

Programa de Prevención y Detección Temprana de Cáncer de Mama y Cuello Uterino de Puerto Rico (PRBCCEDP)



National Breast
and Cervical Cancer
Early Detection
Program



CENTRO
COMPRESIVO
DE
CANCER
universidad de puerto rico

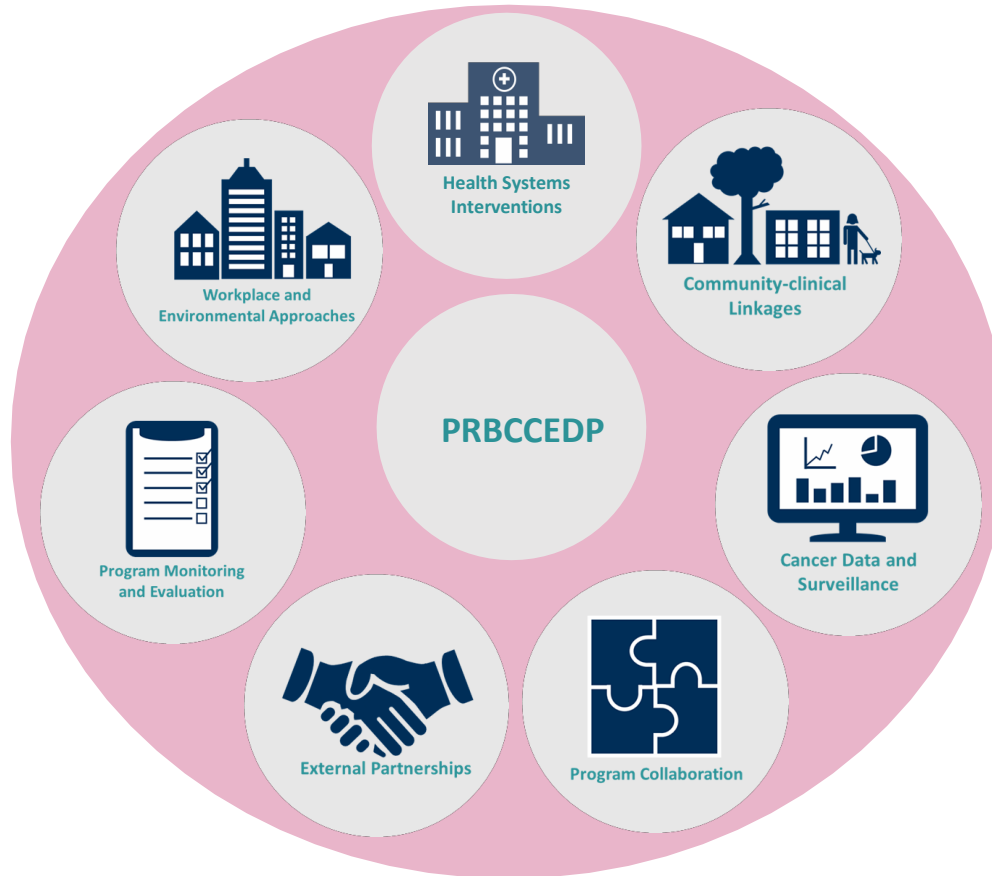
Trasfondo

- El Congreso de los Estados Unidos autorizó la creación del ***Programa Nacional de Detección Temprana de Cáncer de Mama y Cuello uterino*** (NBCCEDP) en el 1990.
- El NBCCEDP es administrado por la **División de Prevención y Control del Cáncer (DCPC)** de los **Centros para el Control y Prevención de Enfermedades (CDC)**.
- Desde el **1991**, el programa ha crecido hasta cubrir los **50 estados** de la unión, **5 territorios** (entre ellos Puerto Rico), el **Distrito de Columbia** y **11 tribus u organizaciones tribales**.
- Se estima que entre **7-13%** de las mujeres en los Estados Unidos son **elegibles para participar** del programa.

