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Exploring Cancer

Examining the Role of Biology, Race, Class, and Socioeconomics

August 25 Welcome to Cancer(s) and Health Disparities 101 - The Introduction

September 1 Radiation Oncology - What Is It, and What Is It Good For?

September 8 New Strategies in Treating GI Cancers October 13 Breast Cancer Health Disparities

October 20 Pancreatic Cancer

October 27 Careers in Cancer

Fridays

11:00 - 11:50 AM EST/EDT

November 10 Precision Medicine and Immunotherapy

November 17

Expanding Cancer Care Quality and Delivery in Sub-Saharan Africa: a collaborative approach





Checo Rorie, PhD

Checo J. Rorie, PhD, is originally from Marshville, NC, was raised by his grandparents and is a first-generation college student. Dr. Rorie attended Clark Atlanta University in Atlanta, GA, where he majored in Biology and graduated in 1998 with a bachelor of science degree. As an undergraduate, Dr. Rorie was a MARC Scholar and conducted breast cancer research in a Cancer Cell Biology laboratory. Dr. Rorie then attended the University of North Carolina at Chapel Hill's Curriculum in Toxicology graduate program earning a PhD in 2004. After graduating from UNC-CH, Dr. Rorie completed a postdoctoral fellowship at New York University, and then participated in a second postdoctoral fellowship back at UNC-CH in the Seeding Postdoctoral Innovators in Research & Education (SPIRE) program.

Dr. Rorie has been at North Carolina Agricultural and Technical State University since 2008, and is currently the Professor and Chair of the Department of Biology. Dr. Rorie has a Cancer Genetics and Cell Biology laboratory where his lab studies the mechanisms of breast cancer health disparities in African American Women.

Professional Highlights

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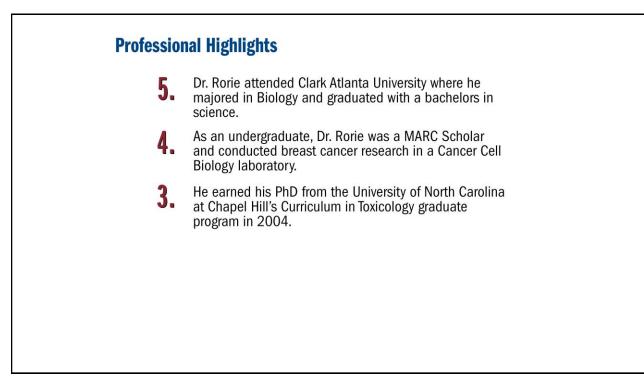
Professional Highlights

5. Dr. Rorie attended Clark Atlanta University where he majored in Biology and graduated with a bachelors in science.

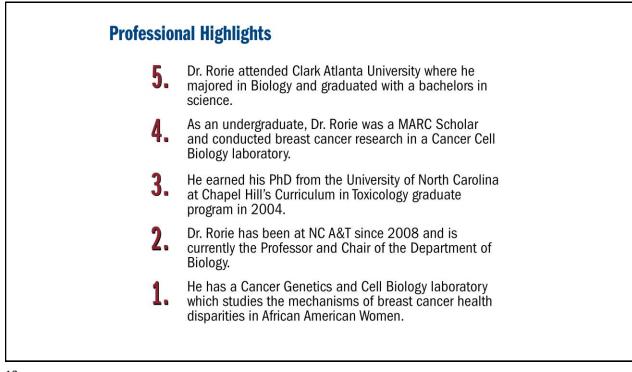
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Breast Cancer Health Disparities

Checo J. Rorie, PhD Professor & Chair Department of Biology



- Originally from Marshville, North Carolina; Forest Hills High School 1994
- Attended Clark Atlanta University, Atlanta, Georgia;
 B.S. Biology 1998 (John Browne)
- Attended UNC Chapel Hill; Ph.D. Toxicology 2004 (Bernard "Buddy" Weissman)
- Postdoc at New York University, New York, New York; Biochemistry 2005 (James "Jim" Boroweic)
- Postdoc in the SPIRE Program at UNC Chapel Hill; Radiation Oncology 2008 (YanPing Zhang)
- Currently: Professor and Chair of Biology, NC A&T State University

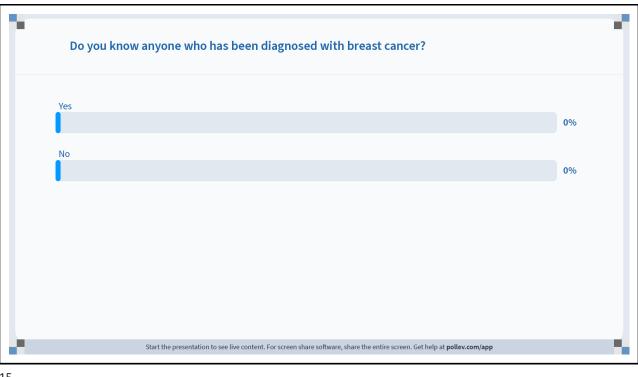
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My Journey



