

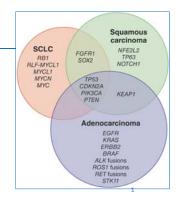
NEW SITE-SPECIFIC FIELDS "REQUIRED FOR STAGING" AJCC 8TH ED.



FCDS Annual Educational Conference

Orlando, Florida July 28, 2017

Steven Peace, CTR



Outline

- 2018 Transition from CS SSFs to Individual Site-Specific Prognostic Factor Fields (SSFs)
- Locating SSFs in the AJCC Cancer Staging Manual, 8th edition
- Historical SSFs Still Required for Staging Moved into New Fields
- New Site Specific Fields Shared Across Chapters Moved into New Fields
- New Site Specific Prognostic Factor Fields Required to Assign Stage Group
- Pending Site Specific Prognostic Factor Fields Recommended for Clinical Care
- Using the Site Specific Fields Required for Staging to Assign Stage Group
- What if the Required SSF is missing? Can I still assign Stage Group?
- QC NOTE: Analytic Cases with No Attempt to Code SSFs Increasing in Frequency
- Prognostic Factors Manual & Training
- Questions



Transition from CS SSFs to Individual Fields

Appendix H - 2016 FCDS Required CSv02.05 Site Specific Factors (SSFs)

Schema Number	Schema Name	TNM/SS Required	2013 FCDS Required	Additional CoC Required
116	AdnexaUterineOther	None	None	None
147	AdrenalGland	None	None	None
66	AmpullaVater	None	None	None
59	Anus	None	None	None
50	Appendix	2,11	2,11	1,3
65	BileDuctsDistal	25	25	None
61	BileDuctsIntraHepat	10	10	1,2,11
63	BileDuctsPerihilar	25	25	11
68	BillaryOther	None	None	None
128	Bladder	2	2	1,3
95	Bone	None	None	3
143	Brain	None	1	4,5,6
106	Breast	3,4,5	1,2,3,4,5,8,9,11,13,14,15,16	6,7,10,12,21,22,23
25	BuccalMucosa	1	1	3,4,5,6,9,11
51	CarcinoidAppendix	2	2	None
110	Cervix	None	None	1
144	CNSOther	None	1	4,5,6
53	Colon	2	2	1,3,4,6,8,9
131	Conjunctive	1	1	None
112	CorpusAdenosarcoma	2	2	1,3,4,5,6
111	CorpusCarcinoma	2	2	1,3,4,5,6
118	CorpusSarcoma	2	2	1,3,4,5,6
64	CysticDuct	25	25	None

New Manual to Include ALL Site-Specific Fields Description of Test, Instructions and Codes ALL Previous CS SSFs in New Fields ALL NEW Required for Staging Fields - Approved New Optional Prognostic Site-Specific Fields - Not Approved - Yet