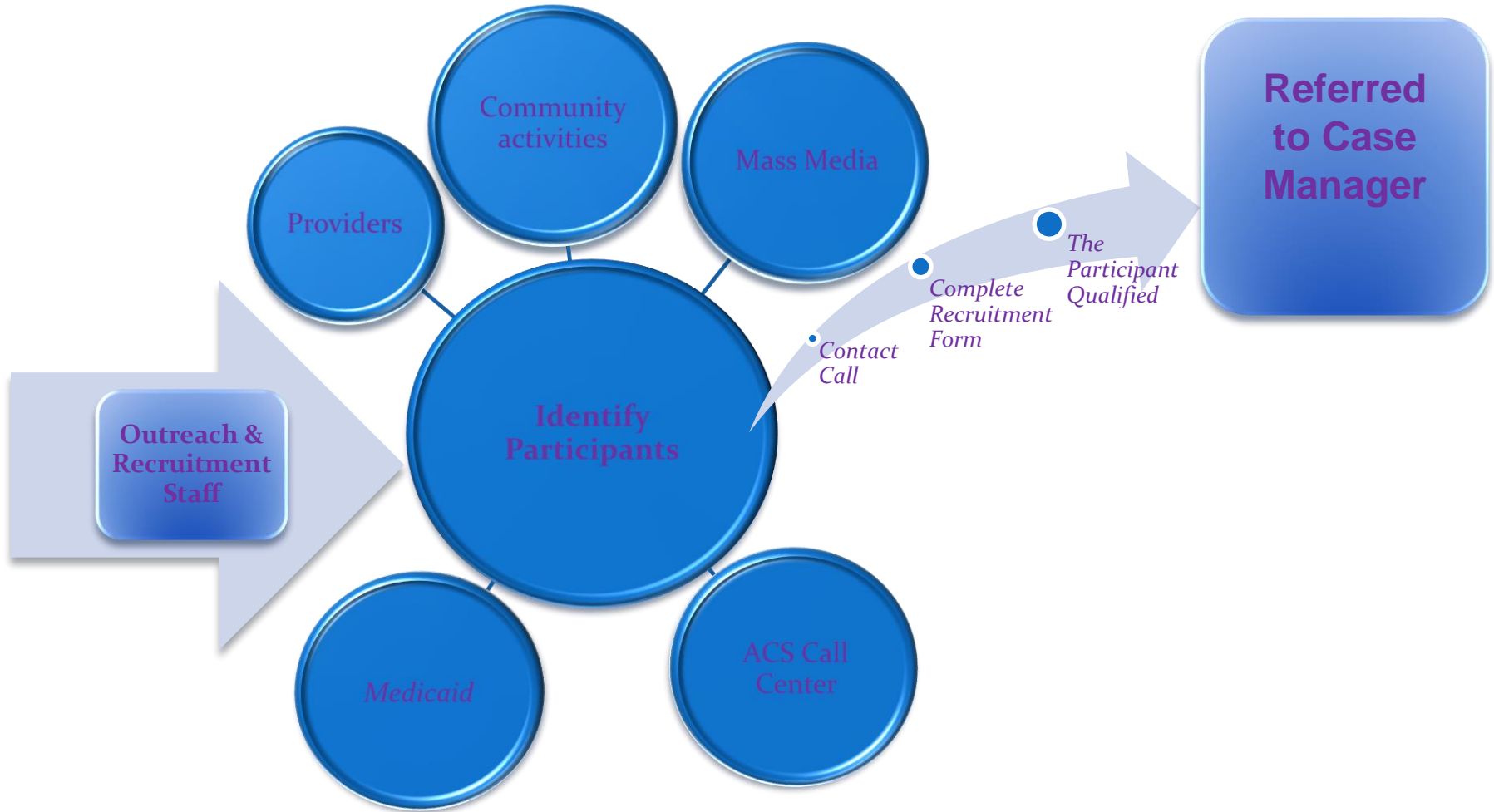
Trasfondo

- En Puerto Rico, desde el año **1991 hasta el 2005**, este programa estuvo adscrito al **Departamento de Salud**. Este era conocido como el Programa “*Compañeras*”.
- Desde el **2007** el programa es manejado por el **Recinto de Ciencias Médicas** de la Universidad de Puerto Rico.
- Actualmente, el NBCCEDP es parte del componente de servicios del **Centro Comprensivo de Cáncer** de la Universidad de Puerto Rico (UPRCCC).
- El programa en la isla lleva el nombre de “*Programa para la Prevención y Detección Temprana de Cáncer de Mama y Cuello Uterino de Puerto Rico*”.

Estrategias



Reclutamiento



Elegibilidad

- Mujeres de **21 a 64 años** de edad
- Clasifiquen entre el **200 y 250% de nivel de pobreza** de acuerdo a sus ingresos y número de personas en el núcleo familiar
- **No** califiquen para la **Reforma de Salud** y no puedan costear la cubierta de un **seguro de salud privado**
- Mujeres de **65 años o más** que **no** tengan **Medicare** o no cuentan con **Medicare Parte B**



Mamografía y
Papanicolaou
GRATIS
para mujeres elegibles
al programa

REQUISITOS:

- Estar entre las edades de 21 a 64 años
- No cualificar para la Reforma de Salud
- No tener plan médico privado
- Cumplir con los requisitos de ingreso económico
- Mujeres de 65 años o más que no cuentan con Medicare, o no tienen Medicare Parte B

Si desea saber si es elegible comuníquese al:

(787) 522-3265

Parámetros

- *Servicios para cernimiento/diagnóstico de Cáncer de Mama*

- El programa sigue como parámetros las guías y recomendaciones para la detección temprana de cáncer de mama publicadas por U.S. Preventive Services Task Force.
- La **prioridad** para estos servicios son las participantes de **50 a 64 años** de edad.
 - Estas deben de representar **no menos del 75% de todas las beneficiarias** de los servicios de mamografía.
- Las mujeres **menores de 50 años** pueden recibir los servicios de **mamografía**.
 - Estas **no deben de exceder el 25%** del total de las beneficiarias de los servicios de mamografía.
 - Las mujeres de **40 a 49 años asintomáticas** pueden beneficiarse de los servicios de mamografía, pero siempre **recordando la regla del 25%**.
 - Mujeres **menores de 40 años sintomáticas** pueden recibir servicios del programa (**examen clínico de mama, mamografía de diagnóstico y consulta de cirujano, de ser necesario.**)
- Las **nuevas guías de cáncer de mama** dictadas por el **United States Preventive Services Task Force (USPSTF, por sus siglas en ingles)** fueron publicadas en **enero de 2016**.

Parámetros

- *Servicios para cernimiento/diagnóstico de Cáncer de Cuello Uterino*
 - Mujeres entre las edades de 21 a 64 años de edad.
 - El programa sigue como parámetros las guías y recomendaciones para la detección temprana de cáncer de cuello uterino publicadas por la United States Preventive Services Task Force.
 - La prioridad son las participantes que nunca se han realizado un Pap.
 - Mujeres que desconocen conservar el cuello uterino luego de la histerectomía, se puede pagar una visita médica para realizar un examen pélvico y determinar si cuenta con el órgano.
 - A partir **de marzo de 2012** se comenzó a implementar la **nuevas guías** de cernimiento para cáncer de cuello uterino .

Guías de Cernimiento Cáncer de Mama

Population	Recommendation ¹
Women aged 40 to 49 with average risk	The decision to start screening mammography in women prior to age 50 years should be an individual one. Women who place a higher value on the potential benefit than the potential harms may choose to begin biennial screening between the ages of 40 and 49 years.
Women aged 50 to 74 with average risk	Biennial screening mammography is recommended.
Women aged 75 or older with average risk	Current evidence is insufficient to assess the balance of benefits and harms of screening mammography in women aged 75 years or older.
Women with dense breasts	Current evidence is insufficient to assess the balance of benefits and harms of adjunctive screening for breast cancer using breast ultrasonography, magnetic resonance imaging (MRI), digital breast tomosynthesis (DBT), or other methods in women identified to have dense breasts on an otherwise negative screening mammogram.
Women at higher than average risk	Women with a parent, sibling, or child with breast cancer are at higher risk for breast cancer and thus may benefit more than average-risk women from beginning screening in their 40s.
Additional issues relevant for all women	Current evidence is insufficient to assess the benefits and harms of digital breast tomosynthesis (DBT) as a primary screening method for breast cancer.