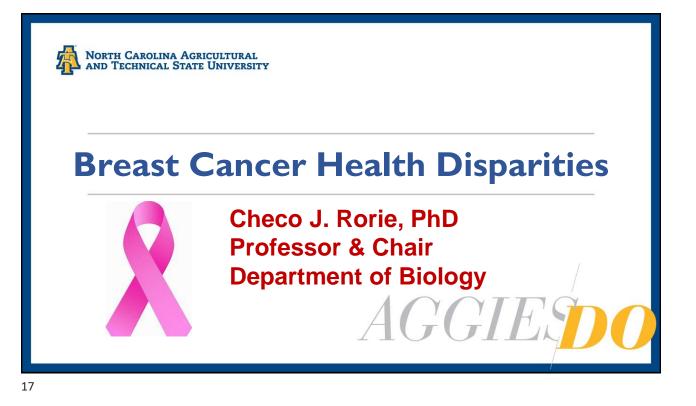
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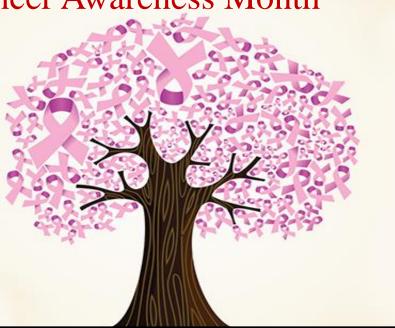






Breast Cancer Awareness Month

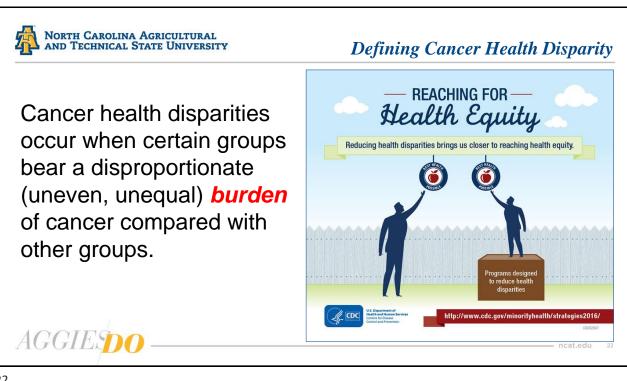
In honor of Aunt Francis and our friend Amanda