PROPOSED Full Data Item Name	PROPOSED NAACCR Data Item Name =	Chapter Assignment =
Adenopathy	Adenopathy	
Alpha Fetoprotein (AFP) Post-Orchiectomy Lab Value	AFP Post-Orch Lab Value	
Alpha Fetoprotein (AFP) Post-Orchiectomy Range	AFP Post-Orch Range	
Alpha Fetoprotein (AFP) Pre-Orchiectomy Lab Value	AFP Pre-Orch Lab Value	
Alpha Fetoprotein (AFP) Pre-Orchiectomy Range	AFP Pre-Orch Range	
Alpha Fetoprotein (AFP) Pretreatment Interpretation	AFP PreTX Interpretation	Liver
Alpha Fetoprotein (AFP) Pretreatment Lab Value	AFP PreTX Lab Value	Liver
Adenoid Cystic Carcinoma - Presence of Basaloid Pattern	Basaloid Pattern	
Anemia	Anemia	
Bilirubin Pretreatment Total Lab Value	Bilirubin PreTX LabValue	Liver
Biliribun Pretreatment Unit of Measure	Bilirubin PreTX Meas	Liver
Bone Invasion	Bone Invasion	Soft Tissue Sarcomas
Carbohydrate Antigen 125 (CA-125) Pretreatment Lab Value	CA-125 PreTX Lab Value	
Carcinoembryonic Antigen (CEA) Pretreatment Intrepretation	CEA PreTX Interpretation	Colon and Rectum
Carcinoembryonic Antigen (CEA) Pretreatment Lab Value	CEA PreTX Lab Value	Colon and Rectum
Chromogranin A (CgA) Pretreatment Lab Value	CgA PreTX Lab Value	NET
Chromosome 1p: Loss of Heterozygosity (LOH)	Chromosome 1p: LOH	
Chromosome 3 Status	Chromosome 3 Status	
Chromosome 8q Status	Chromosome 8q Status	
Chromosome 19q: Loss of Heterozygosity (LOH)	Chromosome 19q: LOH	
Circumferential Resection Margin (CRM)	CircumResecMargin	Colon and Rectum
	Adenopathy Alpha Fatoprotein (AFP) Post-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Post-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Post-Orchiectomy ange Alpha Fatoprotein (AFP) Pros-Orchiectomy tab Value Alpha Fatoprotein (AFP) Pros-Orchiectomy tab Value Alpha Fatoprotein (AFP) Pretraction Interpretation Alpha Fatoprotein (AFP) Pretraction Interpretation Alpha Fatoprotein (AFP) Pretraction Interpretation Alpha Fatoprotein (AFP) Pretraction Alpha Fatoprotein (AFP) Alpha Malaria Billiribun Pratraction (Total Lab Value Carrionembyonic Arntigen (CEA) Pretreatment Lab Value Carrionembyonic Arntigen (CEA) Pretreatment Lab Value Chromosome 15: Loss of Heteroxygosity (LOH) Chromosome 63 Status Chromosome 63 Status Chromosome 69 Status	Adenopathy Adenopathy Alpha Fatoprotein (AFP) Poat-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Poat-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Poat-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Pre-Orchiectomy Lab Value Alpha Fatoprotein (AFP) Pretratement Interpretation Alpha Fatoprotein (AFP) Pretratement Lab Value Blirobin Pretratement Total Lab Value Blirobin Pretratement Total Lab Value Blirobin Pretratement Interfatorial Carlo (CA) Pretratement Lab Value Carlo (CA) Alpha Fatoprotein (CA) Pretratement Lab Value Carlo (CA) Pretratement CA) Pretratement Lab Value Carlo (CA) Pretratement CA) Pretratement Lab Value Carlo (CA) Pretratement Lab Value Chromogome 1p. Loss of Heteroxygosity (LOH) Chromosome 3p. Status Chromosome 8g Status Chromosome 8g Status Chromosome 8g Status Chromosome 1p. LOH Chromosome 1p. LOH Chromosome 1p. LOH Chromosome 8g Status Chromosome 1p. LOH Chromosome 8g Status

- 3

Questions that can be answered by cancer biomarkers Prognestic What type of optimal does port in this this cancer? Pharmacodynamics Becurrence What's the optimal does from my body? What is the optimal does cancer? Fredictive What's the optimal does from my body? What is the optimal does cancer? Fredictive What's the optimal does cancer? Fredictive What is the optimal does cancer? Fredictive Fredictive HEPATOCELLULAR CARCINOMA HEPATOCELLULAR CARCINOMA

Tumor Marker or Genetic Alteration

Tumor Marker

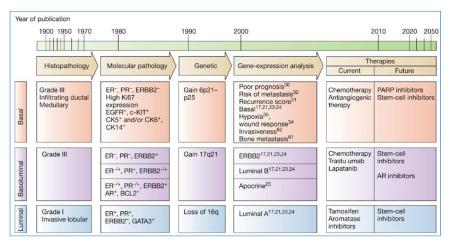
- Tumor Markers are indicators of cellular, biochemical, molecular or genetic alterations by which neoplasia can be recognized.
- Tumor markers detect the presence of tumor based on quantitative and/or qualitative measurements in blood or secretions found in cells, tissues or body fluids
- These surrogate measures of the biology of the cancer provide insight in the clinical behavior of the tumor.
- Biochemical or immunologic counterparts of differentiation states of tumor.

Genetic Alteration

- Cancer is a multigene disease that arises as a result of mutational and epigenetic changes coupled with activation of complex signaling intra and extra cellular networks.
- Alterations in 3 Classes of Genes
 - ProtoOncogenes
 - Tumor Suppressor Genes
 - DNA Repair Genes
- Types of Mutations
- Gene Rearrangement
- Point Mutations
- Gene Amplification
- Resultant effects on <u>death mechanisms</u> embedded within cells coupled with dysregulation of cell proliferation events.