¹Siu AL; U.S. Preventive Services Task Force. Screening for breast cancer: U.S. Preventive Services Task Force recommendation statement. Annals of Internal Medicine 2016; 164(4):279-296.

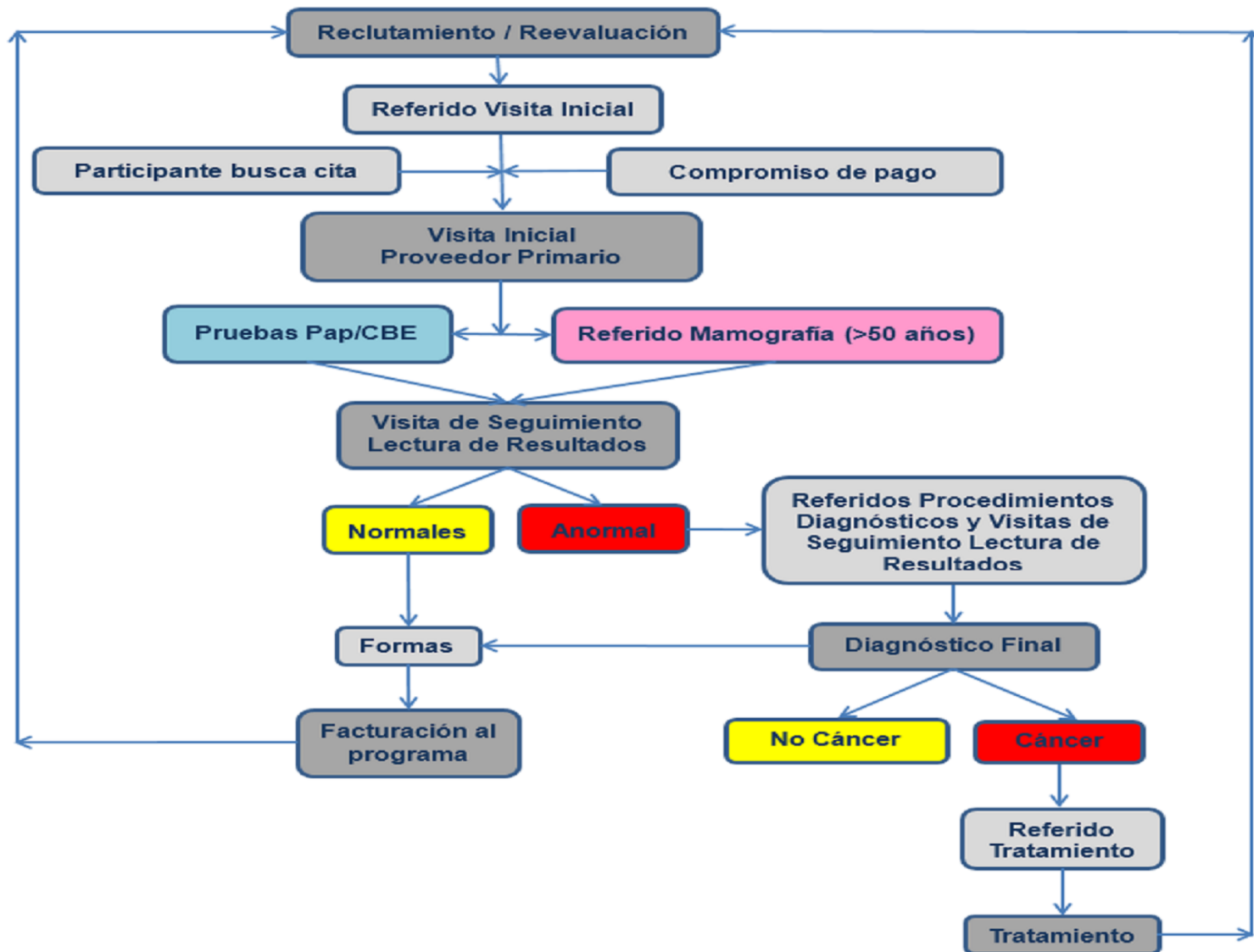
Guías de Cernimiento Cáncer de Cuello Uterino

Population	Recommendation ²
Women 21 to 65 (Pap Smear) or 30-65 (in combo with HPV testing)	The USPSTF recommends screening for cervical cancer in women age 21 to 65 years with cytology (Pap smear) every 3 years or, for women age 30 to 65 years who want to lengthen the screening interval, screening with a combination of cytology and human papillomavirus (HPV) testing every 5 years. See the Clinical Considerations for discussion of cytology method, HPV testing, and screening interval.
Women younger than 30 years, HPV testing	The USPSTF recommends against screening for cervical cancer with HPV testing, alone or in combination with cytology, in women younger than age 30 years.
Women younger than 21	The USPSTF recommends against screening for cervical cancer in women younger than age 21 years.
Women Older than 65, who have had adequate prior screening	The USPSTF recommends against screening for cervical cancer in women older than age 65 years who have had adequate prior screening and are not otherwise at high risk for cervical cancer. See the Clinical Considerations for discussion of adequacy of prior screening and risk factors.
Women who have had a hysterectomy	The USPSTF recommends against screening for cervical cancer in women who have had a hysterectomy with removal of the cervix and who do not have a history of a high-grade precancerous lesion (cervical intraepithelial neoplasia [CIN] grade 2 or 3) or cervical cancer.

²Cervical Cancer: Screening. U.S. Preventive Services Task Force. July 2015.

