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#### Q&A

Please submit all questions concerning the webinar content through the Q&A panel.

If you have participants watching this webinar at your site, please collect their names and emails.

We will be distributing a Q&A document in about one week. This document will fully answer questions asked during the webinar and will contain any corrections that we may discover after the webinar.

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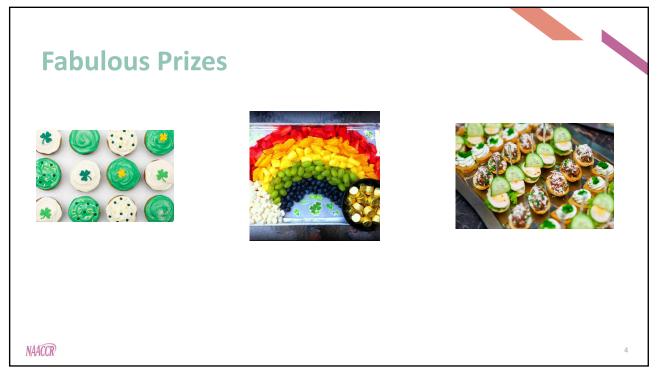
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#### **Resources**

- Text Template
- SSDI Manual: Breast Chapter

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#### **Guest Presenters**

- Nancy Etzold, CTR
  - Director, Cancer Registry, Oklahoma University Health
- Laney Bomberger-Schmotzer, CTR
  - Cancer Registry Coordinator, Oklahoma University Health

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#### **Terrific Text**

- Abstract
- Codes
  - Benefits
  - Drawbacks

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#### **Webinar Agenda**

- Overview Text
- Quiz 1 Text
- Overview Casefinding & Sequence
- Quiz 2 Casefinding & Sequence
- Overview Class of Case
- Quiz 3 Class of Case
- BREAK
- Quiz 4 Terminology
- Overview Breast Site-Specific Data Items
- Quiz 5 Breast SSDI's
- Quiz 6 Wrap up

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#### **TEXT Guidelines**

- Supports and Validates
- Standardization
- CCO
- Just the facts
- Lay the groundwork

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#### **History & Physical Exam Text**

- History of present illness
- Physical exam findings
  - Dates
  - Primary Site
  - Lymph Nodes
  - Extent of disease
- Don't include

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#### **Radiology Endoscopic Procedures Text**

- Endoscopic Procedures
- Radiology Procedures
  - Dates
  - Primary Site
  - Lymph Nodes
  - Extent of Disease
  - Diagnostic Impression

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#### **Laboratory Tests & Tumor Marker Text**

- Laboratory Tests
  - Date
  - Test name/type
  - Results
  - Normal value/range
- Tumor Markers
  - Screen, Monitor, or Detect
  - Determine
  - Provide

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#### **Operative Findings Text**

- Surgical Procedures
- Surgeon Observation
- Most complete assessment
  - Organs/tissues removed
  - Invasion of other tissues/organs
  - Multiple tumors
  - Discontinuous spread
  - Stage
  - Gross tumor remaining

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#### **Pathology Text**

- Microscopic Tissue Examination
- Microscopic Cell Examination
- What to record
- CoC Compliance

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#### **Treatment Text Fields**

- Unique for Treatment Modalities
- 1000 Characters
- Standard Abbreviations
- Don't rely on computer text
- No duplication
- Less is more
- Note what's missing
- · Document all related data fields

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#### **Surgery Text**

- Purpose of Surgery performed
- Type of surgical procedures
- What to record

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#### **Systemic Treatment Text**

- Chemotherapy
- Hormone
- Immunotherapy
- What to record

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#### **Transplant/Endocrine Procedure Text**

- No dedicated text field
- Procedures
- Hormone/Endocrine Surgery & Radiation
- What to record