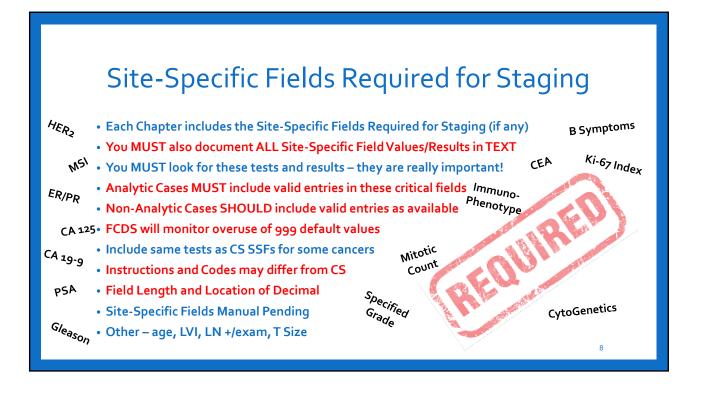
5

Comparison of the histopathology, molecular pathology, genetic, and gene-expression analysis methods used to delineate breast cancer tumor subtypes and suggested current and future therapies in a historical context



http://www.nature.com/article-assets/npg/nrclinonc/journal/v4/ng/images/ncponcogo8-f1.jpg

Locating SSFs in AJCC Staging Manual, 8th ed. AJCC Cancer Staging Manual, 8th Edition – Chapter Outline Summary of major changes and applicable disease Cancers Staged Using This Staging System Cancers NotS Staged Using This Staging System Summary of Changes ICD-O-32 Topography Codes WHO Histology Codes eral information on the disease site, such as background, trends, and recent Introduction **Chapter Specific** Anatom Primary Site(s) Regional Lymph Nodes Metastatic Sites Rules for Classification **Prognostic Factors** Imaging Pathologica o Imageue "atthological "atthological tion and discussion of non-TNM prognostic factors important in each disease Prognostic Factors Required for Stage Grouping Additional Factors Recommended for Clinical Care Emerging Factors for Clinical Care (Web Only) ostic and previous models satisfacted by the ACCS-acceptance criteria for Section in Chapter Prognostic Factors **Registry Data** Collection Variables oticine - Updates are available at www.cancerstaging.org commended factors for partitioning patients entering a clinical trial (web only) - Definition of Primary Tumor (T) - Definition of Regional tymph Node (N) - Definition of Distant Metastasis (M) Listed in Chapter AJCC Prognostic Stage Grouping Registry Data Collection Variable Grating system-to-be usued Discussion or listing of histopathologic types Survival data are the basis for anatomic stage and prognostic groups Additional figures illustrating anatomic extent of disease



Sample New SSFs - Required for Staging

Full Data Item Name: Esophagus and EGJ, Squamous Cell (including adenosquamous), Tumor Location Recommended NAACCR Data Item Name: Ecop/EGJ Tumor Epicenter Recommended NAACCR Data Item Name: Esopy(sG) Tumor Epic Data Item Length: 1 Required for AICC 8th Edition Staging; Yes Source-New data Item, submitted to UDS in 2016 and approved

16.1 Esophagus and EGJ, Squamous Cell (including adenosquamous), Tumor Location

Note 1: Physician statement of the epicenter tumor location can be used to code this data item.

Physician statement of upper, middle, or lower takes, precedence over any individual results or

Note 2: Location is defined by the position of the epicenter of the tumor in the esophagus. Information is most likely to be obtained from CT scan or operative notes.

Note 3: This information is used for pathologic staging of squamous cell carcinomas of the esophagus

Code	Description	AJCC ID
	Blank	All except 16.1
1	U: Upper (Cervical/Proximal esophagus to lower border of azygos vein)	
2	M: Middle (Lower border of azygos vein to lower border of inferior pulmonary vein)	
3	L: Lower (Lower border of Inferior pulmonary vein to stomach, including EGJ and proximal 2cm cardia of stomach)	
9	X: Esophagus, NOS Specific location not documented in medical record Specific location not assessed	

Full Data Item Name: Serum Albumin Pretreatment Level Recommended NAACCR Data Item Name: Serum Alb PreTX Level Data Item Length: 1 Required for ALCC 8* Edition Staging: Yes Source: NEW, approved by UDS in 2016

Note 1: Physician statement of serum albumin 23.5 g/dL can be used to code this data item.

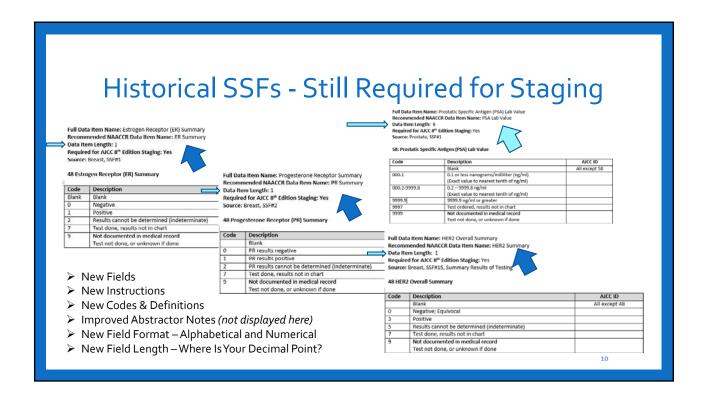
Use the cut points listed below regardless of the lab's reference range.