[http://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/cervical-cancer-screening?ds=1&s=cervical cancer screening](http://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/cervical-cancer-screening?ds=1&s=cervical%20cancer%20screening)

Flujograma de Servicios Clínicos



Navegación y Manejo

- *Los servicios ofrecidos tienen un **tiempo determinado para cumplir satisfactoriamente con las guías establecidas por el programa.***
- **Cáncer de Mama:**
 - **≤60 días :** intervalo de tiempo entre el **cernimiento inicial** y el **diagnóstico** de un cernimiento **anormal de mama.**
 - **≤60 días:** intervalo de tiempo entre **el diagnóstico** y el inicio del **tratamiento** para **cáncer de mama.**
- **Cáncer de Cuello Uterino:**
 - **≤90 días:** intervalo de tiempo entre el **cernimiento inicial** y el **diagnóstico** de un cernimiento **anormal de cuello uterino.**
 - **≤60 días:** intervalo de tiempo entre **el diagnóstico** y el inicio del **tratamiento** para **cáncer de cuello uterino invasivo.**
 - **≤90 días:** intervalo de tiempo **entre el diagnóstico** y el inicio del **tratamiento** para **neoplasia intraepitelial.**

Ciclos

- Ciclo de Cernimiento/Diagnóstico
 - Desde visita inicial hasta visita de diagnóstico
 - ≤ 60 días: Cáncer de Mama
 - ≤ 90 días: Cáncer de Cuello Uterino
- Si la participante comienza el ciclo pero no asiste a su última visita de diagnóstico para cerrarlo, el médico puede completar la forma clínica (incluyendo un comentario o razón para completar el ciclo sin la visita de diagnóstico) y facturar las visitas.

Servicios para Cáncer de Mama

● Cernimiento

- Examen Clínico de Mama (CBE)
- **Mamografías** de cernimiento
 - El programa no cubrirá métodos automatizados para realizar las mamografías: CAD y CT.

● Diagnóstico

- Mamografía diagnóstica (Vistas adicionales o “spot compression”) unilateral o bilateral
- Sonomamografía unilateral o bilateral
- Consulta de un cirujano
- Biopsias
- MRI – cumplir con los criterios establecidos para esta prueba

Servicios para de Cáncer de Cuello Uterino

- Cernimiento

- Examen pélvico
- Prueba de **Pap** (Convencional y “Thin Prep”)
 - El programa no cubrirá métodos automatizados para realizar los Pap.
- Prueba de **Pap** & Prueba de **HPV** de acuerdo a las guías de cernimiento para cáncer de cuello uterino (Algoritmos ASCCP).

- Diagnóstico

- Consultas ginecológicas
- Colposcopías: sin biopsia, con biopsia, con Endocervical Curettage (ECC)
- Endocervical Curettage (ECC) solo
- Loop Electrosurgical Excision Procedure (LEEP), Cold-Knife Conization (CKC) y Laser Conization.
- Otras biopsias (no colposcopia)

Formas de Cernimiento



University of Puerto Rico Comprehensive Cancer Center
Puerto Rico Breast and Cervical Cancer Prevention and Early Detection Program
PMB 371 PO Box 70344, San Juan PR 00936-8344
Phone: 787-772-8300 ext. 1116

Breast Cancer Screening Data Collection Form (Follow Cancer Screening Guidelines provided)

Program Use Only

Patient ID: _____

Cycle #: _____

A. Patient Information					
1a. Last Names	1b. First Name	1c. Initial	2. SSN	3. DOB	4. Age
5a. Postal Address	5b. Municipality	5c. State	5d. Zip Code	6. Phone Number	
7. Provider #	8. Record #	9. Municipality of Screening			
B. Breast Screening History					
10a. Has the patient had a mammogram before? If Yes, <input type="radio"/> Yes <input type="radio"/> No			11. Does the patient have breast implants? <input type="radio"/> Yes <input type="radio"/> No		
10b. Date of previous mammogram: _____			12. The patient reported breast symptoms? <input type="radio"/> Yes <input type="radio"/> No		
C. Breast Screening Tests					
13. CBE Date: _____		<input type="radio"/> Bloody / Serous Nipple Discharge <input type="radio"/> Nipple / Areolar Scarsness <input type="radio"/> Skin Dimpling / Retraction <input type="radio"/> Previous normal CBE in past 12 months – CBE not done today <input type="radio"/> CBE not done today – other / Unknown Reason <input type="radio"/> CBE refused			
14. CBE Results:		<input type="radio"/> Normal <input type="radio"/> Benign Finding <input type="radio"/> Discrete Palpable Mass – Suspicious for Cancer <input type="radio"/> Discrete Palpable Mass – Previously Diagnosed Benign			
15a. Purpose of the Initial Mammogram:		18. Date Initial Mammogram: _____		D. Diagnostic Procedures:	
<input type="radio"/> Routine screening mammogram <input type="radio"/> Evaluate symptoms, positive CBE, or previous abnormal mammogram <input type="radio"/> Already done by a non-program provider, patient referred in for diagnostic evaluation		19. Initial Mammogram Results:		20. Diagnostic Work-up Plan: <input type="radio"/> Planned <input type="radio"/> Not Planned	
15b. Date of Referral: _____		<input type="radio"/> Negative (BI-RADS 1) <input type="radio"/> Benign (BI-RADS 2) <input type="radio"/> Probably Benign (Short Interval follow-up suggested; BI-RADS 3) <input type="radio"/> Suspicious Abnormality (Consider Biopsy; BI-RADS 4) <input type="radio"/> Highly Suggestive of Malignancy (BI-RADS 5)		21. Additional Mammography Views: <input type="radio"/> Yes <input type="radio"/> No	
<input type="radio"/> Not done (Patient proceeded directly for other imaging or diagnostic workup) <input type="radio"/> Not done (Cervical record only)		<input type="radio"/> Assessment is Incomplete (Need additional imaging; BI-RADS 0) <input type="radio"/> Unsatisfactory (Cycle complete) <input type="radio"/> Unknown (Presumed Abnormal, mammogram from non-program provider) <input type="radio"/> Film Comparison Required		22. Ultrasound: <input type="radio"/> Yes <input type="radio"/> No	
16. Initial Mammogram Type: <input type="radio"/> Conventional <input type="radio"/> Digital		23. Film comparison to evaluate an incomplete assessment: <input type="radio"/> Yes <input type="radio"/> No		24. Bill to PRBCCEDP: <input type="radio"/> Yes <input type="radio"/> No	
17. Bill to PRBCCEDP: <input type="radio"/> Yes <input type="radio"/> No		25. Final Imaging Date: _____		26. Final Imaging Outcome: <input type="radio"/> Negative <input type="radio"/> Benign finding <input type="radio"/> Probably Benign (Short Interval follow-up suggested) <input type="radio"/> Suspicious Abnormality (Consider Biopsy) <input type="radio"/> Highly Suggestive of Malignancy <input type="radio"/> Unsatisfactory (Cycle complete) <input type="radio"/> Known Biopsy-Proven Malignancy	
27. Additional Diagnostic Procedures (Complete the Breast Cancer Diagnosis Data Collection Form):					
<input type="radio"/> Diagnostic Mammography <input type="radio"/> Consultant repeat CBE <input type="radio"/> Fine Needle Aspiration Biopsy <input type="radio"/> Surgical Consultation			<input type="radio"/> Large Core Needle Biopsy <input type="radio"/> Open Surgical Biopsy <input type="radio"/> Other procedure (Specify): _____		
28a. Follow-up: <input type="radio"/> 2 years <input type="radio"/> 1 year <input type="radio"/> Short-Term			28b. Specify Short-Term months: _____		
29. Comments: _____			31. Date: _____		
30. Provider's Name and Signature: _____			31. Date: _____		



