

2024 Updates: Site Specific Data Items Grade

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IOWA CANCER REGISTRY

DECEMBER 2024

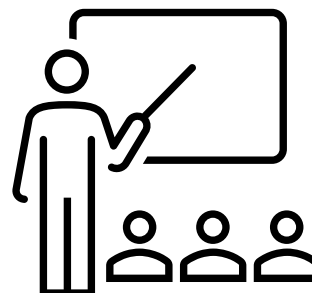


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Overview

2024 Major Updates for:

- Site Specific Data Items (SSDI)
- Grade



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Location of Updates

NAACCR Website

- Central Registry Standards
- Site Specific Data Items (SSDI)/Grade

https://apps.naaccr.org/ssdi/list/?_gl=1*1nrce0i*_ga*NDk5ODY0NzI3LjE3MzE5NDIwMTA.*_ga_V7J8GWYK5P*MTczMjlyMTIzNC4xMy4wLjE3MzlyMjEyMzQuNjAuMC4w

RESOURCES

Version 3.1 For use with cases diagnosed 2018 forward after registry software conversion to the NAACCR Data Standards and Data Dictionary, Version 24)

- » [SSDI Manual](#)
- » [SSDI Manual Appendix A](#)
- » [SSDI Manual Appendix B](#)
- » [SSDI Manual Appendix C](#)
- » [Grade Manual](#)
- » [Change Log](#) ←

Comments or suggestions concerning the SSDI's are welcome and can be posted at the American College of Surgeons [CAnswer Forum](#).



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2024 UPDATES

Site Specific Data Items (SSDI)



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Site Specific Data Items (SSDI)

Reminders

- It is critical to accurately assign primary site and histology
 - Behavior as well for specific sites
- The above information, along with Schema Discriminator when applicable, defines the following:
 - Schema ID
 - AJCC ID
 - EOD Schema
 - Summary Stage chapter
 - SSDIs
 - Grade
 - Surgery codes



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New Schemas for 2024+

No changes to SSDIs or Grade for these:

- NET Stomach 09290
- NET Duodenum 09301
- NET Ampulla of Vater 09302
- NET Jejunum and Ileum 09310
- NET Appendix 09320
- NET Colon and Rectum 09330
- NET Pancreas 09340

SSDI – p16 required for 2024+

- Vulva 09500
 - Currently collected for Cervix (2021+) and Anus (2023+)



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New SSDI

Brain Primary Tumor Location (Brain 2023+ Schema) – Cases 2024+

- This SSDI distinguishes between the Pons and other subsites with Primary Site code C717
- Information regarding Pons primary site is very important, especially for pediatric brain tumors
- Dangerous site to biopsy or have surgical resection, most of the information will come from imaging or physician's statement

Description	Code
Pons	1
Subsite other than Pons; Multiple subsites in Brain stem listed	2
Not applicable (based on standard setter)	8
Brain Stem, NOS; Unknown subsite of Brain Stem	9
Primary site is NOT C717; Diagnosis year prior to 2024	Blank



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Brain Stem Anatomy

Midbrain

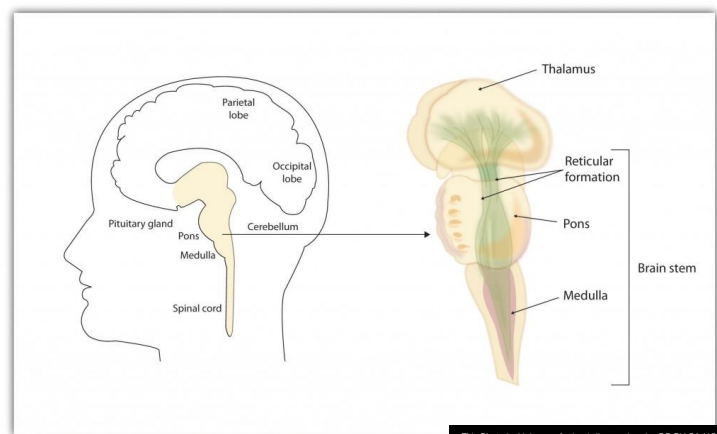
- Most superior portion and acts as a conduit between the forebrain above and pons below

Pons

- Largest portion and located above the medulla and below midbrain
- Group of nerves that function as a connection between the cerebrum and the cerebellum

Medulla Oblongata

- Most inferior portion and houses ascending and descending nerve tracts as well as brain stem nuclei



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New Codes Added

Brain Molecular Markers (2024+)

- New histologies added:
 - **9385/3, 9396/3, 9421/1, 9430/3, 9500/3**
 - **Codes 10-23** can **ONLY** be used **1/1/2024+**
 - **Code 85** (NA due to histology) updated based on new histology codes
 - Histology is **not**: 9385/3, 9396/3, 9400/3, 9401/3, 9430/3, 9440/3, 9450/3, 9451/3, 9471/3, 9478/3, 9421/1, 9430/3, 9500/3
 - **Code 86** – benign or borderline tumor updated
 - *Excludes 9421/1* (newly included histology)
- Based on CNS WHO Blue Book updates, released 2022



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Brain Molecular Markers 2024+