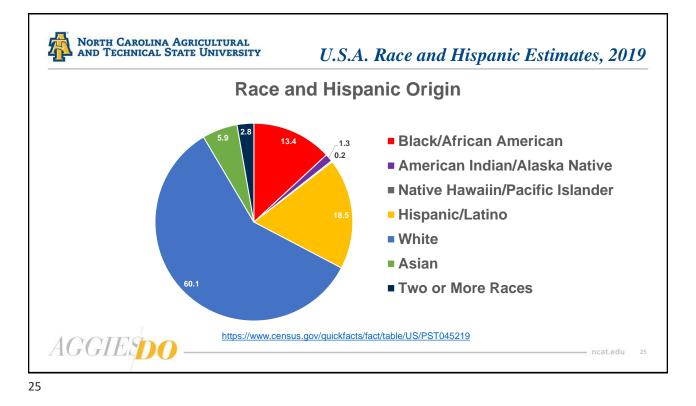


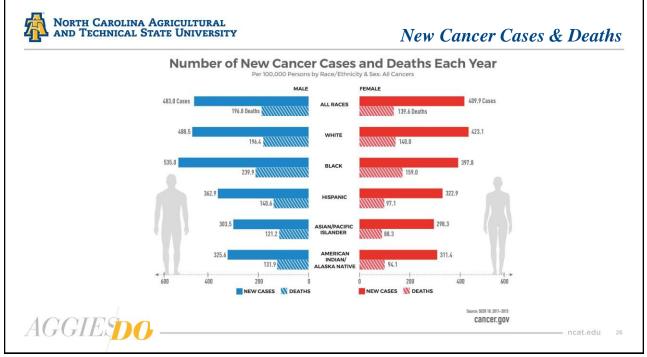


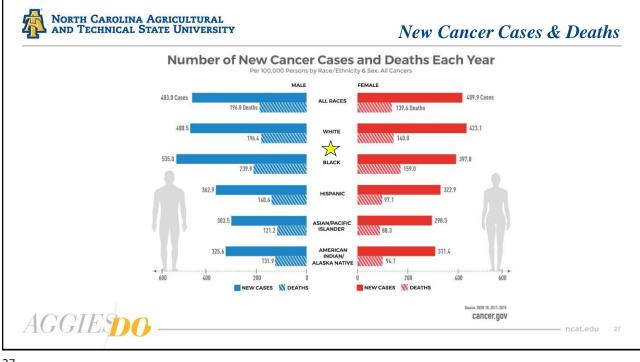
	Male				Female				
	Prostate	161,360	19%			Breast	252,710	30%	
50	Lung & bronchus	116,990	14%	- X -		Lung & bronchus	105,510	12%	
se	Colon & rectum	71,420	9%			Colon & rectum	64,010	8%	
ü	Urinary bladder	60,490	7%			Uterine corpus	61,380	7%	
ŝ	Melanoma of the skin	52,170	6%			Thyroid	42,470	5%	
z	Kidney & renal pelvis	40,610	5%			Melanoma of the skin	34,940	4%	
Ę	Non-Hodgkin lymphoma	40,080	5%			Non-Hodgkin lymphoma	32,160	4%	
na	Leukemia	36,290	4%			Leukemia	25,840	3%	
Estimated New Cases	Oral cavity & pharynx	35,720	4%			Pancreas	25,700	3%	
ш	Liver & intrahepatic bile duct	29,200	3%			Kidney & renal pelvis	23,380	3%	
	All sites	836,150	100%			All sites	852,630	100%	
	Male					Female			
	Lung & bronchus	84,590	27%		_	Lung & bronchus	71,280	25%	
	Colon & rectum	27,150	9%			Breast	40,610	14%	
Ś	Prostate	26,730	8%		- 57 -	Colon & rectum	23,110	8%	
Estimated Deaths	Pancreas	22,300	7%			Pancreas	20,790	7%	
) Se	Liver & intrahepatic bile duct	19,610	6%			Ovary	14,080	5%	
뉟	Leukemia	14,300	4%			Uterine corpus	10,920	4%	
ate	Esophagus	12,720	4%			Leukemia	10,200	4%	
<u>, e</u>	Urinary bladder	12,240	4%			Liver & intrahepatic bile duct	9,310	3%	
ŝ	Non-Hodgkin lymphoma	11,450	4%			Non-Hodgkin lymphoma	8,690	3%	
	Brain & other nervous system	9,620	3%			Brain & other nervous system	7,080	3%	
-		318,420	100%			All sites	282,500	100%	



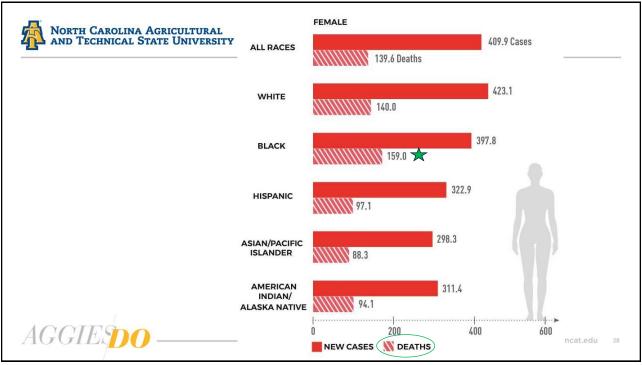
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imat			404		Non-Hodgkin lymphoma	8,690	3%
Estimated Deaths	Non-Hodgkin lymphoma	11,450	4%		Non-Hodgkin tymphoma	0,000	0.0
Estimat	-	11,450 9,620	4% 3%		Brain & other nervous system	7,080	3%

















What is Breast Cancer Health Disparity?

Breast cancer health disparities result when there are *differences* in the *expectations* of cancer measurements and outcomes

- Incidence or new cases diagnosed
- Prevalence or existing cases in a population
- Mortality or death related to cancer
- Survivorship or quality of life after cancer treatment
- Screening rates

