GENITOURINARY SYSTEM

SUBJECT : Pathology GUS

LEC NO. : 16

DONE BY : محمد احمد محمد العجو

وَقُل رَّبِّ زِدْنِي عِلْمَاً

تفريغ الـ

وَقُل رَّبِّ زِدْنِي عِلْمَاً

عزيزي القارئ ، في هذا التفريغ ، انا بشرح و بلخص جزء من المحاضرة بعديد بعرض محتوى السلايدات ، لكن تلخيصي للمادة شامل جدا و بتقدر تعتمد عليه في دراسة المادة

اذكروني بدعوة في ظهر الغيب

وَقُل رَّبِّ زِدْنِي عِلْمًا

بسم الله الرحمن الرحيم ،، اول اشي بدك تعرف انه سرطان الثدي يصنف ثاني اكثر سرطان بسبب الوفاة عند النساء بعد سرطان الرئة و بتم تقسيم سرطان الثدي الى نوعين ، الاول و هو نادر و مش مهم و ما بنركز عليه ، يلي هو sarcoma ، يلي بطلع من ال stromal component of the Brest ، يعني من ال blood , fibroblast vessels , و من الأمثلة عليه phyllodes tumors and angiosarcoma

اما النوع الاهم هو ال carcinoma يلي بطلع من ال epithelial component of the breast ، يلي هم الخلايا يلي بتغطي ال lobules and terminal ducts (يلي هي المكونات المسؤولة عن تكوين الحليب)

بعد هاي الفقرة في معومات كلها عبارة عن ارقام ، زي الكومينتي فما رح احكي عنها

هسة بدنا نيجي لتقسيم السرطانات (التقسيم الاول كان بناء على الخلايا يلي طلع منها السرطان)
هسة بدنا نيجي للتقسيم المهم و المستخدم اكثر و هو انه نصنف السرطانات حسب ال expression of hormone receptors

بتتذكروا بالفارما بس اخذنا دوا نسيت شو اسمه بس يلي بستهدف السرطانات يلي عليها مستقبلات استروجين ، اه اسمه tamoxifen ،، بس هسة بدنا نصنف السرطانات حسب الreceptors تاعتها

ممكن نلاقي : estrogen receptors , progesterone receptors , human epidermal growth factor receptor 2

و بناء على ما ذكرت بنقسم سرطانات الثدي الى ٣ مجموعات:

ER positive (HER2 negative; 50%–65% of cancers)
HER2 positive (ER positive or negative; 10%–20% of cancers)
Triple negative (ER, PR, and HER2 negative; 10%–20% of cancers) اخطر نوع

معلومة انذكرت بالسلايدات :

HER2 is a receptor tyrosine kinase that promotes the proliferation and suppress apoptosis

معلومة حكتها الدكتورة :
اي فحص لسرطان الثدي لازم نشوف هذول ال ٤ :
ER,PR,HER2,ki67

و حديثا صاروا يشوفوا p53

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GENITOURINARY SYSTEM

تمام نيجي لتقسيم ثالث ،، التقسيم يلي بدي احكي عنه هسة بنستخدمه بالابحاث ، و فيه بنقسم السرطانات الى ٤ انواع

1. Luminal A :

Majority of cases are lower grade
ER positive and HER negative
Has low levels of protein Ki-67
Grow more slowly
Good prognosis

2. Luminal B :

Higher grade
ER positive HER 2 negative cancers
Has high levels of protein Ki-67(progesterone negative)
Faster growth of cancer cells

3. HER2-enriched :

Overexpress HER2,ER and PR negative.
Grow faster than luminal cancers يعني أسوأً من يلي فوق
Worse prognosis
Usually successfully treated with targeted therapy medicine aimed at the HER2

4. Triple negative or Basal-like:

ER,PR,HER2 negative
Gene expression profiling resemble nasally located myoepithelial cells
More common in:
1. a BRCA1 mutation
2. Younger women
3. Black women

وقل رب زدني علماً

هسة رح نحكي عن ال Risk factors بدنا نحكي عن العوامل يلي بتتحكم بحدوث سرطان الثدي

العمر : بين ٣٠-٦٠ ، و بصير عند النساء اكثر لكن اذا صار عند الرجال بكون قاتل ، و برضه بصير عند النساء البيض اكثر و اذا صار عند السود بكون قاتل ، برضه ال family history ، اذا وحدة بعائلتها في حالات سرطان ثدي فرصة حدوث السرطان عندها عالية جدا
او اذا وحدة عندها سرطان بالثدي سابقا او عاملة استئصال لكتلة برضه الفرصة لحدوث سرطان عندها بزيد ، او اذا وحدة عاملة radiation على منطقة الصدر او الوجه قبل سن الثلاثين برضه بتزيد الفرصة ، او اذا وحدة عندها تغييرات بالصدر زي ECD او duct papilloma ، و برضه اذا في obesity او lack of exercise ، او اذا في بلوغ مبكر (menarche) او يلي ما العوانس المش متزوجات (nulliparity) او انقطاع متأخر للدورة الشهرية هذول لانهم بتعرضوا لنسبة عالية من الاستروجين و لفترة اطول ، كل هاي عوامل بتأثر على حدوث المرض ، و برضه الجينات بتلعب دور ، مثلا يلي عندها طفرة HER2/new ، ممكن يصير عندها سرطان ثدي

برضه ال hormonal replacement therapy بتزيد الفرصة ، بتصير عند النساء او الرجال يلي يتعالجوا بال oestrogen ، زي يلي عندهم سرطان بروستات و يوخذوا استروجين ، او يلي يتحولوا جنسيا 🏳️‍🌈 ، او النساء يلي بتاخذ استروجين .

