral | 02/19/2024 11:39 AM | bilat lesions | Abnormal mammogram |
| | | | PCP/CC Providers: Welschedel, Anne-Katrin |
| | | | |

DIAGNOSTIC ULTRASOUND

CLINICAL INFORMATION: Diagnostic evaluation of bilateral breast findings detected on prior outside ultrasound.

COMPARISON: 1/25/2024

TECHNIQUE: Targeted bilateral ultrasound was performed.

FINDINGS:

Right:

At the 11:00 axis, 6 cm from the nipple, there is a $0.4 \times 0.3 \times 0.4$ cm hypoechoic mass with posterior acoustic enhancement and no vascularity. This was previously recommended for biopsy.

Left:

In the upper inner quadrant no correlate for mammographic focal asymmetry is seen.

IMPRESSION:

- Right breast 11:00 axis mass, previously recommended for biopsy.
 Patient is scheduled for same day blopsy.
- No correlate for a left breast mammographic finding. Patient is scheduled for same day stereotactic biopsy.

ASSESSMENT:

BI-RADS 4 - Suspicious.

RECOMMENDATION: Patient is scheduled for same day ultrasound and stereotactic biopsies. Findings and recommendations were discussed with the patient at the time of visit.

Our office contacts patients directly to arrange additional mammographic views, supplemental ultrasounds, and short interval mammographic follow-up.

Signed By: Kudrat Gill, MD on 2/19/2024 10:52 AM

Ordered On 2/19/2024 9:56 AM EST

Surgical Pathology Report

PATIENT NAME:

MED. REC. #:

ACCOUNT #: 100235865692 DOB (AGE):

DATE OBTAINED: 2/19/2024 LOCATION:

DATE RECEIVED: 2/19/2024 SUBMITTING PROVIDER: APRIL M. DUCKWORTH, MD

DATE REPORTED: 2/21/2024 CC: KUDRAT GILL, MD

DIAGNOSIS

A. CORE BIOPSY RIGHT BREAST AT 11:00: BENIGN BIPHASIC NEOPLASM, FAVOR BENIGN PHYLLODES TUMOR. SEE COMMENT.

B. CORE BIOPSY LEFT BREAST ASYMMETRY AT 11:00: INVASIVE MAMMARY CARCINOMA, DUCTAL TYPE, GRADE 1, 4.0 MM. IN GREATEST DIMENSION.

CANCER CASE SUMMARY, BREAST: INVASIVE MAMMARY CARCINOMA

Procedure:

CORE BIOPSY.

Tumor Site:

LEFT BREAST, 11:00.

Histologic Type:

INVASIVE DUCTAL CARCINOMA.

Core Length Of Tumor:

4.0 MM.

Tumor Grade (Nottingham):

GRADE I (NOTTINGHAM SCORE 5 OF 9).

Tubule Formation

2.

Nuclear Pleomorphism Mitotic Count 2.

Lymphovascular Invasion:

NOT IDENTIFIED.

Perineural Invasion:

NOT IDENTIFIED.

Microcalcifications:

PRESENT.

In Situ Component:

NOT IDENTIFIED.

Additional Findings:

Immunoperoxidase stain technical preparations were performed at Hartford Hospital Laboratories and interpreted by this pathologist as indicated.

Smooth muscle myosin: Loss of myoepithelial cell staining

Estrogen Receptor (SP1):

Percentage of cells exhibiting nuclear staining:

100%

Intensity of staining:

Strong

Interpretation:

POSITIVE

Progesterone Receptor (PR-16):

Percentage of cells exhibiting nuclear staining:

95%

intensity of staining:

Strong

Interpretation:

POSITIVE

HER2 IHC (EP3):-----Interpretation: NEGATIVE Score: 1+

All controls show appropriate reactivity.

CAP/ABCO HER2 Scoring Criteria Ref: Wolff AC, Hammond ME, Hicks DG, et al. Recommendations for human epidermal growth factor receptor 2 testing in breast cancer: American Society of Clinical Oncology -College of American Pathologists (ASCO/CAP) Clinical Practice Guideline Update (Arch Pathol Lab Med. 2014;138;241–256; doi: 10.5858/arpa.2013-0953-SA)

This test was developed and its performance characteristics determined by HCC Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is cartified under the Clinical Laboratory improvement Amendments of 1986 (CLIA-88) as qualified to perform high complexity clinical laboratory testing." PMTHC

**Electronically Signed Out On 2/21/2024 14:59 **

SUSAN PARKER, MD

COMMENT

A. Core biopsy right breast at 11:00: A Ki67 IHC stain performed on part A shows increased stromal proliferative activity (~3-5%). The control shows appropriate reactivity. The histologic features favor benign phyllodes tumor.

Dr. A. Vdovenko has also reviewed representative slides from parts A and B.

The toels used in the work-up of this specimen may include "Analyta-Specific Reagents" (ASRs). The Immunopathology/Morphologic Proteomics Laboratory at Hartford Hospital has established the performance characteristics of these reagents. They have not been cleared or approved by the United States Food and Drug Administration (FDA): however, the FDA has determined that such clearance or approval is not necessary for their use. All positive controls show appropriate immunoreactivity.

Clinical Information:

Right breast mass Left breast asymmetry Abnormal mammogram Mass of right breast, unspecified quadrant

Tissue(s) Submitted:

A: Core biopsy right breast at 11:00

B: Core biopsy left breast asymmetry at 11:00

Gross Description:

A. Core biopsy right breast at 11:00: Received in formalin in a specimen jar labeled right breast 11:00 and an accompanying specimen requisition slip labeled right beast 11:00 BiRads 4, 0.5 cm mass size are two cylindrical, fibrofatty, 1.1 x 0.2 cm tissue cores which are filtered and submitted in toto. Per the accompanying specimen requisition slip, the specimen requisition slip states three tissue cores are collected, only two discrete tissue cores are grossly identified with a few finely fragmented portions of fibrofatty soft tissue. The specimen is filtered, the specimen is submitted in toto designated A.

Per the accompanying specimen requisitions slip, part A right breast 11:00 is collected at 11:00 AM, placed in formalin at 11:10 AM on 02/192024, cassetted and placed in 10% formalin that same day, processor 2 - daily run.

B. Core biopsy left breast asymmetry at 11:00: Received in formalin labeled left asymmetry with an accompanying specimen requisition slip labeled left beast 11:00 asymmetry is a 4.0 x 4.0 x 0.3 cm aggregate of fragmented, irregular-to-cylindrical portions of fibrofatty soft tissue which are divided into two cassettes, filtered, and submitted in toto designated B1 and B2.

Per the accompanying specimen requisitions slip, part B left breast 11:00 is collected at 11:41 AM, placed in formalin at 11:43 AM on 02/192024, cassetted and placed in 10% formalin that same day, processor 2 - daily run.