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Cervical Cancer Screening Data Collection Form (Follow Cancer Screening Guidelines provided)

Program Use Only

Patient ID: _____

Cycle #: _____

A. Patient Information			
1a. Last Names	1b. First Name	1c. Initial	2. SSN
3. DOB	4. Age	5a. Postal Address	5b. Municipality
5c. State	5d. Zip Code	6. Phone Number	7. Provider #
8. Record #	9. Municipality of Screening		
B. Cervical Screening History			
10a. Has the patient had a prior Pap Test? If Yes, <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown		12a. Has the patient received an HPV vaccination? If Yes, <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	
10b. Date of prior (last) Pap test: _____		12b. Date of first HPV vaccination: _____	
11. Is there history of the following conditions? (Mark all that apply)		15c. Number of doses received: <input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3	
11a. Dysplasia/cervical cancer <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	11b. HPV <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	13. Is the patient post-menopausal? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	14. Is the patient pregnant? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown
11c. HIV <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	11d. Immune-compromised <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	15a. Has the patient had a hysterectomy? If Yes, <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown	15b. Was the hysterectomy performed for either cervical cancer or Neoplasia? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown
C. Cervical Screening Tests			
16. Test(s) Requested (Mark all that apply): Date of test(s): _____		19. Type of Pap Test (Specimen Type):	
<input type="checkbox"/> Pelvic exam: <input type="checkbox"/> Pap test: <input type="checkbox"/> HPV test:		<input type="radio"/> Conventional Smear <input type="radio"/> Liquid Based <input type="radio"/> Other <input type="radio"/> Unknown	
17. Pelvic exam results: <input type="radio"/> Normal <input type="radio"/> Abnormal pelvic <input type="radio"/> Abnormal-not suspicious for cancer <input type="radio"/> Abnormal-suspicious for cancer		20. Specimen Adequacy: <input type="radio"/> Satisfactory <input type="radio"/> Unsatisfactory <input type="radio"/> Unknown	
18a. Indication for Pap test to be performed on this screening:		21. Pap Test results:	
<input type="radio"/> Routine Pap Test <input type="radio"/> Patient under surveillance for a previous abnormal test <input type="radio"/> Already done by a non-program provider, patient referred in for diagnostic evaluation		<input type="radio"/> Negative <input type="radio"/> ASC-US <input type="radio"/> LSIL <input type="radio"/> HSIL <input type="radio"/> ASC-H <input type="radio"/> Squamous Cell Carcinoma <input type="radio"/> AGC <input type="radio"/> Adenocarcinoma <input type="radio"/> AIS <input type="radio"/> Result unknown, presumed abnormal Pap done by a non-program provider	
18b. Date of referral: _____		22. HPV Test Result: (If positive, specify type)	
<input type="radio"/> Not done, Patient proceeded directly for diagnostic work-up or HPV test <input type="radio"/> Not done, Breast record only		<input type="radio"/> Positive: <input type="radio"/> LR <input type="radio"/> HR <input type="radio"/> Unknown <input type="radio"/> Negative <input type="radio"/> Unknown <input type="radio"/> Not done	
25. Bill to PRBCCEDP: <input type="radio"/> Yes <input type="radio"/> No		26. Comments: _____	
27. Provider's Name and Signature: _____		28. Date: _____	
23. Diagnostic Work-up Plan: <input type="radio"/> Planned <input type="radio"/> Not Planned			
24a. Follow up: <input type="radio"/> Short term <input type="radio"/> Specify Short-Term months: _____			
<input type="radio"/> Pap in 1 year <input type="radio"/> Pap in 3 years <input type="radio"/> Pap in 5 years <input type="radio"/> Additional Diagnostic Procedures (Complete the Cervical Cancer Diagnosis Data Collection Form): <input type="checkbox"/> Gynecologic Consultation <input type="checkbox"/> Colposcopy w/o Biopsy <input type="checkbox"/> Colposcopy with Biopsy <input type="checkbox"/> Colposcopy with ECC <input type="checkbox"/> ECC (Only) <input type="checkbox"/> LEEP <input type="checkbox"/> CKC <input type="checkbox"/> Laser Coization <input type="checkbox"/> Other biopsy-not colposcopy <input type="checkbox"/> Other procedure (Specify): _____			

Formas de Diagnóstico



University of Puerto Rico Comprehensive Cancer Center
Puerto Rico Breast and Cervical Cancer Prevention and Early Detection Program
PMB 371 PO Box 70344, San Juan PR 00936-8344
Phone: 787-772-8300 ext. 1116