و برضه مهم اسأل عن ال history of pregnancy and breast feeding ، لانه الولادة و الرضاعة بعملوا وقاية من سرطان الثدي فاذا في وحدة تأخرت بالزواج فرصة انه يصير عندها سرطان ثدي اعلى

الكحول بتزيد فرصة حدوث سرطان اما بالنسبة للتدخين تأثيره مش كثير على سرطان الثدي لكن بأثر على علاج السرطان ، فمثلا لو وحدة عندها سرطان ثدي و بدها تعمل radiation ، ما بتستعمل هذا الاشعاع لانه الرئة عندها تالفة

معلومة كمان ، انه بصير عند الامريكان اكثر من الاسيويين

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هسة يا حلو بدنا ندخل بال pathogenesis ، بالصفحة يلي قبل حكينا عن عوامل الخطر و حكينا من ضمنهم ان الجينات ، يعني انت صرت ماخذ فكرة عن الكلام يلي رح احكيه هسة ،، في عنا ٣ عوامل بتساهم بشكل مباشر بال pathogenesis لسرطان الثدي و هم:

1. Genetic factors

في عندي جينات يلي هم BRCA2, BRCA1 هذول عبارة عن Tumor suppressor genes و سرطان الثدي بصير بسسس اذا صار mutation بهذول الجينات (كلاهما) ، طبعا هاي الجينات يا حلو وظيفتها انه تعمل repair of DNA damage

اعرف انه BRCA2 mutations هو ER + ، اما BRCA1 mutations هو triple negative و بالتالي يعتبر اخطر

شو في كمان مشاكل جينية بتعملي سرطان ثدي؟ TP53 and PTEN

2. Hormonal factors

الاستروجين يعتبر من العوامل المهمة يلي بتسبب سرطان بالثدي لانه بعمل تحفيز لانتاج growth factors promoting the tumor development
و برضه كمان اذا كان عندي precursor الاستروجين بخليه يصير fully malignant

و برضه مضادات الاستروجين زي tamoxifen ، بتقلل حدوث سرطان الثدي بس يكون ER+

3. Environmental factors

Morphology

Location:

Upper outer quadrant (50%)   Central portion(20%).
 Lower outer quadrant 10%
 Upper inner quadrant 10%
 Lower inner quadrant 10%

4% have bilateral primary tumors or sequential lesions in the same breast.

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هسة نط لصفحة رقم ١٢

السلايدات:

Breast Cancer

There are many types of breast cancers, and correctly identifying each one is important to determine the proper treatment.

Breast cancers can be divided into two main overarching groups: the carcinomas and the sarcomas.

Carcinomas are cancers that arise from the epithelial component of the breast. The epithelial component consists of the cells that line the lobules and terminal ducts; under normal conditions, these epithelial cells are responsible for making milk. Carcinomas comprise the vast majority of all breast cancers, and will be further discussed below.

Sarcomas are rare cancers that arise from the stromal (connective tissue) components of the breast. These stromal component cells include myofibroblasts and blood vessel cells, and cancers arising from these "supportive" cells include phyllodes tumors and angiosarcoma.

من هون و انزل المادة
بتصير community

Sarcomas account for less than 1% of primary breast cancers.

In the US, invasive Ca B is 2nd to lung cancer as a cause of cancer death in women, & despite advances in diagnosis & treatment, 1/4 of women who develop Ca B will die of it.

The lifetime risk of Ca B is one in eight (1/8) for women in the US, with 75% of cases older than age 50.

about 5-6% are younger than the age of 40.

About 9% of all new cases of breast cancer in the United States are found in women younger than 45 years of age.

For unknown reasons (possibly related in some part to earlier detection via mammography) there has been worldwide increase in the incidence of Ca B.

Is The most common non-skin malignancy of women.

2nd most common cause of cancer deaths in women, following carcinoma of the lung.

The worldwide incidence and mortality are increasing at an alarming rate. This trend is due to social changes especially in the developing countries.

Those social changes Include delayed childbearing, fewer pregnancies, and reduced breastfeeding and with lack of access to optimal health care.

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Since 1980s the mortality rate has dropped from 30%
to <20% due to improvement in detecting cancers
before they metastasize through screening (mammographic screening) and more effective systemic treatment.
Almost all breast malignancies are adenocarcinomas (>95%).