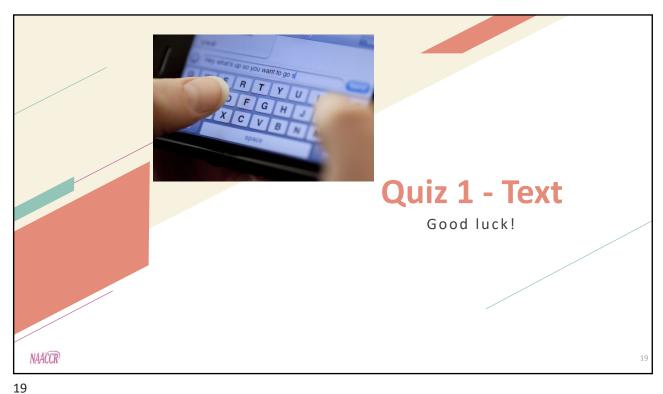
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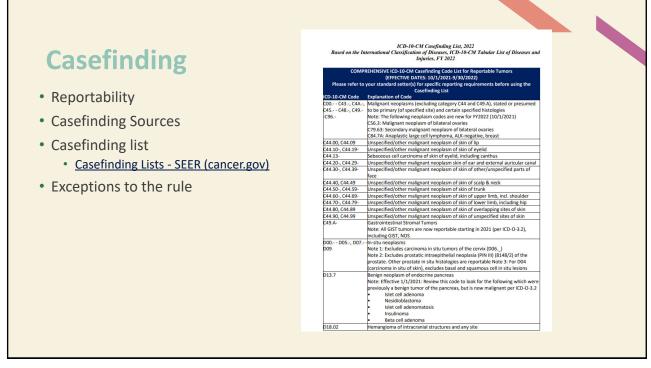
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#### **Other Therapy Text and Remarks**

- Cancer-directed treatment doesn't fit elsewhere
- What to record
- Remarks

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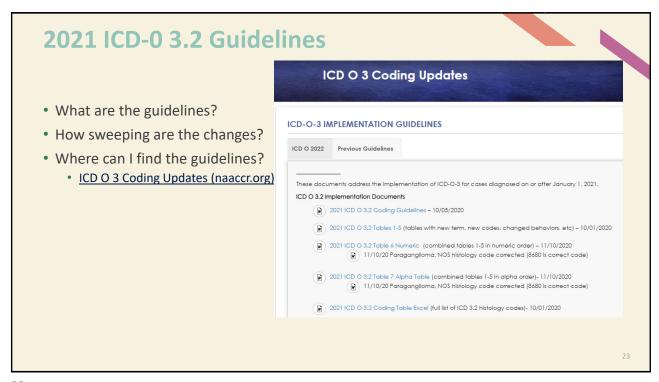


# Casefinding Methods & Sources • Active vs passive casefinding • Source documents may vary • Reliance on multiple sources

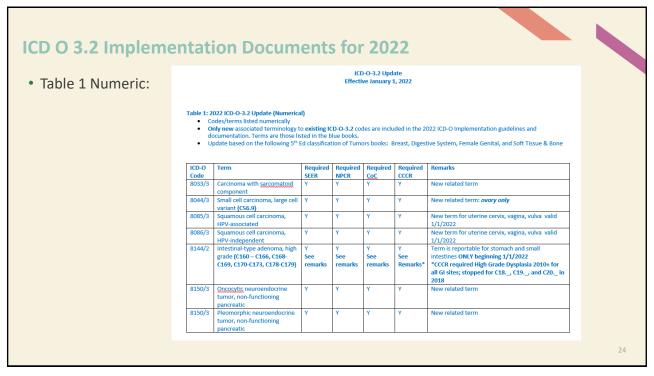
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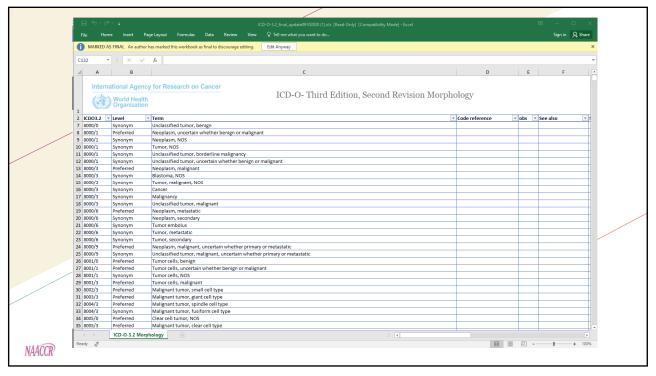


