# AJCC 8th Edition Staging

## **Breast Staging**

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American Joint Committee on Cancer

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# Learning Objectives

Select appropriate stage group table

Examine prognostic stage group criteria

Inspect clarifications for assigning categories

Identify changes in breast staging



#### Learning Assessments

- Testing effect or retrieval practice
  - Testing yourself on idea or concept to help you remember it
- Many experts have agreed for centuries
  - Act of retrieving info over and over, makes it retrievable when needed
  - Aristotle: exercise in repeatedly recalling strengthens memory
- Why retrieval/quizzing slows forgetting, helps remembering
  - Memory is dynamic (keeps changing), retrieval helps it change
  - Test often for better results
- Quizzes
  - Pretest as part of registration
  - Quiz during lecture
  - Posttest emailed weeks later to assess retention
  - Also assesses clarity of instruction and instructor



# Selecting Appropriate Stage Group Table



- May never use anatomic stage group table
  - Even if prognostic factor categories are missing
  - Even if stage group will be unknown
  - Will skew stage group data

- ONLY for global regions where biomarker tests unavailable
- Cancer registries in U.S. must use prognostic tables



## Stage Group Tables

- Clinical prognostic stage group table
  - Used for all patients with diagnostic workup for ca

- Pathological prognostic stage groups
  - Used when surgical resection is initial treatment
  - Does not apply to surgical resection following neoadjuvant therapy

No stage group table for posttherapy staging



# Genomic Profiles in Staging

- Other (non-Oncotype Dx) genomic profiles/multigene panels
  - Only used for patient care
  - Because Level I data not available at this time
  - These profiles not used for assigning prognostic stage

- Specific chapter wording for other multigene panels:
  - ...low-risk score/range, regardless of T size, places the tumor into the same prognostic category as T1a–T1b N0 M0.



## Genomic Profiles in Staging

- Pathological prognostic staging with Oncotype Dx score <11</li>
  - Modify stage group to IA as indicated
  - Assign with Genomic Profile for Pathologic Prognostic Staging table

And TNM is A	And Grade is	And HER2 Status is	And ER Status is	And PR Status is	Then the Pathological Prognostic Stage Group is
T1 N0 M0 A	Any	Negative	Positive	Any	IA

- Other genomic profiles used to guide patient care
  - May not be used at this time to modify stage group
  - Assign using Pathological Prognostic Stage table
  - Do not use genomic profile table



#### Scenario

- Pt with 21mm UOQ tumor and negative axilla. Ductal ca, SBR 6 points, HER2 neg, ER/PR positive, breast bx.
  - Lumpectomy and SLNB showing 21mm tumor, SBR 7 points, 3 sentinel nodes negative.

- Clinical: cT2 cN0 cM0 Gr2 HER2- ER/PR+ stage grp IB
- Pathological: pT2 pN0(sn) cM0 Gr2 HER2- ER/PR+ stage grp IA
- Note different stage groups for clinical and pathological



# Prognostic Stage Group Criteria



# Grade Category in Stage Group

- Must be Nottingham for invasive ca, not nuclear grade
  - Assign G1 G3

- Nuclear grade
  - Just one of three components of Nottingham
  - Least reproducible of three components
  - Must not use for grade category to assign stage group
  - If nuclear grade, code as A-D, stage group not assigned



#### Grade for Invasive & In Situ

#### **Invasive**

G	G Definition
GX	Grade cannot be assessed
G1	Low combined histologic grade (favorable), SBR score of 3–5 points
G2	Intermediate combined histologic grade (moderately favorable); SBR score of 6–7 points
G3	High combined histologic grade (unfavorable); SBR score of 8–9 points

#### In Situ

G	G Definition
GX	Grade cannot be assessed
G1	Low nuclear grade
G2	Intermediate nuclear grade
G3	High nuclear grade



## Biomarkers for Stage Group

Collection of Biomarkers (Hormone receptor assays and HER2 assay)

The Expert Panel determined that all invasive carcinomas should have estrogen receptor (ER), progesterone receptor (PR), and human epidermal growth factor receptor 2 (HER2) status determined by appropriate assays whenever possible.

- Biomarkers consist of
  - Hormone receptors ER & PR
  - HER2
- Assay results for assigning stage
  - Most often only tested on biopsy specimen
  - If retested on resection specimen, positive results take priority
  - If bx specimen not tested, use resection tests for clinical staging



#### Scenario

 Pt with 21mm UOQ tumor and negative axilla. Ductal ca, nuclear gr 2, breast bx.

Lumpectomy and SLNB showing 21mm tumor, SBR 7 points, 3 sentinel nodes negative, HER2 neg, ER/PR positive.

- Clinical: cT2 cN0 cM0 GrX HER2- ER/PR+ stage grp 99
- Pathological: pT2 pN0(sn) cM0 Gr2 HER2- ER/PR+ stage grp IA
- Grade clinical may be coded as B, but not used for AJCC



# Clarifications



## Stage Groups Tables Differ

- Stage group tables different for clinical & pathological
  - Same TNM G Biomarker combinations not same c & p group
  - Based on outcome data
- Examples
  - Clinical
     T2 N0 M0 G3 HER2- ER/PR+ stage IIA
  - Pathological T2 N0 M0 G3 HER2- ER/PR+ stage IB
  - Clinical
     T3 N0 M0 G2 HER2- ER/PR- stage IIIB
  - Pathological T3 N0 M0 G2 HER2- ER/PR- stage IIB
- Reason for differences between clinical & pathological group
  - Tumor size may vary between imaging and resection
  - Negative nodes on exam/imaging contain mets when resected