Breast Cancer Diagnosis Data Collection Form

Program Use Only

Patient ID:
Cycle #:

A. Patient Information					
1a. Last Names	1b. First Name	1c. Initial	2. SSN	3. DOB	4. Age
5a. Postal Address	5b. Municipality	5c. State	5d. Zip Code	6. Phone Number	
7. Provider #	8. Record #	9. Municipality of Diagnosis			
B. Diagnostic Procedures (Mark all that apply)					
10a. Diagnostic Mammography	<input type="checkbox"/>	10b. Date of Procedure	10c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
11a. Consultant-Repeat CBE	<input type="checkbox"/>	11b. Date of Procedure	11c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
12a. Fine Needle Aspiration Biopsy	<input type="checkbox"/>	12b. Date of Procedure	12c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
13a. Surgical Consultation	<input type="checkbox"/>	13b. Date of Procedure	13c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
14a. Large Core Needle Biopsy	<input type="checkbox"/>	14b. Date of Procedure	14c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
15a. Open Surgical Biopsy	<input type="checkbox"/>	15b. Date of Procedure	15c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
16a. Other Breast Procedures:	<input type="checkbox"/>	16b. Date of Procedure	16c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
C. Diagnosis Information					
17a. Status of Final Diagnosis:					
<input type="radio"/> Work-up Complete <input type="radio"/> Lost to Follow Up <input type="radio"/> Work-up Refused					
18. Final Diagnosis:					
<input type="radio"/> Breast Cancer not Diagnosed/Normal breast Tissue <input type="radio"/> Invasive Breast Cancer <input type="radio"/> Lobular Carcinoma In Situ (LCIS)-(Stage 0) <input type="radio"/> Ductal Carcinoma In Situ (DCIS)-(Stage 0) <input type="radio"/> Hyperplasia <input type="radio"/> Atypical Ductal Hyperplasia (ADH)		<input type="radio"/> Other Final Diagnosis (Specify): <div style="border: 1px solid black; height: 100px; width: 100%;"></div>			
D. Treatment Information					
19a. Status of Treatment:		19b. Date of Treatment:		20a. Follow-up:	
<input type="radio"/> Treatment Started <input type="radio"/> Treatment Pending <input type="radio"/> Treatment not Needed		<input type="radio"/> Treatment Refused <input type="radio"/> Lost to Follow-up (includes death)		<input type="radio"/> 2 years <input type="radio"/> 1 year <input type="radio"/> Short-Term 20b. Specify Short-Term months: <input type="text"/>	
21. Comments: <input type="text"/>					
22. Provider's Name and Signature:					23. Date:



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Cervical Cancer Diagnosis Data Collection Form

Program Use Only

Patient ID:
Cycle #:

A. Patient Information					
1a. Last Names	1b. First Name	1c. Initial	2. SSN	3. DOB	4. Age
5a. Postal Address	5b. Municipality	5c. State	5d. Zip Code	6. Phone Number	
7. Provider #	8. Record #	9. Municipality of Diagnosis			
B. Diagnostic Procedures (Mark all that apply)					
10a. Gynecologic Consultation	<input type="checkbox"/>	10b. Date of Procedure	10c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
11a. Colposcopy w/o Biopsy	<input type="checkbox"/>	11b. Date of Procedure	11c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
12a. Colposcopy with Biopsy	<input type="checkbox"/>	12b. Date of Procedure	12c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
13a. Colposcopy with ECC	<input type="checkbox"/>	13b. Date of Procedure	13c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
14a. Endocervical Curettage Only (ECC)	<input type="checkbox"/>	14b. Date of Procedure	14c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
15a. Loop Electrosurgical Excision Procedure (LEEP)	<input type="checkbox"/>	15b. Date of Procedure	15c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
16a. Cold-Knife Cone (CKC)	<input type="checkbox"/>	16b. Date of Procedure	16c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
17a. Laser Conization	<input type="checkbox"/>	17b. Date of Procedure	17c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
18a. Other biopsy-not colposcopy	<input type="checkbox"/>	18b. Date of Procedure	18c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
19a. Other Cervical Procedures:	<input type="checkbox"/>	19b. Date of Procedure	19c. Bill to PRBCCEDP:	<input type="radio"/> Yes <input type="radio"/> No	
C. Diagnosis Information					
20a. Status of Final Diagnosis:					
<input type="radio"/> Work-up Complete <input type="radio"/> Lost to Follow Up <input type="radio"/> Work-up Refused					
21. Final Diagnosis:					
<input type="radio"/> Normal / Benign Reaction / Inflammation <input type="radio"/> HPV / Condylomata / Atypia <input type="radio"/> CIN I / Mild Dysplasia (Biopsy Diagnosis) <input type="radio"/> CIN II / Moderate Dysplasia (Biopsy Diagnosis) <input type="radio"/> CIN III / Severe Dysplasia / Carcinoma in situ (Stage 0) (Biopsy Diagnosis)		<input type="radio"/> Invasive Cervical Carcinoma (Biopsy Diagnosis) <input type="radio"/> HSIL <input type="radio"/> LSIL <input type="radio"/> Adenocarcinoma <input type="radio"/> Other Final Diagnosis (Specify): <div style="border: 1px solid black; height: 100px; width: 100%;"></div>			
D. Treatment Information					
22a. Status of Treatment:		22b. Date of Treatment:		23a. Follow-up:	
<input type="radio"/> Treatment Started <input type="radio"/> Treatment Pending <input type="radio"/> Treatment not Needed		<input type="radio"/> Treatment Refused <input type="radio"/> Lost to Follow-up (includes death)		<input type="radio"/> 5 years <input type="radio"/> 3 year <input type="radio"/> 1 year <input type="radio"/> Short-Term 23b. Specify Short-Term months: <input type="text"/>	
24. Comments: <input type="text"/>					
25. Provider's Name and Signature:					26. Date:

Formas de Referido


 Universidad de Puerto Rico
 Centro Comprensivo de Cáncer
 Programa para la Prevención y Detección Temprana del
 Cáncer de Mama y Cáncer de Cuello Uterino Puerto Rico


Forma de Referido
Procedimientos de Cernimiento/Diagnóstico de Cáncer de Cuello Uterino

Paciente: _____ Lugar de Referido: _____

<u>Procedimientos de Cernimiento</u>	<u>Procedimientos de Diagnóstico</u>
Citología Convencional de Pap <input type="checkbox"/>	Colposcopia sin Biopsia <input type="checkbox"/>
Thin Prep. (LBC) de Pap <input type="checkbox"/>	Colposcopia con Biopsia <input type="checkbox"/>
Sure Path (LBC) de Pap <input type="checkbox"/>	Colposcopia con ECC <input type="checkbox"/>
Otro Procedimiento: <input type="checkbox"/>	ECC solo <input type="checkbox"/>
_____	LEEP <input type="checkbox"/>
	CKC <input type="checkbox"/>
	Otro Procedimiento: <input type="checkbox"/>

Referida por: _____ Lic: _____ Fecha de Referido: _____
 Procedimiento realizado por: _____ Fecha de Procedimiento: _____
 Interpretado por: _____ Lic: _____ Fecha de Interpretación: _____

Last Revised: 12/2013 Original (Facturación al Programa) – 1ra Copia (Records Laboratorio) – 2da Copia (Records Médico) Form RDC101-S


 Universidad de Puerto Rico
 Centro Comprensivo de Cáncer
 Programa para la Prevención y Detección Temprana del
 Cáncer de Mama y Cáncer de Cuello Uterino de Puerto Rico

Forma de Referido
Procedimientos para Cernimiento/Diagnóstico de Cáncer de Mama

Paciente: _____ Lugar de Referido: _____

<u>Procedimientos de Cernimiento</u>	<u>Procedimientos de Diagnóstico</u>
Izquierdo Derecho Ambos	Izquierdo Derecho Ambos
Mamografía <input type="radio"/> <input type="radio"/> <input type="radio"/>	Diagnostic Mammography <input type="radio"/> <input type="radio"/> <input type="radio"/>
Sonmamografía <input type="radio"/> <input type="radio"/> <input type="radio"/>	Fine Needle Aspiration Biopsy <input type="radio"/> <input type="radio"/> <input type="radio"/>
Otro Procedimiento: <input type="radio"/> <input type="radio"/> <input type="radio"/>	Large Core Needle Biopsy <input type="radio"/> <input type="radio"/> <input type="radio"/>
_____	Open Surgical Biopsy <input type="radio"/> <input type="radio"/> <input type="radio"/>
	Otro Procedimiento: <input type="radio"/> <input type="radio"/> <input type="radio"/>

Referida por: _____ Lic: _____ Fecha de Referido: _____
 Procedimiento realizado por: _____ Fecha de Procedimiento: _____
 Interpretado por: _____ Lic: _____ Fecha de Interpretación: _____

Last Revised: 12/2013 Original (Facturación al Programa) – 1ra Copia (Records Laboratorio) – 2da Copia (Records Médico) Form RDB101-S

Laboratorio

- Indicar en boleta o requisición que el método de pago será el programa.
 - Escribir UPRCCC en la sección de método de pago (“Medical Insurance”)
- También, acompañar la muestra con copia del compromiso de pago.

Tarifas

- Tarifas a base de Medicare Physician Fee Schedule (Non-Facility Fee) y Clinical Laboratory Fee Schedule.
- Envío del documento “NBCCEDP Allowable Procedures and Relevant CPT® Codes” a todos los proveedores anualmente.
- **No se permite el cobro de deducible o diferencia a las participantes del programa.**

Facturación

- Documentos para facturación al programa:
 - Factura (1500, UB-04 u otro método)
 - Nombre de la participante, fecha de nacimiento, información del proveedor (nombre, direcciones física/postal y NPI) fecha del servicio, códigos CPT y tarifas.
 - Formas clínicas del programa debidamente completadas (Médicos)
 - Resultados (Laboratorios y Centros de Imágen)
 - Copias de compromisos de pago asociadas a los servicios ofrecidos
- Enviar los documentos durante los primeros 7 días laborables de cada mes.