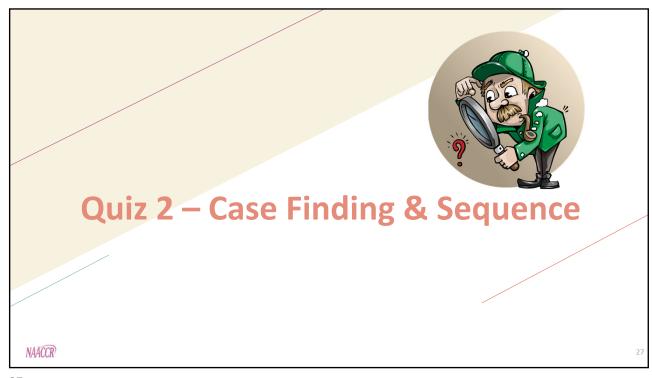
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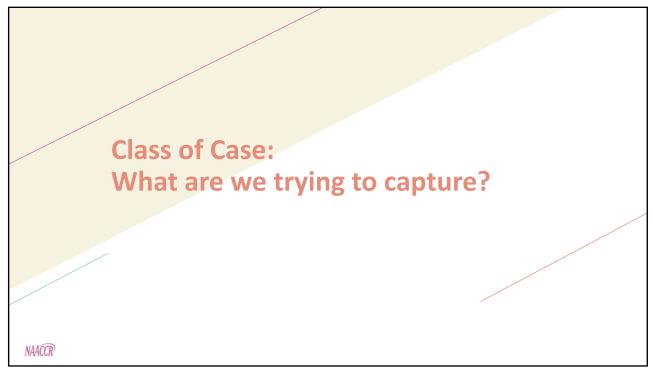


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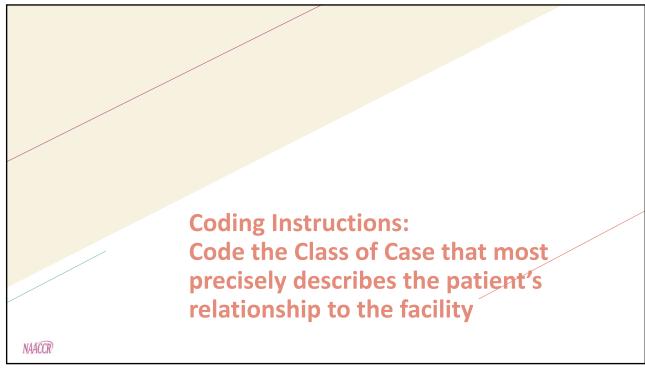
#### **Two Groups:**

• Analytic (codes 00-22)

Code	Label		
Analyt	Analytic Classes of Case (Required by CoC to be abstracted by accredited programs)		
Initial (	diagnosis at reporting facility or in a staff physician's office		
00 Initial diagnosis at the reporting facility AND all treatment or a decision not to treat elsewhere			
10	Initial diagnosis at the reporting facility or in an office of a physician with admitting privileges AND part or all of first course treatment or a decision not to treat was at the reporting facility, NOS		
11	Initial diagnosis in an office of a physician with admitting privileges AND part of first course treatment was done at the reporting facility		
12 Initial diagnosis in an office of a physician with admitting privileges AND all first of treatment or a decision not to treat was done at the reporting facility			
13	Initial diagnosis at the reporting facility AND part of first course treatment was done at the reporting facility; part of first course treatment was done elsewhere		
14	Initial diagnosis at the reporting facility AND all first course treatment or a decision not to treat was done at the reporting facility		
Initial (	diagnosis elsewhere		
20 Initial diagnosis elsewhere AND all or part of first course treatment was done at the facility, NOS			
21	Initial diagnosis elsewhere AND part of first course treatment was done at the reporting facility; part of first course treatment was done elsewhere.		
22	Initial diagnosis elsewhere AND all first course treatment or a decision not to treat was done at the reporting facility		