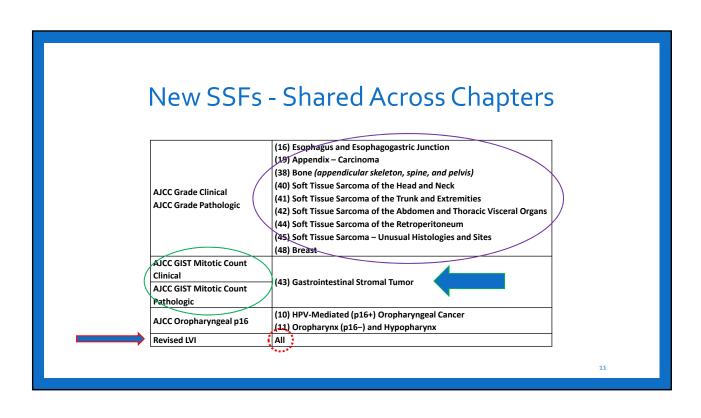
Note 2: Elevated serum albumin is defined by ≥3.5 g/dL and is part of the staging criteria.

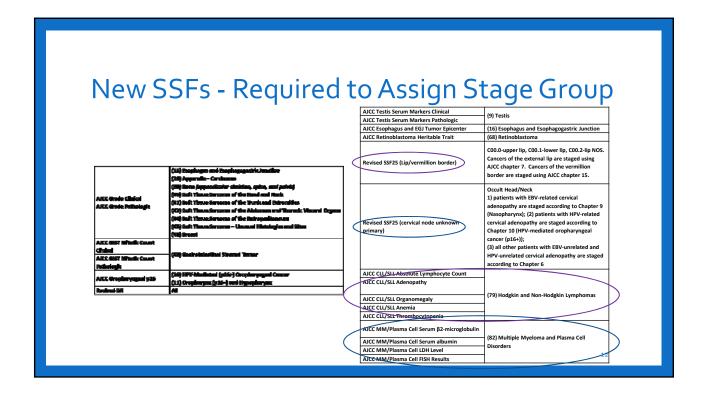
Note 3: Record this data item based on physician statement or blood test lab values at diagnosis (pre-treatment). Do not use findings from a urine test.

Note 4: If the serum albumin level determined by available lab values differs from the physician statement of < or ≥ 3.5 g/dL, the physician's statement takes precedence.

Code	Description	AJCC ID
	Blank	All except 82
1	Serum albumin <3.5 g/dL	
2	Serum albumin ≥3.5 g/dL	
7	Test done, results not in chart	
9	Not documented in medical record	
	Not tested for serum albumin, or unknown if assessed	







New SSFs - Required to Assign Stage Group

Full Data Item Name: Esophagus and EGJ, Squamous Cell (including adenosquamous), Turnor Location Recommended NAACCR Data Item Name: Esop/EGJ Turnor Epicenter Recommended NAACCR Data Item Name: Esopy(sG) Tumor Epic Data Item Length: 1 Required for AICC 8th Edition Staging; Yes Source-New data Item, submitted to UDS in 2016 and approved

16.1 Esophagus and EGJ, Squamous Cell (including adenosquamous), Tumor Location

Note 1: Physician statement of the epicenter tumor location can be used to code this data item.

Physician statement of upper, middle, or lower takes, precedence over any individual results or

Note 2: Location is defined by the position of the epicenter of the tumor in the esophagus. Information is most likely to be obtained from CT scan or operative notes.

Note 3: This information is used for pathologic staging of squamous cell carcinomas of the esophagus

Code	Description	AJCC ID
	Blank	All except 16.1
1	U: Upper (Cervical/Proximal esophagus to lower border of azygos vein)	
2	M: Middle (Lower border of azygos vein to lower border of inferior pulmonary vein)	
3	L: Lower (Lower border of Inferior pulmonary vein to stomach, including EGJ and proximal 2cm cardia of stomach)	
9	X: Esophagus, NOS Specific location not documented in medical record Specific location not assessed	

Full Data Item Name: Serum Albumin Pretreatment Level Recommended NAACCR Data Item Name: Serum Alb PreTX Leve Data Item Length: 1 Required for AICC 8° Edition Staging: Yes Source: NEW, approved by UDS in 2016

Note 1: Physician statement of serum albumin ≥3.5 g/dL can be used to code this data item.