Classification system

The most clinically used classification system for breast cancer depends on the expression of hormone receptors
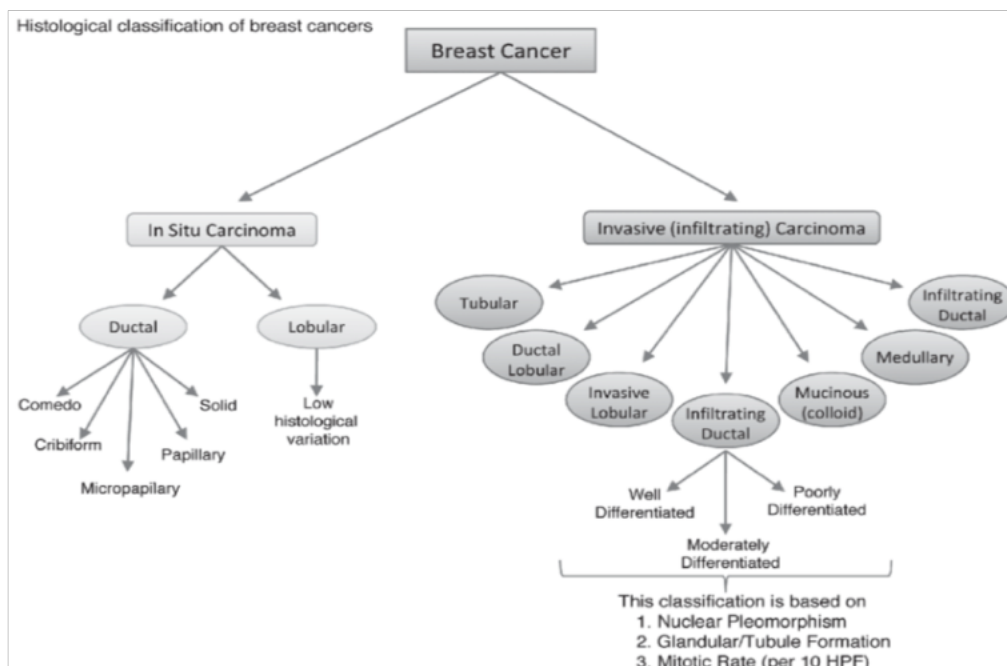
hormone receptors are:

Estrogen receptor (ER), progesterone receptor (PR) & human epidermal growth factor receptor 2 (HER2, or ERBB2)

Can be classified according to expression of hormone receptors into three major groups:

ER positive (HER2 negative; 50%–65% of cancers)
HER2 positive (ER positive or negative; 10%–20% of cancers)
Triple negative (ER, PR, and HER2 negative; 10%–20% of cancers)

Histological classification of breast cancers

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                          Breast Cancer

         In Situ Carcinoma              Invasive (infiltrating) Carcinoma

       Ductal      Lobular         Tubular                    Infiltrating
                                                                Ductal
                                  Ductal
  Comedo      Solid    Low        Lobular              Medullary
              histological
  Cribiform   variation      Invasive            Mucinous
         Papillary           Lobular  Infiltrating  (colloid)
                                      Ductal
   Micropapilary
                                  Well              Poorly
                              Differentiated      Differentiated

                                     Moderately
                                    Differentiated

                          This classification is based on
                          1. Nuclear Pleomorphism
                          2. Glandular/Tubule Formation
                          3. Mitotic Rate (per 10 HPF)
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The three groups show striking differences in patient characteristics, pathologic features, treatment response, metastatic patterns, time to relapse, and outcome

Within each group are additional histologic subtypes, some of which also have clinical importance.

An alternative classification system relies on gene expression profiling

used mainly in clinical research
divides breast cancers into four major types:

1.Luminal A. majority of cases are lower grade, ER-positive & HER2 negative cancers, has low levels of the protein Ki-67, which helps control how fast cancer cells grow. Luminal A cancers tend grow more slowly than other cancers, be lower grade, and have a good Px

2.Luminal B.Majority of cases are higher grade ER positive +ve
and HER2 negative cancers .Has High Ki67 protein which indicate faster growth of cancer cells Or progesterone Negative

3. HER2-enriched. overexpress HER2 and ER-negative.ER and PR Negative , Her2 Positive.HER2-enriched cancers tend to grow faster than luminal cancers and can have a worse prognosis, but are usually successfully treated with targeted therapy medicines aimed at the HER2

4.Triple Negative or Basal-like.
Triple-negative or basal-like breast cancer is estrogen receptor-negative, progesterone receptor-negative, and HER2-negative. Gene expression profiling resemble basally located myoepithelial cells

Triple-negative breast cancer is more common in people with:
1-a BRCA1 mutation
2-younger women
3-Black women

Triple-negative breast cancer is considered more aggressive than either luminal A or luminal B

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Risk Factors

Age:
It is considered rare in women younger than 25 and incidence increase after the age of 30. more than two thirds of women with breast cancer are older than the age of 50 and only 5% are younger than the age of 40.

Gender:
The incidence in men is only 1% of that in women.