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Stage at diagnosis

https://www.cancer.gov/about-cancer/understanding/disparities

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Breast Cancer



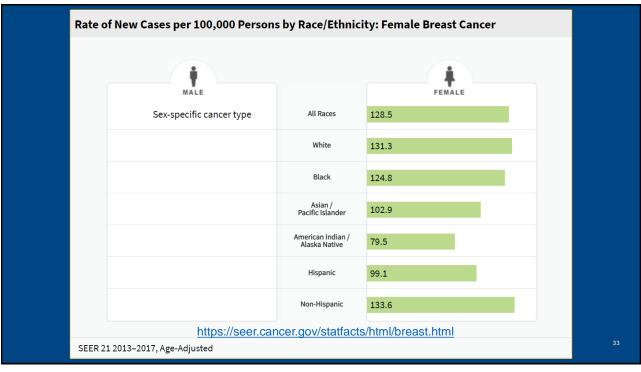


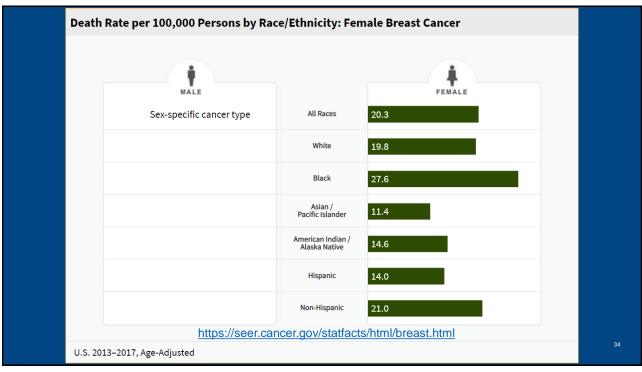
- Most breast cancers begin in the lobules (milk glands) or in the ducts that connect the lobules to the nipple.
- Typically has no symptoms when the tumor is small and most easily treated, which is why screening is important for early detection.
- Most common physical sign is a painless lump.
- Men get breast cancer too (less than 1%)

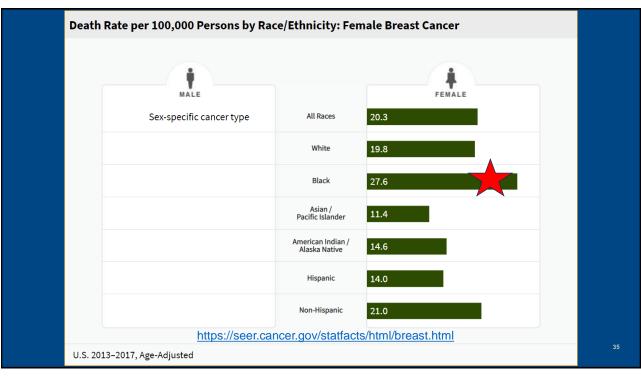
Duct	Lobule
Invasive lobular cancer (carcinoma) Ductal carcinoma in situ (DCIS) (non-invasive	
invasive ductal cancer (carcinoma)	Z

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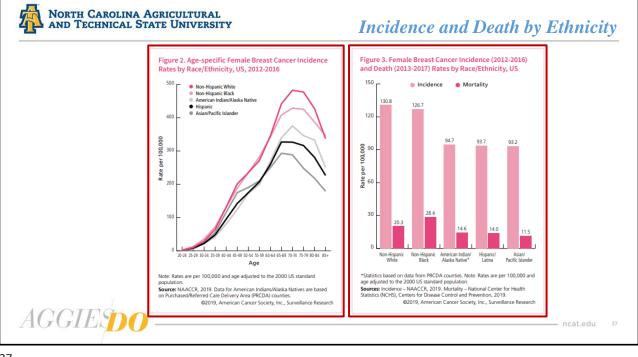
exspecific cancer type All Races 128.5 All Races 20.3 All Races 20	ases per 100,000 Person	is by Race/Ethnie	city: Female Breast Cancer	Death Ra	ate per 100,000 Persons by Ra	ce/Ethnicity: Fe	male Breast Cancer
exspecific cancer type All Races 128.5 All Races 20.3 All Races 20	+		-		-		-
Black 124.8 Pacific blander 102.9	MALE Sex-specific cancer type	All Races				All Races	FEMALE
Adian / Adian / Pacific Islander 102.9 Pacific Islander 11.4		White	131.3			White	19.8
Pacific Islander 11.4 Pacific Islander 11.4		Black	124.8			Black	27.6
Amarican Indian /		Asian / Pacific Islander	102.9			Asian / Pacific Islander	11.4
Alaska Native 79.5 American Indian / Alaska Native 14.6		American Indian / Alaska Native	79.5			American Indian / Alaska Native	14.6
Hispanic 99,1 Hispanic 14,0		Hispanic	99.1			Hispanic	14.0
Non-Hispanic 133.6 Non-Hispanic 21.0		Non-Hispanic	133.6			Non-Hispanic	21.0
	17, Age-Adjusted			U.S. 2013-	2017, Age-Adjusted		