# Posttherapy Staging Critical

- Critical posttherapy staging regardless of response
  - Even if tumor does not respond, stays the same
  - Even if tumor larger or more nodal involvement
  - Not considered progression of disease
  - Posttherapy staging must be assigned
- Need data on all patients undergoing neoadjuvant therapy
  - Not just those with partial or complete response
  - Do not skew data by eliminating those with no response
- Assign ypT, ypN, and c/pM categories
- No posttherapy stage group



# Breast Neoadjuvant Therapy

- Breast neoadjuvant therapy
  - Must meet standard guidelines, such as NCCN or ASCO
  - Usually 4-6 cycles of chemo, sometimes more
  - Usually 4-6 months of endocrine therapy, may be up to 1 year
  - Short course endocrine therapy does NOT qualify
  - Rule for staging, not for registry treatment data items
- Must assign posttherapy staging
  - Even if chemo changed to different group of chemo drug
  - Even if endocrine therapy changed
- CoC states surgical resection coded even if no response



#### Stage Data

Stage is more than just the group

- Assign T, N, M, and prognostic factor categories
  - Even if stage group doesn't exist
  - Especially with missing info and stage group can't be assigned

- Value not tied only to stage group
  - Studies performed on TNM data, not just stage groups
  - Critical comparisons between cT cN and ypT ypN



#### **Breast Sentinel Node Procedure**

- Sentinel lymph node procedure includes
  - Sentinel nodes with dye/radiotracer
     and
  - Non-sentinel nodes, palpably abnormal, without dye/radiotracer
  - Use (sn) suffix for N category
- Sentinel node procedure results
  - Pathologist reports to surgeon in Operating Room
  - Surgeon needs results to decide if node dissection needed
- Waiting for results
  - If you don't wait for SLN path results, no reason to perform it
  - Next steps based on frozen sections of those sentinel nodes



#### Scenario

Pt with 48mm UOQ tumor and enlarged axillary node.
 Ductal ca, SBR 6 points, HER2 neg, ER/PR positive, breast bx. Axillary node FNA positive for mets.

Neoadjuvant chemo. No response, tumor 60mm, ycT3.

Mastectomy showing 58mm tumor, SBR 7 points, 4/10 axillary nodes positive.

- Clinical: cT2 cN1(f) cM0 Gr2 HER2- ER/PR+ stage grp IIA
- Posttherapy: ypT3 ypN2a cM0 Gr2 HER2- ER/PR+ stage grp not assigned



# Additional Changes to Breast



#### LCIS

#### Definition of Primary Tumor (T)

Lobular carcinoma *in situ* (LCIS) is removed as a pTis category for T categorization. Lobular carcinoma *in situ* is treated as a benign entity and is removed from TNM staging.

- Lobular carcinoma in situ no longer assigned a stage
  - Please discuss reportability with your standard setter

- In situ category may only be
  - Tis (DCIS)
  - Tis (Paget)



#### Rounding Caution for Tumor Size

Definition of Primary Tumor (T)

The general rules for rounding to the nearest millimeter do not apply for tumors between 1.0 and 1.5 mm, so as to not classify these cancers as microinvasive (T1mi) carcinomas (defined as invasive tumor foci 1.0 mm or smaller). Tumors >1 mm and <2 mm should be reported rounding to 2 mm.

- Breast exception T category
  - ->1.0 mm to 1.4 mm rounded to 2 mm
  - Avoid assigning "microinvasion" category to cancer >1.0 mm
  - Other sizes rounded for T category assignment
    - Round down between 1 and 4
    - Round up between 5 and 9
- Critical for prognosis and data analysis
  - T1mi "microinvasion" must only represent ≤1 mm



#### Scenario

- Pt's screening mammogram identified 1.1mm tumor with negative axilla. Ductal ca, intermediate combined histologic grade, HER2 neg, ER/PR positive, breast bx.
  - Lumpectomy and SLNB showing no residual tumor, 2 sentinel nodes negative, 1 non-sentinel node negative.

- Clinical: cT1a cN0 cM0 Gr2 HER2- ER/PR+ stage grp IA
- Pathological: pT1a pN0(sn) cM0 Gr2 HER2- ER/PR+ stage grp IA



# Information and Questions on AJCC Staging

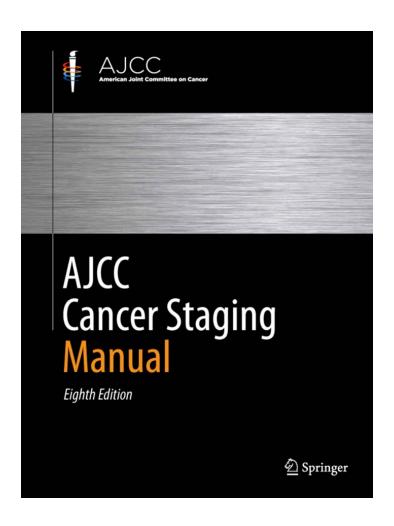


#### **AJCC** Web site

https://cancerstaging.org

- Ordering information
  - Cancerstaging.net

- General information
  - Education
  - Articles
  - Updates





#### **CAnswer Forum**

- Submit questions to AJCC Forum
  - NEW 8th Edition Forum
  - 7<sup>th</sup> Edition Forum will remain
  - Located within CAnswer Forum
  - Provides information for all
  - Allows tracking for educational purposes

http://cancerbulletin.facs.org/forums/





# Quiz



# Summary



## Summary

Comprehend appropriate stage group table usage

Identify prognostic stage group criteria

Interpret clarifications for assigning categories

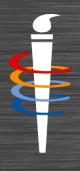
Examine changes in breast staging



# Thank you

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