Family History of Breast Cancer:
The greatest risk is for individuals with multiple affected first-degree relatives with early onset breast cancer mostly related to various combinations of low penetrance or weak cancer genes.
However, in about 5 to 10% of cases a highly penetrance germline mutations in the tumor suppressor genes is associated with lifetime risk greater than 90%

Geographic Factors:
higher in the Americas and Europe than in Asia and Africa
The mortality rates of breast cancer in America is 5 times greater than japan .
Immigration studies showed that immigration from low incidence to high incidence areas tends to acquire the rates of their new home countries.
In this context, diet, reproductive patterns, and breast feeding practices are thought to be involved .
Breast cancer rates appear to be raising in parts of the world
that are adapting the western habits.

Race/Ethnicity:
highest rate in women of European descent because of higher incidence of ER-positive cancers.
Hispanic and African American develop cancer at a younger age and develop aggressive tumors.
This is thought to result from combination of differences in genetic social factors and access to health care.

Reproductive History:
Including Early age of menarche, nulliparity, absence of breastfeeding, with older age at first pregnancy are all associated with increased risk due to increased the exposure of the epithelial cells of the breast to estrogenic stimulation.

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Ionizing Radiation:
Chest Radiation especially if the breast is developing.

Other Risk Factors:
Postmenopausal obesity
postmenopausal hormone replacement therapy
mammographic density
alcohol consumption

Pathogenesis
Factors that contribute directly to the development of breast cancer can be grouped into:

1. Genetic
2. Hormonal
3. Environmental

Genetic Factors

BRCA1 and BRCA2: Are classic tumor suppressor genes and the cancer only occur if both alleles are defected encode proteins that are required for repair of DNA damage. most carriers develop breast cancer by the age of 70 years

For unclear reasons, BRCA2 mutations are primarily associated with ER-positive tumors, whereas BRCA1 mutations are associated with triple-negative cancers

Other mutated genes: TP53 and PTEN

P53 (A gene that makes a protein that is found inside the nucleus of cells and plays a key role in controlling cell division and cell death. Mutations (changes) in the p53 gene may cause cancer cells to grow and spread in the body. It is guardian of the genome

The pathways in which familial breast cancer genes function also are often disturbed in sporadic cancers

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HER2 gene amplification :
Cancers that overexpress HER2 are highly proliferative.
In the past they had a poor prognosis; Nowadays, the availability of therapeutic agents targeting HER2 has improved the prognosis.
It is a receptor tyrosine kinase that promotes the cell proliferation and suppress apoptosis

Hormonal factors

Estrogens are considered an important hormonal factors since they stimulate the production of growth factors promoting the tumor development.

Estrogen receptors regulate other genes in an estrogen dependent fashion. Some of those genes are important for the tumor development or growth.

Estrogens also drives the proliferation from precursor regions to a fully malignant and metastatic carcinoma.

Estrogen antagonists: reduce the development of ER-positive cancers in women at high risk and are mainstays in the treatment of established ER-positive tumors.

Morphology

Location:
Upper outer quadrant (50%) Central portion(20%).
 Lower outer quadrant 10%
 Upper inner quadrant 10%
 Lower inner quadrant 10%

4% have bilateral primary tumors or sequential lesions in the same breast.

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حكينا انها epithelial component

هسة بدنا نحكي عن تقسيمات ال Brest carcinoma

Non invasive

Invasive

Intradutal
(ductal carcinoma in situ)

Lobular
(Lobularl carcinoma in situ)

Bilateral
Multicentic
good prognosis

Ductal ⟿ 80 %
Lobular ⟿ 10 %
tubular ⟿ 6%
Medullary ⟿ 2%
Mucinous ⟿ 2%

tumor cells proliferate incide duct lumen

Small

large

- Uniform cells
- Central small nuclei
- loosely Cohesive
- indistinct cell boundaries

Four patterns:

solid type

Comedotype ⟿ like acne

papillary type

Cribriform type

تمام يا حلوين ، بهاي الصفحة بدي بس تاخذوا نظرة عامة و بدي تعرفوا انه ال epithelial component of the breast بتكون عبارة عن ducts اوlobules ، و بكلامها ممكن يصير كانسر و ممكن يكون invasive او لا

أهم جاجة انهم مش عاملين

Invasion

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بسم الله الرحمن الرحيم ، نبلش تلخيص باقي المحاضرة ، اول اشي يا حلوين احنا ما رح ندرس كل الانواع يلي كتبتهم فوق و اصلا الدكتورة ما حكت عنهم كلهم و انا جايبهم من الدكتور سامح غازي ،، رح نناقش تقريبا ٥ انواع ،،

و عشان اريحكم في ٣ انواع بشبهوا بعض ببعض الصفات لهيك رح احطهم جمب بعض و ندرسهم مع بعض و هيك بنكون خلصنا ٣ من ٥ ، كيفني معكم ؟

Invasive ductal carcinoma (NOS)	Invasive lobular carcinoma	Tubular carcinoma
Most common	Associated with lobular carcinoma in situ	Irregular mamographic density
Associated with ductal carcinoma in situ	Almost all these carcinoma express hormone receptors but HER2 over expression is very rare or absent	Well formed tubules with low grad nuclei
Produce desmoplastic response lead to mamographic density		Lymph node metastases are very rare
immunohistochemical breast cancer prognostic and therapeutic markers used include: estrogen receptor, human epidermal growth factor receptor-2, Ki-67, progesterone receptor, and p53.	Loss of E-Catherine is a specific bio marker	Excellent prognosis
	Metastases reach CSF, serosal surface ,bone marrow ,ovary, uterus	HER2 negative
Hard palpable , irregular mass	Discohesive	Sometimes mistaken for benign sclerosing lesions. Calcification may present in the tumor lumen
Cohesive		

