

Name: _____ DOB: ____/____/____ Lead Region: _____

BREAST

4. Breast Risk Information and History (Yes/No answers should be chosen if risk assessed and determined by provider)

High Risk for Breast Cancer: Yes No Not Assessed/Unknown Symptoms: Yes No

Has client ever had: breast cancer? Yes No a previous mammogram? Yes No Date of previous mammogram: ____/____/____

5. Clinical Breast Exam

CBE Result: Not performed Normal/Benign Abnormality Suspicious for Cancer

CBE Provider #: _____
CBE Screening Date: ____/____/____
CBE Paid by FBCCEDP: Yes No

Additional Breast Procedures for CBE
 Additional procedures needed or planned
 Additional procedures not needed or planned

6. Mammogram

Mam Provider #: _____
Mam Screening Date: ____/____/____
Mam Paid by FBCCEDP: Yes No

Indication for Mammogram

- Screening
- Diagnostic
- Non-program mammogram.
Referred in for diagnostic evaluation
Breast Diagnostic Referral Date: ____/____/____
- No mammogram
- No breast service
- Unknown

Mammogram Result

- Negative (BI-RADS 1)
- Benign Finding (BI-RADS 2)
- Probably Benign/STFU suggested (BI-RADS 3)
- Unsatisfactory
- Result Pending
- Result unknown, presumed abnormal, mam from non-funded source

Suspicious Abnormality (BI-RADS 4)
 Highly Suggestive of Malignancy (BI-RADS 5)
 Need evaluation or film comparison (BI-RADS 0)

Additional Procedures for Mammogram

- Additional procedures needed or planned
- Additional procedures not needed or planned
- Need or plan for additional procedures not yet determined.

Next mammogram date: ____/____/____

6. Screening MRI (high-risk only)

MRI Pre-Authorization Date: _____
Central Office Nurse: _____

Screening MRI Provider#: _____
Screening MRI Date: ____/____/____
Screening MRI Paid by FBCCEDP: Yes No

Screening MRI Result

- Negative (BI-RADS 1)
- Benign Finding (BI-RADS 2)
- Probably Benign indicated (BI-RADS 3)
- Unsatisfactory
- Result Pending
- Not done

Suspicious (BI-RADS 4)
 Highly Suggestive of Malignancy (BI-RADS 5)
 Known Malignancy (BI-RADS 6)
 Need Additional Imaging Evaluation (BI-RADS 0)

Additional Procedures for Screening MRI

- Additional procedures needed or planned
- Additional procedures not needed or planned
- Need or plan for additional procedures not yet determined.

Name: _____ DOB: ____/____/____ Lead Region: _____

8. Additional Breast Procedures

<u>Breast Imaging Procedures</u>	<u>Date</u>	<u>Paid</u>		<u>Provider Number</u>
		<u>Yes</u>	<u>No</u>	
<input type="checkbox"/> Additional Mammographic Views	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Ultrasound	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Film Comparison	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____

MRI Pre-Authorization Date:	_____	Central Office Nurse:	_____
<input type="checkbox"/> Magnetic Resonance Imaging	_____	<input type="checkbox"/>	<input type="checkbox"/>

Check if additional page 3, Section 7. Additional Breast Procedures, is needed to document more than one imaging procedure:

Final Imaging Outcome

- Negative (BI-RADS 1)
- Benign Finding (BI-RADS 2)
- Probably Benign/STFU suggested (BI-RADS 3)
- Suspicious abnormality (BI-RADS 4)
- Highly suspicious of malignancy (BI-RADS 5)
- Known Malignancy (BI-RADS 6)
- Results pending

Final Imaging Date: _____

<u>Breast Diagnostic Procedures</u>	<u>Date</u>	<u>Paid</u>		<u>Provider Number</u>
		<u>Yes</u>	<u>No</u>	
<input type="checkbox"/> Repeat Breast Exam/ <input type="checkbox"/> Surgical Consultation	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Repeat Breast Exam/ <input type="checkbox"/> Surgical Consultation	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Biopsy	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Repeat Biopsy (comment required)	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Fine Needle/Cyst Aspiration	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____

Name: _____ DOB: ____/____/____ Lead Region: _____

9. Breast Final Diagnosis

Status of Breast Final Diagnosis

- Work-up complete
- Work-up pending

<input type="checkbox"/> Lost to follow-up - not otherwise specified	<input type="checkbox"/> Refused - not otherwise specified
<input type="checkbox"/> Lost to follow-up - deceased	<input type="checkbox"/> Refused - ineligible due to income or insurance
<input type="checkbox"/> Lost to follow-up - moved	
<i>If status of final diagnosis is lost to follow-up or refused, complete comments section below.</i>	

Final Breast Diagnosis

- Breast Cancer Not Diagnosed
- Carcinoma In Situ, Other
- Invasive Breast Cancer
- Lobular Carcinoma In Situ (LCIS) (Stage 0)
- Ductal Carcinoma In Situ (DCIS) (Stage 0)
- Atypical Ductal Hyperplasia (ADH)

Left	Right
<input type="checkbox"/>	<input type="checkbox"/>

Final Diagnosis Date: _____

Diagnosis Facility: _____
(include Provider Number)

10. Breast Cancer Treatment Status

- Treatment started
- Treatment pending

<input type="checkbox"/> Lost to follow-up - not otherwise specified	<input type="checkbox"/> Refused - not otherwise specified
<input type="checkbox"/> Lost to follow-up - deceased	<input type="checkbox"/> Refused - ineligible due to income or insurance
<input type="checkbox"/> Lost to follow-up - moved	
<i>If status of final diagnosis is lost to follow-up or refused, complete comments section below.</i>	