Use the cut points listed below regardless of the lab's reference range

Note 2: Elevated serum albumin is defined by ≥3.5 g/dL and is part of the staging criteria.

Note 3: Record this data item based on physician statement or blood test lab values at diagnosis (pre-treatment). Do not use findings from a urine test.

Note 4: If the serum albumin level determined by available lab values differs from the physician statement of c or ≥ 3.5 g/dL, the physician's statement takes precedence.

Code	Description	AJCC ID
	Blank	All except 82
1	Serum albumin <3.5 g/dL	
2	Serum albumin ≥3.5 g/dL	
7	Test done, results not in chart	
9	Not documented in medical record Not tested for serum albumin, or unknown if assessed	

13

New SSFs - Required to Assign Stage Group

Full Data Item Name: Mitotic Rate GIST Clinical Recommended NAACCR Data Item Name: Mitotic Rate GIST Clin Data Item Length: 1
Required for AJCC 8th Edition Staging: Yes
Source: New data item, submitted to UDS in 2016 and approved

43 Mitotic Rate Gastrointestinal Tumor Clinical

Note 1: Record the mitotic rate as 1 (Low) or 2 (High) as indicated on the pathology report or CAP

- 1 (Low) is 5 or fewer mitoses per 5 mm², or per 50 HPF.
- 2 (High) is over 5 mitoses per 5 mm², or per 50 HPF

Note 3: If only one mitotic rate is available, and is assessed during the AJCC clinical staging timeframe, record this value for Mitotic Rate GIST Clinical. Mitotic Rate GIST Pathological will be

Code	Description	AJCC ID
	Blank	All except 43.1, 43.2
1	L: Low (5 or fewer mitoses per 5 mm², or per 50 HPF)	
2	H: High (over 5 mitoses per 5 mm², or per 50 HPF)	
7	Test done, results not in chart	
9	Not documented in medical record	
	Mitotic rate cannot be assessed or unknown if assessed	

Full Data Item Name: Mitotic Rate GIST Pathological Recommended NAACCR Data Item Name: Mitotic Rate GIST Path Data Item Length: 1

Required for AICC 8th Edition Staging: Yes

Source: New data Item, submitted to UDS in 2016 and approved

43 Mitotic Rate Gastrointestinal Tumor Pathological

Note 1: Record the mitotic rate as 1 (Low) or 2 (High) as indicated on the pathology report or CAP

1 (Low) is 5 or fewer mitoses per 5 mm², or per 50 HPF. 2 (High) is over 5 mitoses per 5 mm², or per 50 HPF

Note 3: If only one mitotic rate is available, and is assessed during the AJCC pathological staging timeframe, record this value for Mitotic Rate GIST Pathological. Mitotic Rate GIST Clinical will be coded

Code	Description	AJCC ID
	Blank	All except 43.1, 43.2
1	L: Low (5 or fewer mitoses per 5 mm², or per 50 HPF)	
2	H: High (over 5 mitoses per 5 mm ² , or per 50 HPF)	
7	Test done, results not in chart	
9	Not documented in medical record	
	Mitotic rate cannot be assessed or unknown if assessed	

The "Complete List" of Required SS Fields

	TNM/SS	1	
Schema Name	Required		
GISTPeritoneum	5,10	Penis	17
Testis	4,5,13,15,16	GISTAppendix	11
Melanomalris	4,25	GISTColon	11
Breast	3,4,5	GISTRectum	11
MerkelCellVulva	3,11	Vulva	11
MelanomaCiliaryBody	2,3,4,25	BileDuctsIntraHepat	10
MelanomaChoroid	2,3,4	GISTEsophagus	6
Appendix	2,11	GISTSmallIntestine	6
Scrotum	12,16	GISTStomach	6
Skin	12,16	SkinEyelid	6
Prostate	1,3,8,10	MerkelCellPenis	3
EsophagusGEJunction	1,25	MerkelCellScrotum	3
Nasopharynx	1,25	MerkelCellSkin	3
Peritoneum	1,25	Bladder	2
PharyngealTonsil	1,25	CarcinoidAppendix	2
Stomach	1,25	Colon	2
MelanomaSkin	1,2,3,4,7	CorpusAdenosarcoma	2
MelanomaConjunctiva	1,2	CorpusCarcinoma	2
BileDuctsDistal	25	CorpusSarcoma	2
BileDuctsPerihilar	25	Lymphoma	2
CysticDuct	25	LymphomaOcularAdnexa	2
LacrimalGland	25	NETColon	2
LacrimalSac	25	NETRectum	2
PeritoneumFemaleGen	25	Rectum	2