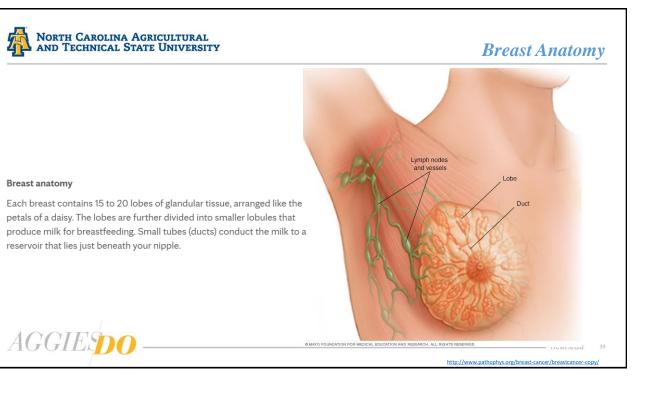
Cases per 100,000 Persor	ns by Race/Ethnie	city: Female Breast Cancer	Death	n Rate per 100,000 Persons by Ra	ce/Ethnicity: Fe	male Breast Cancer
+						+
MALE Sex-specific cancer type	All Races	FEMALE 128.5		MALE Sex-specific cancer type	All Races	FEMALE
	White	131.3			White	19.8
	Black	124.8			Black	27.6
	Asian / Pacific Islander	102.9			Asian / Pacific Islander	11.4
	American Indian / Alaska Native	79.5			American Indian / Alaska Native	14.6
	Hispanic	99.1			Hispanic	14.0
	Non-Hispanic	133.6			Non-Hispanic	21.0
17, Age-Adjusted			U.S. 20	013–2017, Age-Adjusted		



North Carolina Agricultural and Technical State University

Breast Cancer Histology & Subtypes

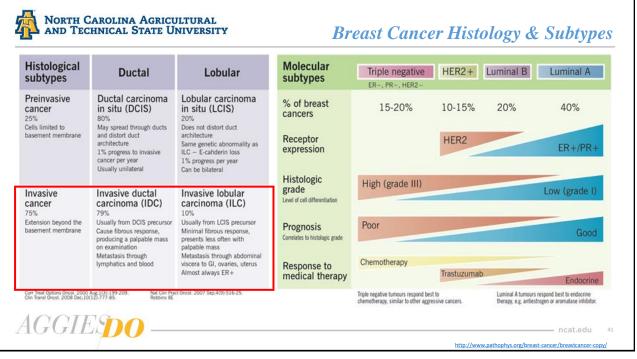
Ductal	Lobular	Molecular	Triple negative	HER2+	Luminal B	Luminal A
		subtypes	ER-, PR-, HER2-			
Ductal carcinoma in situ (DCIS)	Lobular carcinoma in situ (LCIS) 20%	% of breast cancers	15-20%	10-15%	20%	40%
and distort duct architecture 1% progress to invasive cancer per year	architecture Same genetic abnormality as ILC – E-cahderin loss 1% progress per year	Receptor expression		HER2		ER+/PR+
Osually uninateral	Can be bilateral	Histologic	Lich (made III)			
Invasive ductal carcinoma (IDC) 79%	Invasive lobular carcinoma (ILC)	grade Level of cell differentiation	High (grade III)			Low (grade I)
Usually from DCIS precursor Cause fibrous response, producing a palpable mass on examination	Usually from LCIS precursor Minimal fibrous response, presents less often with palpable mass	Prognosis Correlates to histologic grade	Poor			Good
Metastasis through	Metastasis through abdominal	Deserves to	Chemotherapy			
symphatics and blobd	Almost always ER+	Response to medical therapy		Trastuzumal	b	Endocrine
Aug. 1(3):199-209. Nat Clin Prac 12):777-85. Robbins 8E	t Oncol. 2007 Sep;4(9):516-25.		Triple negative tumours respond best chemotherapy, similar to other aggre	l to ssive cancers.		respond best to endocrine trogen or aromatase inhibitor.
	Ductal carcinoma in situ (DCIS) 80% May spread through ducts and distort duct architecture 1% progress to invasive cancer per year Usually unilateral Invasive ductal carcinoma (IDC) 79% Usually from DCIS precursor Cause fibrous response, producing a palpable mass on examination Metastasis through lymphatics and blood	Ductal carcinoma in situ (DCIS) Lobular carcinoma in situ (LCIS) 80% Bostoria 20% May spread through ducts architecture Does not distort duct architecture 20% 1% progress to invasive cancer per year Does not distort duct architecture 30% Ilsually unilateral ILC – E-cahderin loss 1% progress per year Can be bilateral ILC – E-cahderin loss 1% progress per year Can be bilateral Invasive ductal carcinoma (IDC) 10% Usually from DCIS precursor Gause fibrous response, producing a palpable mass Metastasis through lymphatics and blood Invasive lobular carcinoma (ILC) 10% Usually from LCIS precursor Maimal fibrous response, presents less often with palpable mass Metastasis through lymphatics and blood Metastasis through abdominal viscera to Gl, ovaries, uterus Almost always ER+	Ductal Lobular subtypes Ductal carcinoma in situ (DCIS) 80% Lobular carcinoma in situ (LCIS) 20% % of breast cancers May spread through duct and distord duct architecture Lobular carcinoma in situ (LCIS) 20% % of breast cancers Display the second state