<div style="border: 1px solid black; display: inline-block; padding: 2px 5px;">1500</div> HEALTH INSURANCE CLAIM FORM <small>APPROVED BY NATIONAL UNIFORM CLAIM COMMITTEE (NUCC)</small>												PATIENT AND INSURED INFORMATION
<input type="checkbox"/> INDICA <input type="checkbox"/> FICA												
1. MEDICARE: MEDICATED # (Medicare #)		2. PATIENT'S NAME (Last Name, First Name, Middle Initial)		3. PATIENT'S BIRTH DATE (MM DD YY)		4. INSURED'S NAME (Last Name, First Name, Middle Initial)		5. INSURED'S ID NUMBER (For Program in Section 5)		6. INSURED'S ADDRESS (No. Street)		PATIENT AND INSURED INFORMATION
7. PATIENT'S ADDRESS (No. Street)		8. PATIENT RELATIONSHIP TO INSURED (Self, Spouse, Child, Other)		9. PATIENT STATUS (Single, Married, Other)		10. IS PATIENT'S CONDITION RELATED TO (Employment, Auto Accident, Other Accident)		11. INSURED'S POLICY GROUP OR FICA NUMBER		12. INSURED'S DATE OF BIRTH (MM DD YY)		
13. OTHER INSURED'S NAME (Last Name, First Name, Middle Initial)		14. OTHER INSURED'S POLICY OR GROUP NUMBER		15. OTHER INSURED'S DATE OF BIRTH (MM DD YY)		16. EMPLOYER'S NAME OR SCHOOL NAME		17. INSURANCE PLAN NAME OR PROGRAM NAME		18. IS THERE AN ADDITIONAL BENEFIT PLAN (YES/NO)		PATIENT AND INSURED INFORMATION
19. OTHER INSURED'S POLICY OR GROUP NUMBER		20. OTHER INSURED'S DATE OF BIRTH (MM DD YY)		21. EMPLOYER'S NAME OR SCHOOL NAME		22. INSURANCE PLAN NAME OR PROGRAM NAME		23. IS THERE AN ADDITIONAL BENEFIT PLAN (YES/NO)		24. IS THERE AN ADDITIONAL BENEFIT PLAN (YES/NO)		
READ BACK OF FORM BEFORE COMPLETING & SIGNING THIS FORM. 25. PATIENT'S OR AUTHORIZED PERSON'S SIGNATURE (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												PATIENT AND INSURED INFORMATION
26. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
27. DATE OF SERVICE (MM DD YY)		28. DATE OF INJURY (MM DD YY)		29. DATE OF SERVICE (MM DD YY)		30. DATE OF INJURY (MM DD YY)		31. DATE OF SERVICE (MM DD YY)		32. DATE OF INJURY (MM DD YY)		PATIENT AND INSURED INFORMATION
33. NAME OF REFERRING PROVIDER OR OTHER SOURCE		34. NAME OF REFERRING PROVIDER OR OTHER SOURCE		35. NAME OF REFERRING PROVIDER OR OTHER SOURCE		36. NAME OF REFERRING PROVIDER OR OTHER SOURCE		37. NAME OF REFERRING PROVIDER OR OTHER SOURCE		38. NAME OF REFERRING PROVIDER OR OTHER SOURCE		
39. RESERVATION FOR LOCAL USE												PATIENT AND INSURED INFORMATION
40. RESERVATION FOR LOCAL USE												
41. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												PATIENT AND INSURED INFORMATION
42. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
43. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												PATIENT AND INSURED INFORMATION
44. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
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48. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
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60. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
61. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												PATIENT AND INSURED INFORMATION
62. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												
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67. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in total or to the party who accepts assignment shown.)												PATIENT AND INSURED INFORMATION
68. SIGNATURE OF PHYSICIAN OR SUPPLIER (Indicate the release of any medical or other information necessary to process the claim. I also request payment of government benefits either in												

Envío de Documentos

- Se enviarán al UPRCCC
 - Taína De La Torre (Data Manager)
 - Email: tdelatorre@cccupr.org
 - documentos deben estar protegidos con el password que provee el programa
 - Correo regular: Universidad de Puerto Rico
Centro Comprensivo de Cáncer
PRBCCEDP
PO BOX 363027
San Juan PR, 00936-3027
 - Personalmente/mensajero: Universidad de Puerto Rico,
Centro Comprensivo de Cáncer
Piso 2, Complejo del Centro Médico
San Juan, PR 00935
 - **No se aceptan documentos vía fax.**

Aprobación y Pago

- Búsqueda en las bases de datos para asegurar:
 - la paciente es participante del programa
 - los servicios no hayan sido facturados anteriormente
 - los servicios fueron los aprobados por el programa
- Verificar que las formas clínicas y las facturas esten debidamente completadas.
 - Los documentos que tienen información incompleta o errores se devuelven al proveedor para correcciones antes de aprobar el pago.
- UPRCCC paga los servicios de los proveedores que tienen contrato directo.
- SAC paga los servicios de los proveedores que **no** tienen contrato con el UPRCCC.



Bienvenidos a Nuestro Portal

El Programa de Prevención y Detección Temprana de Cáncer de Mama y Cuello Uterino de PR es auspiciado por los Centros de Control y Prevención de Enfermedades (CDC por sus siglas en inglés). El Programa provee servicios de cerimiento y diagnóstico para cáncer de seno (mama) y cérvix (cuello uterino) de manera gratuita a mujeres que sean elegibles, así como servicio de navegación de pacientes y manejo de casos, y actividades educativas para la comunidad y los profesionales de la salud.



Cáncer de Seno

El cáncer de seno en Puerto Rico continúa siendo el cáncer más común y la primera causa de muerte por cáncer en las mujeres. La mamografía es la única prueba recomendada para la detección del cáncer de seno (mama) en sus etapas tempranas.

[Ver Más](#)



Cáncer de Cérvix

El cáncer de cérvix (cuello uterino) continúa siendo de las causas más importantes de cáncer en la mujer en Puerto Rico. Este cáncer es prevenible a través de la vacunación contra el virus de papiloma humano y la prueba de Papanicolaou.

[Ver Más](#)

<http://www.cancerdesenoycuellouterino.com> o
<http://www.cancerdesenoycuellouterino.org>

Página Electrónica

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Navigation menu:

- INICIO
- SOBRE NOSOTROS
- CÁNCER DE SENO
- CÁNCER DE CÉRVIX
- PARTICIPANTES
- PROVEEDORES**
- RECURSOS
- EVENTOS Y NOTICIAS

Main image: A close-up of a doctor's hands holding a pen, with a stethoscope visible on the left.

Proveedores

- Funcionamiento del programa ▾
- Política de pago por servicios de cernimiento y diagnóstico de cáncer de mama y cuello uterino ▾
- Formas clínicas para procedimientos de cernimiento y diagnóstico de cáncer de mama (seno) ▾
- Formas clínicas para procedimientos de cernimiento y diagnóstico de cáncer de cérvix (cuello uterino) ▾
- Formas de referido para procedimientos de cernimiento y diagnóstico ▾
- Ask DR. Miller ▾

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ABOUT

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<http://www.controldecancerpr.org/>

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Control de Cáncer PR
 August 13 ·

¡Conoce más sobre nosotros!
 Control de Cáncer de Puerto Rico reúne varios programas relacionados a la prevención, detección, control y educación relacionada al Cáncer; tales como el Programa de Prevención y Detección Temprana de Cáncer de Mama y Cuello Uterino, el Programa de Control Comprensivo, y el Registro Central de Cáncer, en un solo lugar.
 Conoce más sobre nosotros en <http://www.controldecancerpr.org/>



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Material Educativo



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E-mail: tdelatorre@cccupr.org



National Breast
and Cervical Cancer
Early Detection
Program