	(16) Esophagus and Esophagogastric Junction (19) Appendix – Carcinoma (38) Bone (appendicular skeleton, spine, and
AJCC Grede Clinical AJCC Grede Pathologic	pelvis) (40) 5oft Tissue Sarcoms of the Head and Neck (41) 5oft Tissue Sarcoms of the Trunk and Extremitie: (42) 5oft Tissue Sarcoms of the Abdomen and Thorack Viscored Organs (44) 5oft Tissue Sarcoms of the Retroperitoneum (45) 5oft Tissue Sarcoms of Thusual Histologies and Sites.
AJCC GIST Mitotic Count Clinical	,,
AJCC GIST Mitotic Count Pathologic	(43) Gastrointestinal Stromal Tumor
AJCC Testis Serum Markers Clinical AJCC Testis Serum Markers Pathologic	(9) Testis
AJCC Oropharyngeal p16	(10) HPV-Mediated (p16+) Oropharyngeal Cance (11) Oropharynx (p16-) and Hypopharynx
AJCC Esophagus and EGJ Tumor Epicenter	(16) Esophagus and Esophagogastric Junction
AJCC Retinoblastoma Heritable Trait	(68) Retinoblastoma
Revised LVI	All
Revised SSF25 (Lip/vermillion border)	C00.0-upper lip, C00.1-lower lip, C00.2-lip NOS. Cancers of the external lip are staged using AJCC chapter 7. Cancers of the vermillion border are staged using AJCC chapter 15.
Revised 55F25 (cervical node unknown primary)	Occult Head/Neck J patients with EBV-related cervical adenopath are staged according to Chapter 9 (Hasopharnyi, [2) patients with HPV-related cervical adenopathy are staged according to Chapter 10 (HPV-meditated oropharnyaeal cancer [p.15+1); [3] all other patients with EBV-unrelated and HPV-unrelated corrival adenopathy are staged according to Chapter 6
AJCC CLL/SLL Absolute Lymphocyte Count	
AJCC CLL/SLL Adenopathy	
AJCC CLL/SLL Organomegaly	(79) Hodgkin and Non-Hodgkin Lymphomas
AJCC CLL/SLL Anemia	
AJCC CLL/SLL Thrombocytopenia AJCC MM/Plasma Cell Serum B2-microglobulin	
AJCC MM/Plasma Cell Serum albumin AJCC MM/Plasma Cell LDH Level	(82) Multiple Myeloma and Plasma Cell Disorders

Pending SSFs - Recommended for Clinical Care

Lifestyle Factors

- o Tobacco Use
- o Depression
- o Alcohol

Virus Exposures

- o P16/HPV
- o HIV
- o Hep B or Hep C

Overall Health Status

- o Comorbidity(s)
- o Overall Health
- o Performance Status
 - o Zubrod/ECOG
 - o Karnofsky

Other Anatomic Info

- o Location of Positive Lymph Node(s)
- o Size of Positive Node
- o Extranodal Extension
- o Perineural Invasion
- o TumorThickness
- o Depth of Invasion
- o Surgical Margins
- Clinically Relevant Site Specific Prognostic Variables are in the **AJCC Staging Manual**
- New Site Specific Fields not yet created to store these variables
- ALL are Pending Review
- None are Required in 2018
- None are Optional in 2018
- No Instructions or Codes Yet.