Treatment Start Date: _____

Treatment Facility: _____

Comments

CENTRAL OFFICE USE ONLY – MEDICAID FOR BREAST CANCER				COMMENTS:
REFERRED TO MEDICAID FOR TREATMENT?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	MEDICAID REFERRAL DATE: _____	
PATIENT ENROLLED?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	MEDICAID APPROVAL DATE: _____	

CERVICAL

11. Cervical Cancer Risk Information and History (Yes/No answers should be chosen if risk assessed and determined by provider)

Risk for Cervical Cancer: Yes No Not Assessed/Unknown Previous Dx'd Cervical Cancer? Yes No

Previous Pap Test? Yes No Unknown Date of Previous Pap? _____

12. Pap

Indication for Pap

- Screening
- Surveillance
- Non-program Pap. Referred in for diagnostic evaluation
Cervical Diagnostic Referral Date: ___/___/___
- No Pap
- No cervical service
- Pap after primary HPV+
- Unknown

Pap Provider #: _____

Pap Screening Date: ___/___/___

Pap Paid by FBCCEDP: Yes No

Specimen Type: Conventional Smear Liquid Based

Specimen Adequacy: Satisfactory Unsatisfactory

Pap Result

- Negative for intraepithelial lesion or malignancy
- Infection/Inflammation/Reactive Changes
- Atypical squamous cells of undetermined significance (ASC-US)
- Low Grade SIL (including HPV changes)
- Other _____
- Unsatisfactory
- Result Pending

Atypical squamous cells cannot exclude HSIL (ASC-H)

High Grade SIL

Squamous Cell Carcinoma

Atypical Glandular Cells

Adenocarcinoma In Situ (AIS)

Adenocarcinoma

Result Unknown, presumed abnormal, Pap test from non-program funded source

Diagnostic Work-up Planned for Cervical Dysplasia or Cancer:

- Diagnostic work-up planned on basis of abnormal Pap test or pelvic exam
- Diagnostic work-up not planned
- Diagnostic work-up plan not yet determined

Next Pap date: ___/___/___

13. HPV

Indication for HPV

- Co-Test/Screening
- Reflex (follow-up test after screening Pap)
- Test not done
- Unknown

HPV Provider #: _____

HPV Screening Date: ___/___/___

HPV Paid by FBCCEDP: Yes No

HPV Result

- Positive with genotyping not done/Unknown
- Negative
- Positive with positive genotyping (types 16 or 18)
- Positive with negative genotyping (positive HPV, but not types 16 or 18)
- Unknown

Name: _____ DOB: ____/____/____ Lead Region: _____

14. CERVICAL DIAGNOSTIC PROCEDURES

Cervical Diagnostic Procedures	Date	Paid		Provider Number
		Yes	No	
<input type="checkbox"/> Colposcopy without biopsy	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Colposcopy with biopsy and/or ECC	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
<input type="checkbox"/> Other cervical procedures performed	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____
Please specify: _____				
<input type="checkbox"/> ECC alone	_____	<input type="checkbox"/>	<input type="checkbox"/>	_____

LEEP or Cold Knife Cone		Central Office Nurse:	_____
Pre-Authorization Date:	_____	Yes	No
<input type="checkbox"/> Diagnostic Cold Knife Cone (CKC)	_____	<input type="checkbox"/>	<input type="checkbox"/>
<input type="checkbox"/> Diagnostic LEEP	_____	<input type="checkbox"/>	<input type="checkbox"/>

15. CERVICAL FINAL DIAGNOSIS

Status of Cervical Final Diagnosis

- Work-up complete
- Work-up pending

<input type="checkbox"/> Lost to follow-up - not otherwise specified	<input type="checkbox"/> Refused - not otherwise specified
<input type="checkbox"/> Lost to follow-up - deceased	<input type="checkbox"/> Refused - ineligible due to income or insurance
<input type="checkbox"/> Lost to follow-up - moved	
<i>If status of final diagnosis is lost to follow-up or refused, complete comments section below.</i>	

Final Cervical Diagnosis

- Normal/Benign reaction/inflammation
- HPV/Condylomata/Atypia
- CIN 1/mild dysplasia (biopsy diagnosis)
- CIN 2/moderate dysplasia (biopsy diagnosis)
- CIN 3/severe dysplasia/Carcinoma in situ (Stage 0) or Adenocarcinoma in situ of the cervix (AIS) (biopsy diagnosis)
- Invasive cervical carcinoma (biopsy diagnosis)
- Other cancer diagnosis (only if patient has no cervix due to cervical cancer): _____
- Low grade SIL (biopsy diagnosis)
- High grade SIL (biopsy diagnosis)

Final Diagnosis Date: _____ **Diagnosis Facility:** _____
(include Provider Number)

16. Cervical Cancer Treatment Status

- Treatment started
- Treatment pending

<input type="checkbox"/> Lost to follow-up - not otherwise specified	<input type="checkbox"/> Refused - not otherwise specified
<input type="checkbox"/> Lost to follow-up - deceased	<input type="checkbox"/> Refused - ineligible due to income or insurance
<input type="checkbox"/> Lost to follow-up - moved	
<i>If status of final diagnosis is lost to follow-up or refused, complete comments below.</i>	

Treatment Start Date: _____ **Treatment Facility:** _____

Comments

CENTRAL OFFICE USE ONLY – MEDICAID FOR CERVICAL CANCER				COMMENTS:
REFERRED TO MEDICAID FOR TREATMENT?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	MEDICAID REFERRAL DATE: _____	
PATIENT ENROLLED?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	MEDICAID APPROVAL DATE: _____	