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Code	Nonanalytic (codes 30-49 8		
	ss of Case not required by CoC to be abstracted (May be required by Cancer Committee, state or nal registry, or other entity)		
Patie	nt appears in person at reporting facility		
30	Initial diagnosis and all first course treatment elsewhere AND reporting facility participated in diagnostic workup (for example, consult only, treatment plan only, staging workup after initial	32	Diagnosis AND all first course treatment provided elsewhere AND patient presents at reportin facility with disease recurrence or persistence (active disease)
31	diagnosis elsewhere) Initial diagnosis and all first course treatment elsewhere AND reporting facility provided intransit care; or hospital provided care that facilitated treatment elsewhere (for example, stent placement)	33	Diagnosis AND all first course treatment provided elsewhere AND patient presents at reportir facility with disease history only (disease not active)
		34	Type of case not required by CoC to be accessioned (for example, a benign colon tumor) AND initial diagnosis AND part or all of first course treatment by reporting facility
		35	Case diagnosed before program's Reference Date AND initial diagnosis AND all or part of first course treatment by reporting facility
		36	Type of case not required by CoC to be accessioned (for example, a benign colon tumor) AND initial diagnosis elsewhere AND all or part of first course treatment by reporting facility
		37	Case diagnosed before program's Reference Date AND initial diagnosis elsewhere AND all or part of first course treatment by facility
		38	Initial diagnosis established by autopsy at the reporting facility, cancer not suspected prior to death
		Patier	it does not appear in person at reporting facility
		40	Diagnosis AND all first course treatment given at the same staff physician's office
		41	Diagnosis and all first course treatment given in two or more different offices of physicians wi admitting privileges
		42	Nonstaff physician or non-CoC accredited clinic or other facility, not part of reporting facility, accessioned by reporting facility for diagnosis and/or treatment by that entity (for example, hospital abstracts cases from an independent radiation facility)
		43	Pathology or other lab specimens only
		49	Death certificate only
		99	Nonanalytic case of unknown relationship to facility (not for use by CoC accredited cancer programs for analytic cases).



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#### Class of case 00 vs 10

Is the initial cancer diagnosis rendered at your facility?

**Did the patient receive treatment?** 

Did all or a portion of the treatment take place at your facility?

Did the patient decide at your facility that he or she did not want to receive treatment?

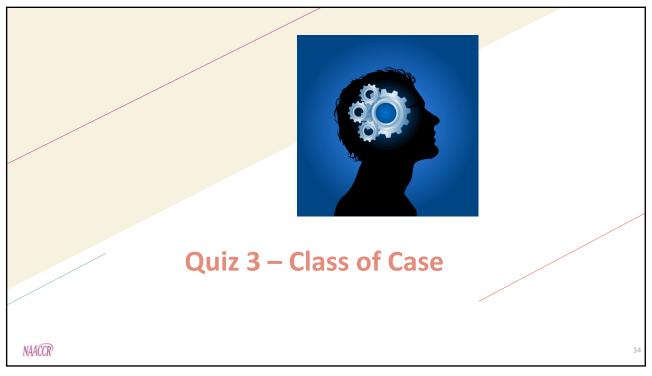
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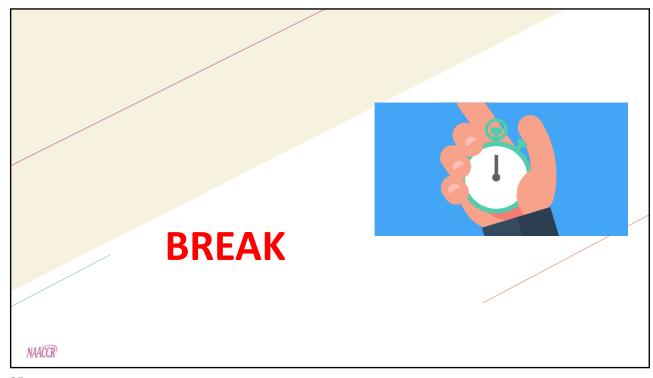
#### **Reportable-by-Agreement Cases**

#### Examples:

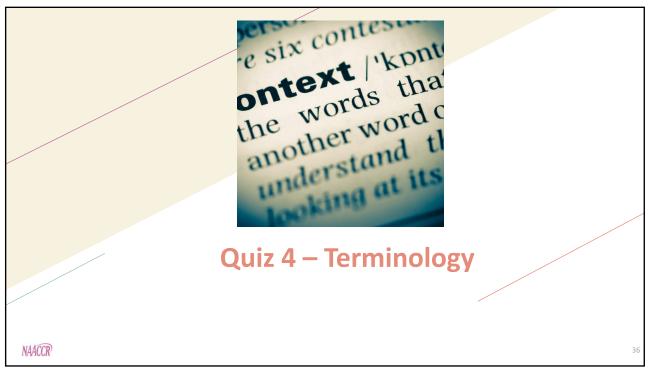
- The cancer committee requests abstracting and follow-up of Class of Case 30 cases.
- The state central registry requests abstracting and reporting of pathology-only cases.