- arranged in groups
لأ بس رح تشوف شكلين من هاي المجموعات بنفس الريت

① indian file pattern
Malignant cells of linear arrangement

② Target like pattern
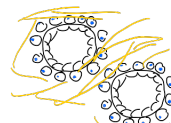
في معلومة حكتها الدكتورة انه بس يكون ال mypepithelial in perephery of duct بكون ductal carcinoma in situ او lobular carcinoma in situ

وقل رب زدني علماً

هسة رح احكي عن كمان ٢ ، و هيك بنكون قربنا نخلص ، لولولوليييش

Medullary carcinoma

Triple negative , and don't over express HER2/new

large anaplastic cells with pushing, well-circumscribed borders with a pronounced lymphocytic infiltrate.

No precancerous lesion

Increase frequency in women with BRCA1

Rounded mass

Colloid mutinous carcinoma

Rare

Produce extra cellular mucin that dissects into the gelatinous

ER positive ,HER2 negative Abundant blue mucin

هسة مشان نكون خلصنا كل الامراض ، ضل اشي واحد نحكي عنه

Inflammatory Carcinoma

clinically present as an enlarged, swollen, erythematous breast (resulting from the blockage of dermal lymphatic spaces by ca cells)usually without or, with ill-define palpable mass or presents with breast erythema and skin thickening

صلى الله و بارك ، اقفز على صفحة ٢١ ، و بدي اياك و انت بتقفز توخذ نظرة سريعة خمس دقائق فقط ، و ركز على الشغلات يلي مظللها (تظلييييش مش تخطيط ، لاني مخطط كثير شغلات ، المهم المظلل عشان تكون لام المادة ، يلا نطواا عشان حكي على اخر موضوع

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Breast carcinoma

Noninvasive:

(confined by a basement membrane and do not invade into stroma or lympho-vascular channels), include:

1. Ductal carcinoma in situ
2. Lobular carcinoma in situ

Invasive (infiltrating):

1. Invasive ductal carcinoma (includes all carcinomas that are of no special type)70% to 80%
2. Invasive lobular carcinoma 10% to 15%
3. Carcinoma with medullary features5%
4. Mucinous carcinoma (colloid carcinoma) 5% 5. Tubular carcinoma5%
6. Other types

1. Invasive ductal carcinoma (NST)(NO SPECIAL TYPE)   وه   أشش هاهـ

70-80%

Arise from Milk Duct , This type of cancer forms in the lining of a milk duct within your breast. The ducts carry breast milk from the lobules, where it's made, to the nipple.

Ductal carcinoma can remain within the ducts as a noninvasive cancer (ductal carcinoma in situ), or it can break out of the ducts (invasive ductal carcinoma).

Previously called Carcinomas "not otherwise specified"((NOS))

Precancerous lesion: usually DCIS

Cases with invasive ductal carcinoma produces desmoplastic response which replaces the normal fat and result in mammographic densities

The most common immunohistochemical breast cancer prognostic and therapeutic markers used include: estrogen receptor, human epidermal growth factor receptor-2, Ki-67, progesterone receptor, and p53.
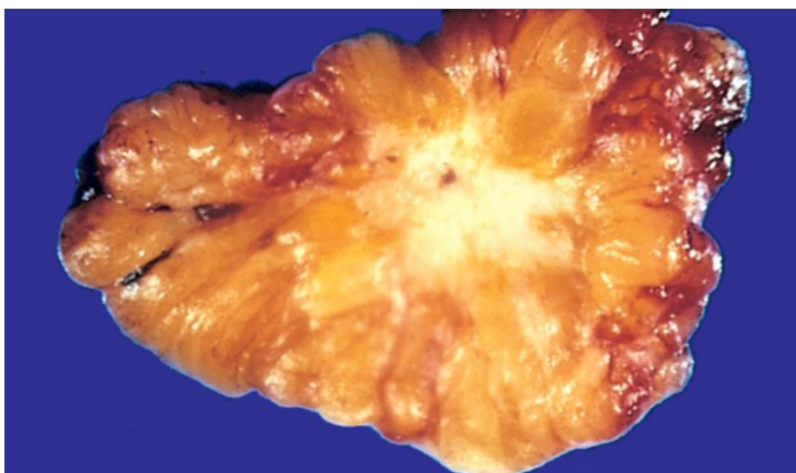
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Clinical presentation:

a mammographic density; a hard, palpable irregular mass,
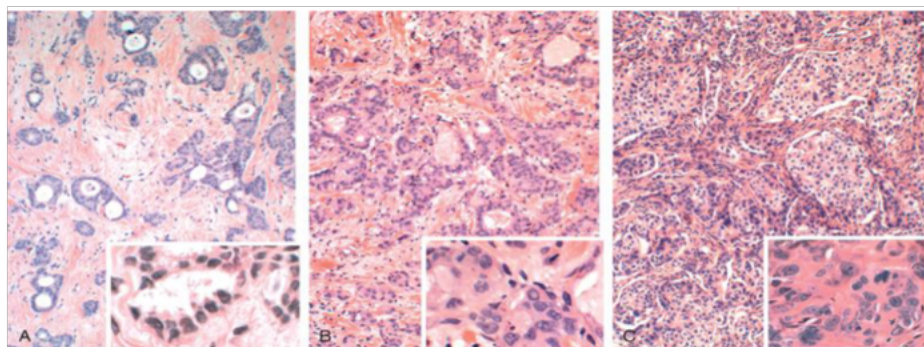nipple retraction, or fixation to the chest wall can be seen in advanced cancers

Receptor profile:
ER (+ve in 50-60%) HER2 (+ve in 20%)
15% are negative for both

Breast carcinoma, not well
circumscribed ,Irregular ,C/S
showing gritty sensation ,
hard ,white because of
desmoplasia

Invasive ductal carcinoma

A-Well differentiated carcinoma consisting of tubules with small monomorphic
nuclei
B-Moderate differentiation carcinoma with less tubular formation more solidness of
cells and monomorphic nuclei
C-Poorly differentiated carcinoma with sheets of pleomorphic cells containing
numerous mitotic figures and central areas of tumor necrosis
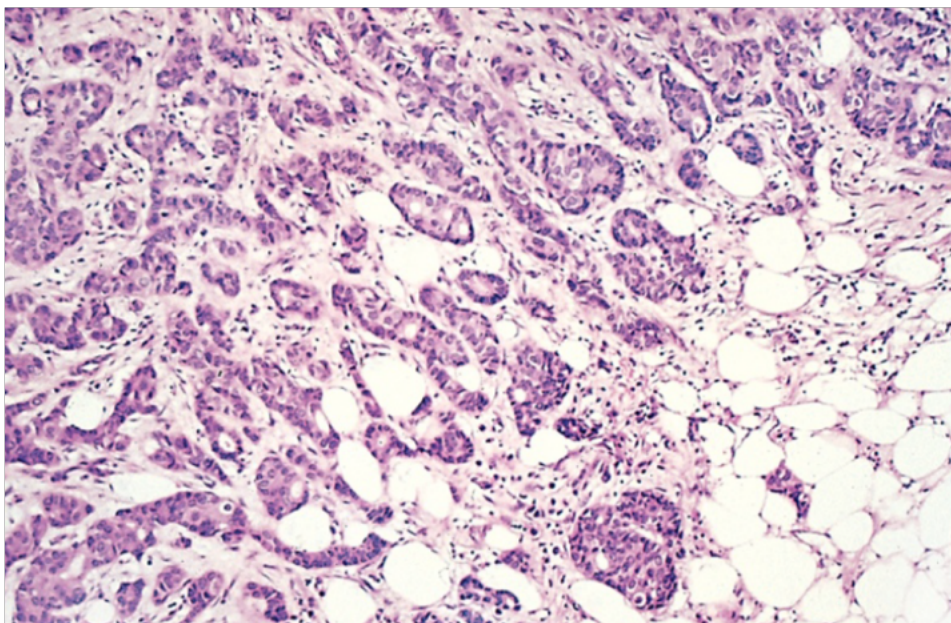
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Breast carcinoma

margin, showing invasion & infiltration of the adjacent fatty tissue (on the right).

Invasive lobular carcinoma

10-15%

Arise from Milk producing lobule, Milk-producing lobules.
Lobular carcinoma starts in the lobules of the breast, where breast milk is produced. When it breaks out of the lobules, it's considered invasive lobular carcinoma. The lobules are connected to the ducts, which carry breast milk to the nipple.

Precancerous lesion: associated with LCIS.

10% to 20% are multicentric and bilateral

Clinical presentation. Most present as palpable masses or mammographic densities

Invasive lobular carcinoma is generally composed of single (CD) small cells arrayed in a linear pattern with a targetoid pattern invading into stroma, TDLU, and adipose tissue of the breast. Neoplastic cells display round nuclei often eccentrically placed with occasional intracytoplasmic vacuoles.

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cells invade stroma individually and often are aligned in "single-file"

Almost all of these carcinomas express hormone receptors, but HER2 overexpression is very rare or absent.

Markers. Loss of E-cadherin is a specific biomarker for invasive lobular carcinoma as opposed to invasive breast carcinoma of no special type which show E.cadherin Positive
Metastasis of lobular carcinoma is unique since it frequently reaches the CSF, serosal surfaces, bone marrow , ovary, and uterus

Carcinoma with Medullary features

5%

Triple negative
Microscopically: large anaplastic cells with pushing, well-circumscribed borders with a pronounced lymphocytic infiltrate.

Precancerous lesions. usually absent

increased frequency in women with BRCA1 mutations,.

Receptor profile. lack hormone receptors and do not overexpress HER2/NEU.