of the second architecture Des not distort duct architecture % of breast cancers 1% progress to invasive cancer per year Usually unilateral Des not distort duct architecture Same genetic abnormality as ILC – E-cahderin loss Receptor expression Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (ILC) 10% Histologic grade Usually from DCIS precursor Cause fibrorus response, presents less often with palpable mass on examination Metastasis through lymphatics and blood Prognosis Correlates to histelogic grade Metastasis through lymphatics and blood Almost always ER+ Response to medical therapy	Ductal Lobular subtypes Imple negative Ductal carcinoma in situ (DCIS) 80% Lobular carcinoma in situ (LCIS) 20% % of breast cancers 15-20% May spread through ducts and distort duct architecture cancer per year Usually unilateral Lobular carcinoma in situ (LCIS) 20% % of breast cancers 15-20% Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (IDC) 79% Invasive lobular carcinoma (IDC) 10% Invasive lobular carcinoma (IDC) 10% Invasive lobular carcinoma (IDC) 10% Histologic grade Level d cell differentiation High (grade III) Usually from DCIS precursor Cause fibrous response, producing a plaptie mass Metastasis through abdominal twiscera to GI, ovaries, uterus Almost always ER+ Prognosis Comelates to histeligic grade Response to medical therapy Poor Statistics and blood Signer once, 200% Metastasis through abdominal twiscera to GI, ovaries, uterus Response to medical therapy Chemotherapy	Ductal Lobular subtypes Imple negative HER2 + Ductal carcinoma in situ (DCIS) 80% Lobular carcinoma in situ (LCIS) 20% No f breast cancers 15-20% 10-15% May spread through ducts and distort duct architecture cancer per year Usually unilateral Does not distort duct architecture cancer per year Can be bilateral % of breast cancers 15-20% 10-15% Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (ILC) 10% Invasive lobular carcinoma (ILC) 10% Histologic grade grade High (grade III) Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (ILC) 10% Prognosis Comelates to histologic grade Poor Usually from DCIS precursor Cause fibrous response, producing a palpable mass metastasis through lymphates and blood Noaries, uterus Almost always ER + Prognosis Comelates to histologic grade Poor Suite 129/2920 BetCht 207 Step.409:616-25. Trastuzuma Chemotherapy Trastuzuma	Ductal Lobular subtypes Imple negative HER2 + Luminal B Ductal carcinoma in situ (DCIS) 80% Lobular carcinoma in situ (LCIS) 20% No f breast cancers 15-20% 10-15% 20% May spread through ducts and distort duct architecture cancer per year Usually unilateral Same genetic abnormality as LC = E-andrein loss % of breast cancers 15-20% 10-15% 20% Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (ILC) 10% Invasive lobular carcinoma (ILC) 10% Histologic grade High (grade III) Invasive ductal carcinoma (IDC) 79% Invasive lobular carcinoma (ILC) 10% Prognosis Constats to histologic grade Poor Invasive ductal carcinoma (IDC) 79% Invasive lobular Usually from DCIS precursor Minimal fibrous response, presents less often with palpable mass Amost always ER + Prognosis Correlates to histologic grade Poor Question fibrous response, producing a palpable mass Amost always ER + Response to medical therapy Chemotherapy gright 192-290 Metch 2920 fbreat. 2007 Sec.409:616-25. Trastuzumab