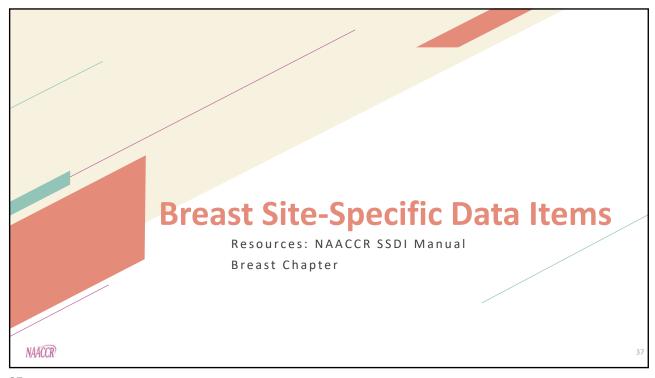
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#### **Site-Specific Data Items (SSDI)**

- STORE Pages 229-230
- NAACCR <a href="https://www.naaccr.org/SSDI/SSDI-Manual.pdf">https://www.naaccr.org/SSDI/SSDI-Manual.pdf</a>
- Cancer Registry Software

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# **Estrogen Receptor Progesterone Receptor**

- Percentage nuclear positivity
- Intensity of staining
- Reported as specific number or range

ER or PR Status	51-60%
Positive	61-70%
Percentage of cells with nuclear positivity#	71-80%
Specify: %	81-90%
-OR-	91-100%
Range (Note A)	+ Average intensity of staining:
1-10% (specify): %#	+ Weak
11-20%	<b></b>
21-30%	+ Moderate
31-40%	+ Strong
41-50%	Negative

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#### **ER/PR Summary**

- Path Interpretation
- Code 0 when reported as negative or normal
- Code 1 when reported as positive or elevated
- Code 7 when test was ordered, but results not available
- Code 9 when
  - Reported as borderline, undetermined whether positive or negative
  - Cannot be determined by pathologist (inadequate specimen)
  - Unknown whether test was performed
  - Patient has only clinical diagnosis (no tissue examined)

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#### **Estrogen Receptor Percent Positive or Range**

- Affects AJCC Staging
- Use same report as ER Summary

Code	Description
000	ER negative, or stated as less than 1%
001-100	1-100 percent
R10	Stated as 1-10%
R20	Stated as 11-20%
R30	Stated as 21-30%
R40	Stated as 31-40%
R50	Stated as 41-50%
R60	Stated as 51-60%
R70	Stated as 61-70%
R80	Stated as 71-80%
R90	Stated as 81-90%
R99	Stated as 91-100%
XX8	Not applicable: Information not collected for this case
	(If this item is required by your standard setter, use of code XX8 will result in an edit error.)
XX9	Not documented in medical record
	ER (Estrogen Receptor) Percent Positive or Range not assessed or unknown if assessed

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#### **Progesterone Receptor % Positive or Range**

- Affects AJCC Staging
- Use same report at PR Summary

Code	Description	
000	PR negative, or stated as less than 1%	
001-100	1-100 percent	
R10	Stated as 1-10%	
R20	Stated as 11-20%	
R30	Stated as 21-30%	
R40	Stated as 31-40%	
R50 Stated as 41-50%		
R60	Stated as 51-60%	
R70	Stated as 61-70%	
R80	Stated as 71-80%	
R90	Stated as 81-90%	
R99	Stated as 91-100%	
XX8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code XX8 will result in an edit	
	error.)	
XX9	Not documented in medical record	
	PR (Progesterone Receptor) Percent Positive or Range not assessed or unknown if	
	assessed	