16

"Complete List" of Clinically Relevant SSFs

Methylation f 6-Methylguanine-Methyltransferase (MGMT) Brain, SSF #4 osome 1p: Loss of Heterozygosity (LH) Brain, SSF #5 Chromosome 19q: Loss of Heterozygosity (LH) Brain, SSF #6 IDH (Is citrate dehydrogenase) gene Brain, new Lymph Nodes Head and Neck, Levels I-III Head and Neck (Cmmn SSF), SSF#3 Lymph Nodes Head and Neck, Levels IV-V ad and Neck (Cmmn SSF), SSF#4 Lymph Nodes Head and Neck, Levels VI-VII Head and Neck (Cmmn SSF), SSF#5 Lymph Nodes Head and Neck, other Head and Neck (Cmmn SSF), SSF#6 Carcinoembryonic Antigen (CEA) Pretreatment Lab Value Colon and Rectum, SSF #3 Alpha Fetoprotein (AFP) Pretreatment Lab Value Liver, SSF #3 Creatinine Pretreatment Lab Value Liver, SSF #4 International Normalized Ratio Prothrombin Time (INR) Liver, SSF #8 Percent Necrosis Post Neoadjuvant Chemotherapy Bone, SSF #3 Serum Lactate Dehydrogenase (LDH) Pretreatment Lab Value Mela Alpha Fetoprotein (AFP) Pre-orchiectomy Lab Value Testis, SSF#6 Human Chorionic Gonadotropin (hCG) Pre-orchiectomy Lab Value Testis, SSF#8 Alpha Fetoprotein (AFP) Post-orchiectomy Lab Value Testis, SSF# 12 Human Chorionic Gonadotropin (hCG) Post-Orchiectomy Lab Value Testis, SSF# 14 Lymph Nodes, Axillary, Number of Positive Ipsilateral Level I-II Breast, SS# 3 Response to Neoadiuvant Therapy Breast, SS# 21 Multigene Signature Method Breast, SSF# 22 Multigene Signature Results Breast, SSF# 23 HER2 IHC Summary Breast 8th edition, CAP guidelines HER2 ISH Summary Breast 8th edition, CAP guidelines Ki-67 (MIB-1) Breast 8th edition, CAP guidelines HIV Status Lymphoma, SSF#1

Alpha Fetoprotein (AFP) Pretreatment Interpretation Liver, SSF #1 Fibrosis Score (Ishak Score) Liver, SSF #2 Creatinine Pretreatment Unit f Measure Liver, SSF #5 Bilirubin Pretreatment Unit f Measure Liver, SSF #7 Tumor Growth Pattern Intrahenatic Rile Duct. SSE#10 Chromosome 3 Status Uveal Melanomas, SSF #5 chrism 8q Status Uveal Melanomas, SSF #7 Extravascular Matrix Patterns, Lps Uveal Melanomas, SSF #11 Separate Tumor Nodules Lung, SSF#1 Visceral Pleural Invasion Lung, SSF #2 Pleura Effusion Pleura (Malignant Pleural Mesothelioma), SSF#1 Carcinoembryonic Antigen (CEA) Pretreatment Interpretation, Colon and Rectum, SSF #1 Tumor Deposits, Colon and Rectum, SSF #4 Perineural Invasion, Colon and Rectum, SSF #8 Estrogen Receptor, Percent Positive Range, Draft, Breast 8th edition, CAP guidelines Estrogen Receptor, Total Allred Score, Draft, Breast 8th edition, CAP guidelines Progesterone Receptor, Percent Positive Range, Draft, Breast 8th edition, CAP guidelines Progesterone Receptor, Total Allred Score, Draft, Breast 8th edition, CAP guidelines Ki-67 (MIB-1), Draft, Breast 8th edition, CAP guidelines Extranodal Extension: Head and Neck ENE (-) r ENE (+), Clinical, H&N SSF#8 Extranodal Extension: Head and Neck ENE (-) r ENE (+), Pathological, H&N SSF#9 Primary Sclerosing Cholangitis (PSC), Intrahepatic Bile Duct, SSF#11 Breslow tumor thickness, Melanoma Skin, SSF #1 Mitotic Rate Melanoma, Melanoma Skin, SSF #7 Mitotic Cunt Uveal Melanoma, Uveal Melanomas, SSF #9

"Complete List" of Clinically Relevant SSFs

- HER2 ISH Dual Probe Ratio, new Draft, Breast 8th edition, CAP guidelines

- HER2 ISH Dual Probe Ratio, new Dualt, Breast 8th edition, CAP guidelines
 HER2 ISH Dual Probe Copy Number
 HER2 ISH Single Probe Copy Number
 Lymph Nodes Size of Metastasis, Head and Neck (Common. SSF.), SSF#1
 Hillrobin Protreatment Total Lab Value, Liver, SSF #6
 Measured Basal Dismeter, Ureal Melancoms, SSF #2
 Measured Hischest, Verul Melancoms, SSF #3
 Entamodal Extension Chricial, Pernis, SSF #17
 Entamodal Extension Pathological, Pernis, SSF #17
 Entamodal Extension Pathological, Pernis, SSF #17
 Microvascular Density, Ureal Melancoms, SSF #3
 Adensid Cysic Basaloid Pattern, Lacrimed Gland, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, SSF #6
 Circumferential or Radial Resection Magin, Colon and Rectum, CAP guidelines
 Omotype Da Risk Level-Drussive, Dard, Breast 8th edition, CAP guidelines
 Omotype Da Risk Level-Drussive, Dard, Breast 8th edition, CAP guidelines
 Omotype Da Risk Level-Drussive, Dard, Breast 8th edition, CAP guidelines
 Omotype Da Risk Level-Drussive, Dard, Breast 8th edition, CAP guidelines
 Omotype Da Risk Level-Drussive, Medical Cell Skin, SSF #22
 Microsatellite Instability, Colon and Rectum, SSF #7
 KRAS, Colon and Rectum, SSF #9
 Kitchey Tumor Eisension, Kidney, SSF#1
 Major veta Involvement, Kidney, SSF#2
 Ipsilatural Adensat Gland Involvement, Kidney, SSF#3
 Sarceomitoid Features, Kidney, SSF#4
 IAX2, Herne Retric, SSF#1