North Carolina Agricultural and Technical State University

Breast Cancer Histology & Subtypes

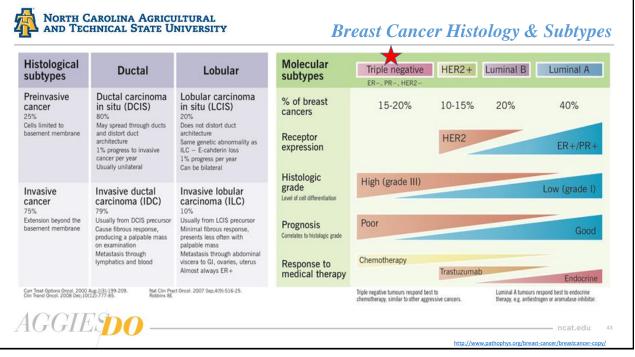
Histological subtypes	Ductal	Lobular	Molecular subtypes	Triple negative ER-, PR-, HER2-	HER2+	Luminal B	Luminal A
Preinvasive cancer 25% Cells limited to basement membrane	Ductal carcinoma in situ (DCIS) 80% May spread through ducts and distort duct architecture 1% progress to invasive cancer per year	Lobular carcinoma in situ (LCIS) 20% Does not distort duct architecture Same genetic abnormality as ILC – E-cahderin loss 11% progress per year	% of breast cancers Receptor expression	15-20%	10-15% HER2	20%	40% ER+/PR+
Invasive cancer	Usually unilateral Invasive ductal carcinoma (IDC)	Can be bilateral Invasive lobular carcinoma (ILC)	Histologic grade Level of cell differentiation	High (grade III)			Low (grade I)
75% Extension beyond the basement membrane	79% Usually from DCIS precursor Cause fibrous response, producing a palpable mass on examination	10% Usually from LCIS precursor Minimal fibrous response, presents less often with palpable mass	Prognosis Correlates to histologic grade	Poor			Good
	Metastasis through lymphatics and blood	Metastasis through abdominal viscera to GI, ovaries, uterus	Response to	Chemotherapy			
		Almost always ER+	medical therapy		Trastuzuma	b	Endocrine
Curr Treat Options Oncol. 2000 Clin Transl Oncol. 2008 Dec;10	Aug:1(3):199-209. Nat Clin Prac (12):777-85. Robbins 8E	t Oncol. 2007 Sep;4(9):516-25.		Triple negative tumours respond best chemotherapy, similar to other aggres			respond best to endocrine trogen or aromatase inhibitor.
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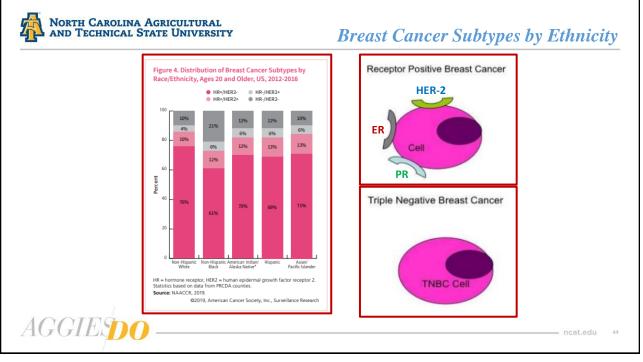


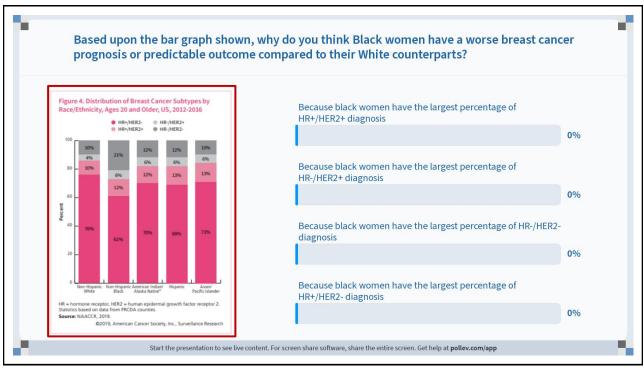
North Carolina Agricultural and Technical State University

Breast Cancer Histology & Subtypes

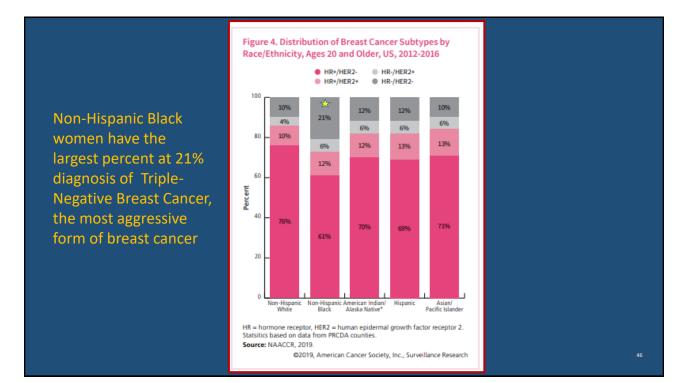
112-1-1-1-1-1			Malandar	1	2	3	4
Histological	Ductal	Lobular	Molecular	Triple negative	HER2+	Luminal B	Luminal A
subtypes			subtypes	ER-, PR-, HER2-			
Preinvasive cancer 25%	Ductal carcinoma in situ (DCIS) 80%	Lobular carcinoma in situ (LCIS) 20%	% of breast cancers	15-20%	10-15%	20%	40%
Cells limited to basement membrane	May spread through ducts and distort duct architecture 1% progress to invasive cancer per year Usually unilateral	Does not distort duct architecture Same genetic abnormality as ILC – E-cahderin loss 1% progress per year Can be bilateral	Receptor expression		HER2		ER+/PR+
			Histologic	High (grade III)			
Invasive cancer 75%	Invasive ductal carcinoma (IDC)	Invasive lobular carcinoma (ILC)	grade Level of cell differentiation	ringin (grade in)			Low (grade I)
Extension beyond the basement membrane	Usually from DCIS precursor Cause fibrous response, producing a palpable mass on examination	Usually from LCIS precursor Minimal fibrous response, presents less often with palpable mass	Prognosis Correlates to histologic grade	Poor	-		Good
	Metastasis through lymphatics and blood	Metastasis through abdominal viscera to GI, ovaries, uterus		Chemotherapy			
	Tymphanes and mood	Almost always ER+	Response to medical therapy		Trastuzuma	b	Endocrine
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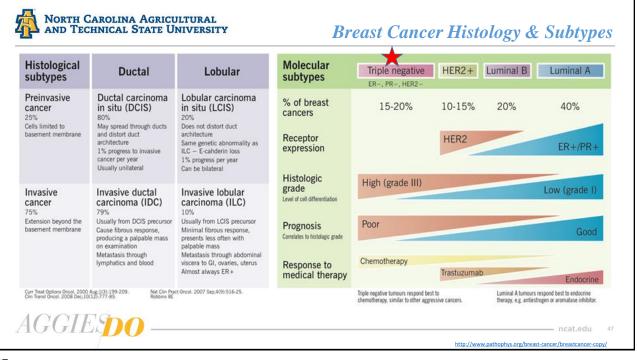