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#### **Allred Score**

## Quantifies ER and PR Scores Calculated by adding Proportion & Intensity Scores

Proportion Score	Positive Cells, %
0	0
1	<1
2	1 to 10
3	11 to 33
4	34 to 66
5	≥67

Intensity	Intensity Score
None	0
Weak	1
Intermediate/Moderate	2
Strong	3

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#### **Estrogen Receptor Allred Score**

- Physician statement
- Use same record as ER Summary

Code	Description
00	Total ER Allred score of 0
01	Total ER Allred score of 1
02	Total ER Allred score of 2
03	Total ER Allred score of 3
04	Total ER Allred score of 4
05	Total ER Allred score of 5
06	Total ER Allred score of 6
07	Total ER Allred score of 7

Code	Description
08	Total ER Allred score of 8
X8	Not applicable: Information not collected for this case (If this item is required by your standard setter, use of code X8 will result in an edit error.)
Х9	Not documented in medical record ER (Estrogen Receptor) Total Allred Score not assessed, or unknown if assessed

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#### **Progesterone Receptor Total Allred Score**

- Physician statement only
- Same Report

Code	Description
00	Total PR Allred score of 0
01	Total PR Allred score of 1
02	Total PR Allred score of 2
03	Total PR Allred score of 3
04	Total PR Allred score of 4
05	Total PR Allred score of 5
06	Total PR Allred score of 6
07	Total PR Allred score of 7

Code	Description
08	Total PR Allred score of 8
X8	Not applicable: Information not collected for this case
	(If this item is required by your standard setter, use of code X8 will result in an edit
	error.)
X9	Not documented in medical record
	PR (Progesterone Receptor) Total Allred Score not assessed, or unknown if
	assessed

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#### HER2

- Human Epidermal Growth Factor Receptor 2 (HER2)
- Gene Amplification
- Prognostic Indicator

Result	Criteria
Negative (Score 0)	No staining observed
	or
	Incomplete, faint/barely perceptible membrane staining in ≤10% of invasive tumor cells
Negative (Score 1+)	Incomplete, faint/barely perceptible membrane staining in >10% of invasive tumor cells*
Equivocal (Score 2+)	Incomplete and/or weak to moderate circumferential membrane staining in >10% of invasive tumor cells
	or
	Complete, intense, circumferential membrane staining in ≤10% of invasive tumor cells*
Positive (Score 3+)	Complete, intense, circumferential membrane staining in >10% of invasive tumor cells*

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### **HER2 IHC & Summary**

Code	Description	
0	Negative (Score 0)	
1	Negative (Score 1+)	
2	Equivocal (Score 2+)	
	Stated as equivocal	
3	Positive (Score 3+)	
	Stated as positive	
4	Stated as negative, but score not stated	
7	Test ordered, results not in chart	
8	Not applicable: Information not collected for this case	
(If this item is required by your standard setter, use of code 8 will result in a		
9	Not documented in medical record	
	Cannot be determined (indeterminate)	
	HER2 IHC Summary not assessed or unknown if assessed	

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#### **HER2 ISH & Summary**

Code	Description	
0	Negative [not amplified]	
2	Equivocal	
3	Positive [amplified]	
7	Test ordered, results not in chart	
8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code 8 will result in an edit	
	error.)	
9	Not documented in medical record	
	Results cannot be determined (indeterminate)	
	HER2 ISH Summary not assessed or unknown if assessed	

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#### **HER2 ISH** Reporting Results of HER2 Testing by In Situ Hybridization (single-probe assay) Criteria Negative (not Average HER2 copy number <4.0 signals/cell amplified) Equivocal Average *HER2* copy number ≥4.0 and <6.0 signals/cell Positive (amplified) Average HER2 copy number ≥6.0 signals/cell Reporting Results of HER2 Testing by In Situ Hybridization (dual-probe assay) Criteria Result Negative (not HER2/CEP17 ratio <2.0 AND average HER2 copy number <4.0 signals/cell amplified) Equivocal HER2/CEP17 ratio <2.0 AND average HER2 copy number ≥4.0 but <6.0 Positive (amplified) HER2/CEP17 ratio ≥2.0 (regardless of average HER2 copy number) Average HER2 copy number ≥6.0 signals/cell (regardless of ratio)