Manual to Include Complete Site-Specific Field Definitions, Instructions & Codes

Do Not Use Old Manuals for New Site Specific Field Definitions, Instructions, or Codes



2018 - New Manual for ALL New Site Specific Fields – Date TBA

Will Include All Required for Staging Fields

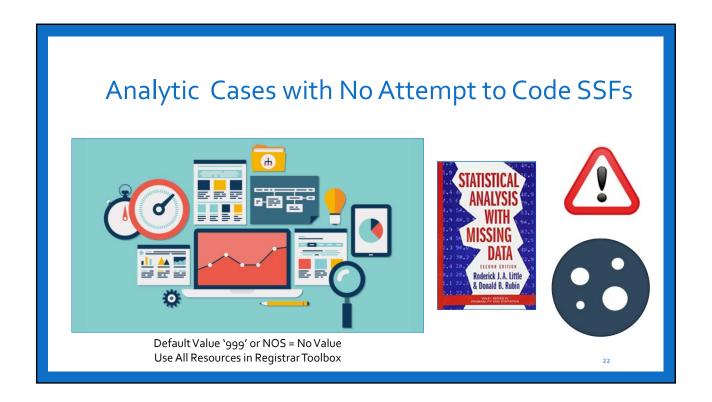
May Include Some Recommended for Clinical Care Fields

Definitions, Instructions, and Codes Included

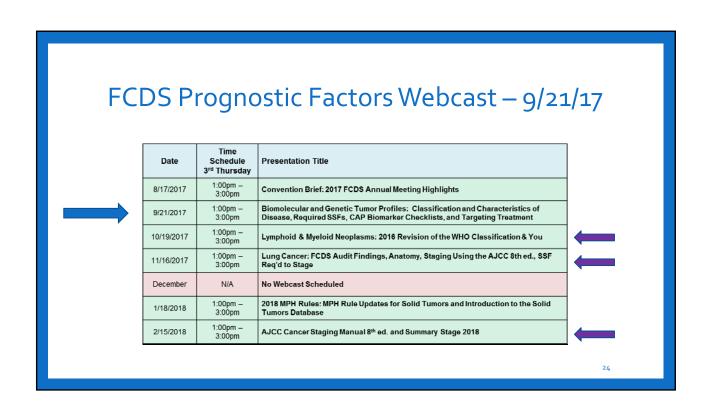
19

Using Required SSFs to Assign Stage Group Table 8. Examples of Revisions to Breast Cancer Staging Using Biomarkers and Oncotype DX T N M G HERZ EN PROSPRING MOSTIC GROUP SAVEVITE EUTION STAGE GROUP STAGE GROUP STAGE GROUP T N M G HERZ EN PROSPRING GROUP T N M G HERZ EN PROSPRING GROUP STAGE GROUP STAGE GROUP T N M G HERZ EN PROSPRING GROUP T N M G HERZ EN PROSPRING GROUP STAGE GROUP STAGE GROUP T N M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ EN PROSPRING GROUP T N M M G HERZ E

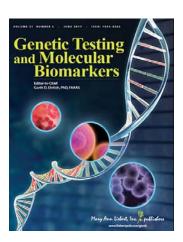








Resources



Genetic Testing and Molecular Biomarkers is the leading peer-reviewed journal covering all aspects of human genetic testing including molecular biomarkers. The Journal provides a forum for the development of new technology; the application of testing to decision making in an increasingly varied set of clinical situations; ethical, legal, social, and economic aspects of genetic testing; and issues concerning effective genetic counseling. This is the definitive resource for researchers, clinicians, and scientists who develop, perform, and interpret genetic tests and their results.

Genetic Testing and Molecular Biomarkers coverage includes:

- o Diagnosis across the life span
- o Risk assessment
- o Carrier detection in individuals, couples, and populations
- o Novel methods and new instrumentation for genetic testing
- o Results of molecular, biochemical, and cytogenetic testing
- o Genetic counseling

25

Questions