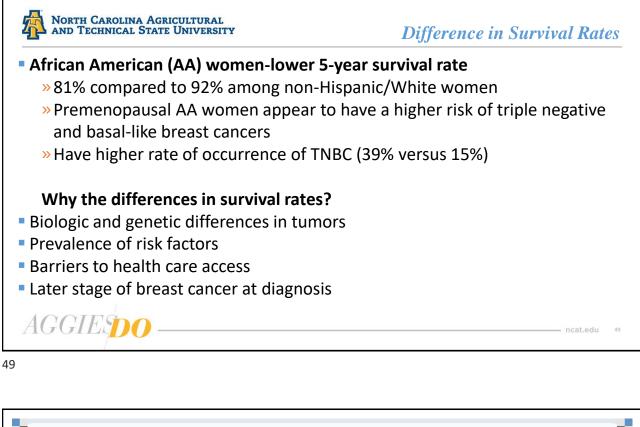




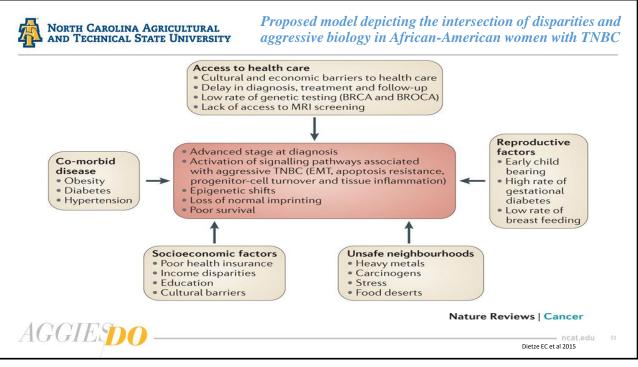
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Background: Triple negative breast cancer

- Lack of estrogen receptors (ER), progesterone receptors (PR) and human epidermal growth factor receptor-2 (HER2) expression resulting in *lack of* targeted therapies
- Typically stains positive for *mutant-p53* (80% of cases)
- Account for 10-17% of all breast cancer
- More prevalent in African-American women with a higher death rate in those age <35</p>
- Significantly more aggressive (high growth rates, highly invasive/metastatic) than other subgroups with a peak *risk of recurrence (survival rate 40%)* in visceral and soft tissue between the first- and third-years following therapy





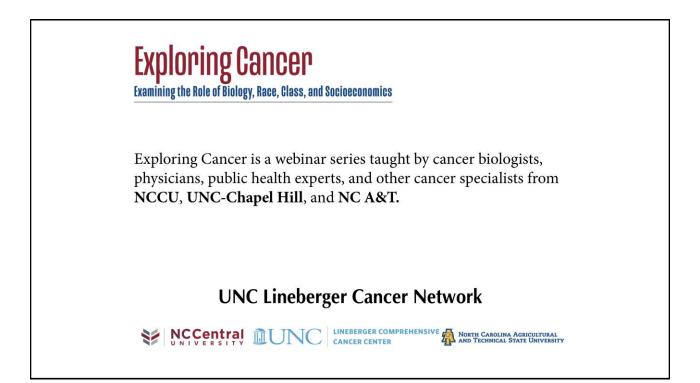




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	Thank You!	

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Exploring Cancer Examining the Role of Biology, Race, Class, and Socioeconomics

Thank you for participating!

You may now return to the UNCLCN Learning Portal to complete a course evaluation and claim your certificate.

For any technical issues or questions, contact us at Email: unclcn@unc.edu Call: (919) 445–1000