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#### **HER2 Overall Summary**

- Pathologist's interpretation
- Prior to neoadjuvant therapy
- Multiple specimens
- Exceptions

Code	Description	
0	Negative (Score 0)	
1	Negative (Score 1+)	
2	Equivocal (Score 2+)	
	Stated as equivocal	
3	Positive (Score 3+)	
	Stated as positive	
4	Stated as negative, but score not stated	
7	Test ordered, results not in chart	
8 Not applicable: Information not collected for this case		
(If this item is required by your standard setter, use of code 8 will result in		
9 Not documented in medical record		
	Cannot be determined (indeterminate)	
	HER2 IHC Summary not assessed or unknown if assessed	

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# HER2 ISH Single & Dual Probe Copy Number

Code	Description	
0.0-99.9 Reported HER2 copy number of 0.0-99.9		
XX.1	XX.1 Reported HER2 copy number of 100 or greater	
XX.7	XX.7 Test ordered, results not in chart	
XX.8	Not applicable: Information not collected for this case (If this item is required by your standard setter, use of code XX.8 will result in an edit	
error.)		
XX.9	Not documented in medical record  Cannot be determined (indeterminate)HER2 ISH Single Probe Copy Number not assessed or unknown if assessed	

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#### **HER2 ISH DUAL PROBE RATIO**

Code	Description	
0.0-99.9	Ratio of 0.0 to 99.9	
XX.2	Less than 2.0	
XX.3	Greater than or equal to 2.0	
XX.7	Test ordered, results not in chart	
XX.8 Not applicable: Information not collected for this case (If this item is required by your standard setter, use of code XX.8 will res		
XX.9	edit error.)  Not documented in medical record	
77.3	Results cannot be determined (indeterminate)HER2 ISH Dual Probe Ratio not assessed or unknown if assessed	

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#### **Multigene Signature Method**

• Node-Negative Female Breast cancer

Code	Description
1	Mammaprint
2	PAM50 (Prosigna)
3	Breast Cancer Index
4	EndoPredict

5	Test performed, type of test unknown	
6	Multiple tests, any tests in codes 1-4	
7	Test ordered, results not in chart	
8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code 8 will result in an edit error.)	
9	Not documented in medical record	
	Multigene Signature Method not assessed or unknown if assessed	

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#### **Oncotype DX**

- Low risk: Recurrence Score result less than 18: The patient has a lower risk of having a
  recurrence, assuming 5 years of hormonal therapy is given. Chemotherapy is likely to have little
  or no benefit.
- Intermediate Risk: Recurrence Score result between 18 and 30: The patient has a tumor that is in the middle of the risk spectrum reflecting that biology is continuous and not all patients have a low or a high recurrence risk, assuming 5 years of hormonal therapy is given. The likelihood of distant recurrence and benefit from chemotherapy increases with an increase in the Recurrence Score result.
- High risk: Recurrence Score result greater than or equal to 31: The patient has a high risk of
  distant recurrence, assuming 5 years of hormonal therapy and is likely to benefit from
  chemotherapy.

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## Oncotype DX Recurrence Score & Risk Level DCIS

Code	Description	
000-100	Enter actual recurrence score between 0 and 100	
XX6	Not applicable: invasive case	
XX7	Test ordered, results not in chart	
XX8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code XX8 will result in an edit	
	error.)	
XX9	Not documented in medical record	
	Oncotype Dx Recurrence Score-DCIS not assessed or unknown if assessed	

Code	Description	
0	Low risk (recurrence score 0-38)	
1	Intermediate risk (recurrence score 39-54)	
2	High risk (recurrence score greater than or equal to 55)	
6	Not applicable: invasive case	
7	Test ordered, results not in chart	
8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code 8 will result in an edit error.)	
9	Not documented in medical record	
	Oncotype Dx Risk Level-DCIS not assessed or unknown if assessed	

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#### **Multigene Signature Method**

• Node-Negative Female Breast cancer

Code	Description
1	Mammaprint
2	PAM50 (Prosigna)
3	Breast Cancer Index
4	EndoPredict