Those carcinoma typically grow as rounded masses that can be difficult to distinguish from benign tumors on imaging

Histology with Carcinoma with medullary features
the tumor in the middle consist of tightly adhesive
clusters of cells
At he periphery there is a dense lymphocytic infiltrate around the island of tumor cells
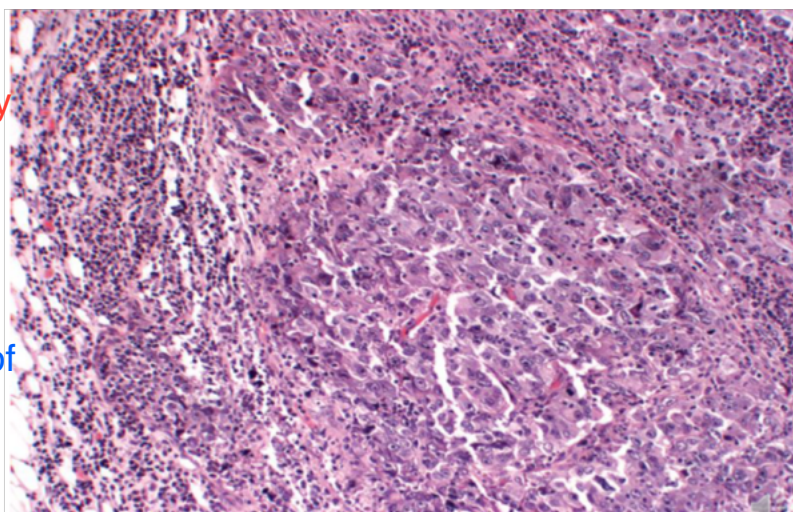
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Colloid mucinous Carcinoma
a rare subtype

Microscopic picture:
The tumor cells produce abundant quantities of extracellular mucin that dissects into the gelatinous.

ER-positive/HER2-negative cancer

Shows abundant blue mucin and the carcinoma cells appears to be floating in those lakes of mucin
This mucin matrix gives the tumor the grossly soft blue to gray appearance
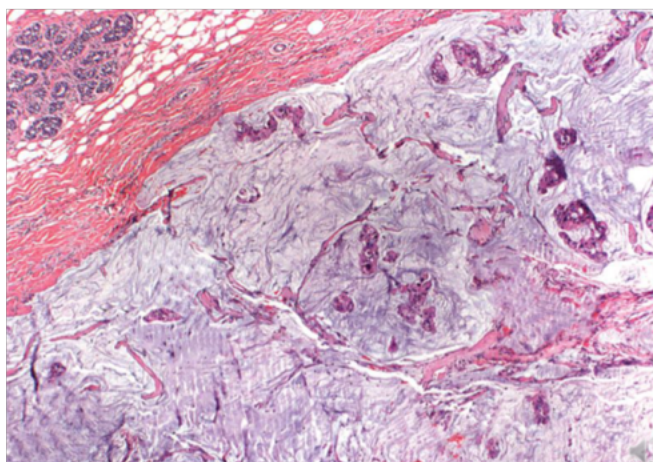
Tubular carcinoma

10% of invasive carcinomas

Clinical presentation. irregular mammographic densities.
Microscopically, well-formed tubules with low-grade nuclei.
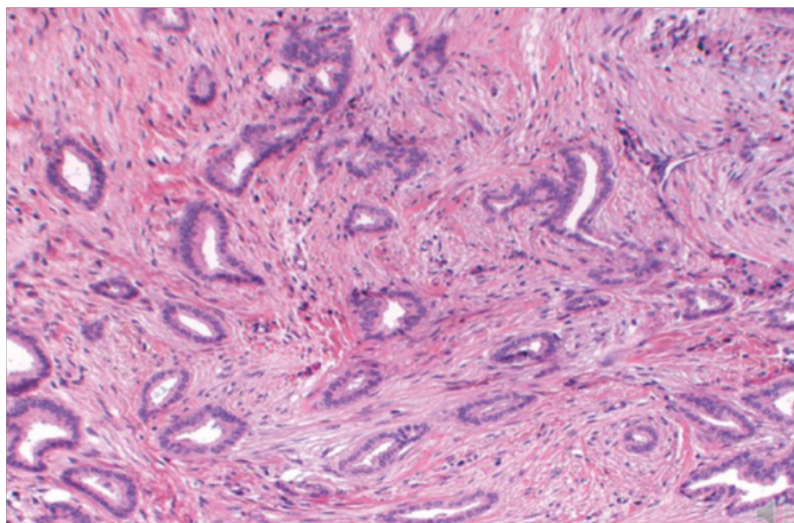Lymph node metastases are rare, and prognosis is excellent.
ER-positive/HER2-negative cancer

Sometimes mistaken for benign sclerosing lesions. Calcification may present in the tumor lumen

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well- differentiated neoplastic cells form a single cuboidal layer in small,round to tear drop shaped ductules widely spaced in a fibrous stroma.

Inflammatory Carcinoma

clinically present as an enlarged, swollen, erythematous breast (resulting from the blockage of dermal lymphatic spaces by ca cells)usually without or, with ill-define palpable mass or presents with breast erythema and skin thickening

The ca is generally poorly differentiated & diffusely invading the breast tissue.
True inflammation is minimal or absent.
Most of these T have distant metastases & the prognosis is extremely poor.
mimics the surface of an orange peel, an appearance referred to as peaud'orange.