5	Test performed, type of test unknown	
6	Multiple tests, any tests in codes 1-4	
7	Test ordered, results not in chart	
8	Not applicable: Information not collected for this case	
	(If this item is required by your standard setter, use of code 8 will result in an edit error.)	
9	Not documented in medical record	
	Multigene Signature Method not assessed or unknown if assessed	

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# Oncotype DX Recurrence Score and Risk Level Invasive

Code	Description	
000-100	Enter actual recurrence score between 0 and 100	
XX4	tated as less than 11	
XX5	Stated as equal to or greater than 11	
XX6	Not applicable: in situ case	
XX7	Test ordered, results not in chart	
XX9	Not documented in medical record	
	Oncotype Dx Recurrence Score-Invasive not assessed or unknown if assessed	

Code	Description
0	Low risk (recurrence score 0-17)
1	Intermediate risk (recurrence score 18-30)
2	High risk (recurrence score greater than or equal to 31)
6	Not applicable: DCIS case
7	Test ordered, results not in chart
8	Not applicable: Information not collected for this case
	(If this item is required by your standard setter, use of code 8 will result in an edit error.)
9	Not documented in medical record
	Oncotype Dx Risk Level-Invasive not assessed or unknown if assessed

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#### Ki- 67

- Physician Statement
- Values
- Nodes or Mets

Code	Description
0.0-100.0	0.0 to 100.0 percent positive: enter percent positive
XXX.7	Test done, actual percentage not stated
XXX.8	Not applicable: Information not collected for this case
	(If this item is required by your standard setter, use of code XXX.8 will result in an edit error.)
XXX.9	Not documented in medical record
	Ki-67 (MIB-1) not assessed or unknown if assessed

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#### **Positive Lymph Node Axillary Level**

- Include
- Exclude

Code	Description
00	All ipsilateral axillary nodes examined negative
01-99	1 - 99 nodes positive
	(Exact number of nodes positive)
X1	100 or more nodes positive
X5	Positive nodes, number unspecified
X6	Positive aspiration or needle core biopsy of lymph node(s)
X8	Not applicable: Information not collected for this case
	(If this item is required by your standard setter, use of code X8 will result in an edit error.)
Х9	Not documented in medical record
	Level I-II axillary nodes not assessed or unknown if assessed

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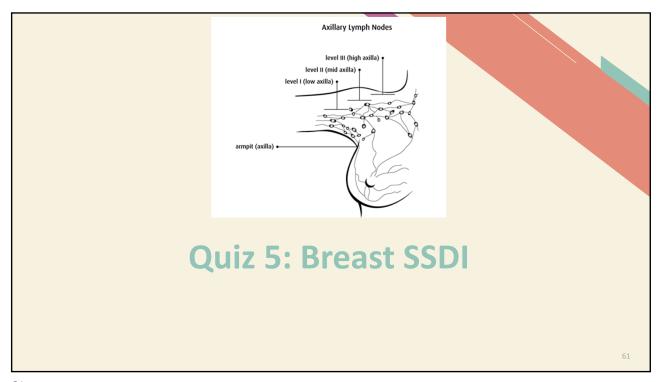
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#### **Response to Neoadjuvant Therapy**

- Systemic or Radiation Treatment
- Clinician's Statement not AJCC 8<sup>th</sup> Edition definitions

Code	Description
0	Neoadjuvant therapy not given
1	Stated as complete response (CR)
2	Stated as partial response (PR)
3	Stated as response to treatment, but not noted if complete or partial
4	Stated as no response (NR)
8	Not applicable: Information not collected for this case (If this item is required by your standard setter, use of code 8 will result in an edit error.)
9	Not documented in medical record Response to neoadjuvant therapy not assessed or unknown if assessed

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#### **CE Certificate Quiz/Survey**

**CE Phrase** 

Link

https://survey.alchemer.com/s3/6563871/Boot-Camp-2022

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#### **Coming UP...**

- Hematopoietic and Lymphocytic Neoplasms 2022
  - Guest Host: Denise Harrison, CTR; Louanne Currence, RHIT, CTR
  - 4/14/2022
- Colon 2022
  - · Guest Host: Janice Smith
  - 5/05/2022

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