Lactating (Inflammatory) carcinoma:

breast. A large greyish-white cancer with extensive central necrosis & hemorrhage.
Clinically, the tumor resembles an acute inflammatory lesion & has a rapid malignant course with extremely poor prognosis.

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Spread of breast cancer

through lymphatic and hematogenous channels.
Favored metastasis are the bone, lungs, skeleton, liver, and adrenals and (less commonly) the brain, spleen, and pituitary.

Metastases may appear many years after apparent therapeutic control of the primary lesion that's why we use screening program

SCREENING:
mammographic screening,Magnetic resonance imaging, MRI

Spread eventually occurs through lymphatic & hematogenous channels.

LN metastases are present in about 50% of ca presenting as palpable masses, but... in fewer than 15% of cases found by mammography.

Outer quadrants & centrally located catypically spread first to the axillary LN.
Ca B in the inner quadrantsoften involve the LN along the internal mammary arteries.

The supraclavicular LN are usually become involved only after the axillary & internal mammary LN are affected, but... sometimes are the primary site of spread (Skipped).

More distant dissemination eventually follows, with metastatic involvement of almost any organ or tissue in the body. Favored locations are the lungs, skeleton, liver, & adrenals & (less commonly) the brain, spleen, & pituitary. However, no site is exempt!
Metastases may appear many years (sometimes 15 years) after apparent therapeutic control of the primary ca

Clinically,Ca B is often discovered by the woman or her physician as a solitary, painless, & fixed mass. At this time, the ca is typically 2 to 3 cmin with involvement of the regional LNs(most often axillary) in about 50% of patients.

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Breast cancer Prognosis

The outcome for women with breast cancer depends on the biologic features of the carcinoma (molecular or histologic type)and the extent to which the cancer has spread (stage) at the time of diagnosis.

Prognostic Factors

Tumor stage:

Invasive carcinoma versus carcinoma in situ.
Distant metastases.
Lymph node metastases.
Tumor size.In cm
Locally advanced disease
Inflammatory carcinoma
Lymphovascularinvasion
Molecular subtype.
Special histologic types.
Histologic grade
Estrogen and progesterone receptors and HER2 expression

مهم

Distance metastases:
Once distant metastases are present, cure is unlikely.

Lymph node metastases:
Axillary lymph node status is the most important prognostic factor for invasive carcinoma in the absence of distant metastases,biopsy is necessary for accurate assessment.

With no lymph involvement the ten years survival is 70-80%
1 -3 lymph involvement 35-40%
If more than 10 lymph nodes 10-15%

Tumor size:
 The risk of axillary lymph node metastases increases with the size of the primary tumor, but both are independent prognostic factors.

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Locally advanced disease:
Carcinomas invading into skin or skeletal muscle are usually large and may be difficult to treat surgically

Lymphovascular Invasion:
strongly associated with the presence of lymph node metastases.
poor prognostic factor

Special Histologic Types:
The survival rate of women with tubular, mucinous, lobular, papillary, and adenoid cystic is greater than that of women with cancers of no special type.

Women with metaplastic carcinoma or micro papillary carcinoma have a poorer prognosis

Histologic grade:
All invasive carcinomas are graded using Histologic Score composed of Nuclear grade, tubule formation, and mitotic rate

Proliferative rate:
measured by mitotic counts.
Highly proliferative tumors have poorer prognosis but may respond better to chemotherapy

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ER,PR,HER2

ER & PR:

Eighty percent of carcinomas that are both ER-and PR- positive respond to hormonal manipulation

40% of CA positive for only ER or PR respond.

Strongly ER-positive cancers are less likely to respond to chemotherapy.

cancers that fail to express either ER or PR have a less than 10% likelihood of responding to hormonal therapy but are more likely to respond to chemotherapy.

HER2:

HER2 overexpression is associated with poorer survival predictor of response to agents that target this receptor.

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Stages of breast ca

Stage 0: DCIS or LCIS, with 5-year survival rate (5YSR):92%

Stage I: Invasive ca up to 2 cm (including ca in situ with micro invasion) without LN involvement (5YSR:87%).

Stage II: Invasive ca up to 5 cm with up to 3 involved axillary LNs or invasive ca more than 5 cm without LN involvement (5YSR:75%).

Stage III. Invasive ca up to 5 cm with 4 or > involved axillary LNs; invasive ca more than 5 cm with LN involvement; invasive ca with 10 or more involved axillary LNs; invasive ca with involvement of the ipsilateral internal mammary LNs; or invasive ca with skin involvement (edema, ulceration, or satellite skin nodules), chest wall fixation, or clinical inflammatory ca (5YSR:46%).

Stage IV. Any Ca B with distant metastases(5YSR: 13%).
Why some cancers recur following postoperative therapy whereas others do not?
Remains unknown & a mystery.

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تم بحمد الله

ادعوا لأهلنا في غزة بالنصر و الرحمة
واذكروني بدعوة

وقل رب زدني علماً