Histology	Description	Code
9385/3 (Pediatric-type diffuse low grade gliomas)	Diffuse hemispheric glioma, H3-G34 mutant	10
	Diffuse midline glioma, H3 K27-altered	11
	Diffuse pediatric-type high-grade glioma, H3-wildtype, and IDH-wildtype	12
	Infant-type hemispheric glioma	13
9396/3 (Ependymal Tumors)	Posterior fossa group A (PFA) ependymoma	14
	Posterior fossa group B (PFB) ependymoma	15
	Spinal ependymoma, MYCN-amplified	16
	Supratentorial ependymoma, YAP 1 fusion-positive	17
	Supratentorial ependymoma, ZFTA fusion-positive	18



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Brain Molecular Markers 2024+

Histology	Description	Code
9421/1 (Pilocytic Astrocytoma/Juvenile Pilocytic Astrocytoma)	Diffuse astrocytoma, MYB- or MYBL 1-altered	19
	Diffuse low grade glioma, MAPK pathway-altered	20
9430/3 (Circumscribed Astrocytic Tumor)	Astroblastoma, MN1-altered	21
9500/3 (Other CNS Embryonal Tumor)	CNS Neuroblastoma, FOXR2-activated	22
	CNS Tumor with BCOR internal tandem duplication	23



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SSDI –
Updated
Notes/Codes



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Updated Notes and Codes

Response to Neoadjuvant Therapy (3922) Breast [schema: 00480]

- **Updated Note 2**
 - This data item should not be coded based on the pathological, radiological, or imaging findings. **This data item should only be coded based on the managing physician's overall interpretation of the results.**
 - Do not confuse these instructions with the SEER Data Item: Neoadjuvant Therapy-Treatment Effect (#1634)
 - For [SEER's data item](#) the field is based on the *pathologic response only* after neoadjuvant therapy.
- Added to **code 9**:
 - Unknown if neoadjuvant therapy done
- *Remember:*
 - If neoadjuvant therapy is not done, that is not going to be documented in the medical record
 - Based on sequence of events (diagnosis, treatment), you can usually determine if neoadjuvant therapy was done or not.



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Updated Notes and Codes

Circumferential Resection Margin (3823) Colon and Rectum [schema: 00200]

- **Note 3 updated:**
 - For **colon primaries**, surgery of primary site **must be a surgical resection**
 - No surgical resection of primary site (i.e. polypectomy), then **code XX.7**
 - For **rectum primaries**, surgery of primary site must be coded as **excisional biopsy, transanal excision, or surgical resection**
 - For excisional biopsy, transanal procedures, *only the peritonealized portion of the rectum* (upper 1/3) is where you can get the CRM
 - Non-peritonealized portion (below the peritoneal reflection) of rectum is involved or it's unknown if peritonealized portion is involved **code XX.7**
 - If surgery of primary site is not excisional biopsy/transanal excision/surgical resection, **code XX.7**



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Updated Notes and Codes

p16 (3956) Anus [09210] Cervix [09520] Vulva [09500]

- **Note 3 updated:**
 - This data item must be based on testing results for p16 overexpression.
 - Testing for HPV by DNA, mRNA, or other methods **should not be coded** in this data item.
 - **Do not confuse p16 with HPV**, which is a specific strain of virus. A statement of a patient being HPV positive or negative is not enough to code this data item.



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Schemas: 00528, 00530, 00541, 00542 Cervix Sarcoma and Corpus

SSDI	Note/Code	Update
3899: Number of Examined Para-aortic Nodes *not routinely examined*	Note 4	Code 00 when <u>no LN</u> are removed (including FNA, core biopsy, SLN Bx, and LND) <ul style="list-style-type: none"> • If LN are removed and only pelvic LN are assessed, or only “nodes” are documented – code 00
	Code X9	Removed “No lymph node dissection performed”
3900: Number of Examined Pelvic Nodes	Note 4	If LN are removed and only “nodes” are documented without specifying pelvic or para-aortic, assume they are pelvic
	Code X9	Removed “No lymph node dissection performed”



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Schemas: 00528, 00530, 00541, 00542 Cervix Sarcoma and Corpus

SSDI	Note/Code	Update
3901: Number of Positive Para-aortic Nodes 3902: Number of Positive Pelvic Nodes	Note 6	Code X9 if no LN removed: <ul style="list-style-type: none"> • Only a <u>FNA or core biopsy</u> is done and positive, code X6 • Only a <u>FNA or core biopsy</u> is done and negative, code X9
	Code X9	<i>Added:</i> <ul style="list-style-type: none"> • No lymph nodes removed <i>Removed:</i> <ul style="list-style-type: none"> • No lymph node dissection performed



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Updated Notes and Codes

Invasion Beyond Capsule (3864) Kidney Parenchyma [00600]

- **Note 3:** Perinephric/sinus fat invasion should be confirmed microscopically
 - Synonyms include renal sinus fat, medial invasion
 - Do **not** code invasion of renal hilum
 - If all you have is renal hilum invasion code to 9 (unknown)



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2024 UPDATES

Grade



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Derived Summary Grade

Derived in central registry *ONLY*

- Applied to cases for 2018+
- Based on information already in the cancer registry
- Data items used to derive Summary Grade:
 - Grade Clinical
 - Grade Pathological



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Ranges in Grade

New guideline included in “General Grade Coding Instructions for Solid Tumors”

1. Code the grade from the primary tumor only
 - a. Do NOT code grade from metastatic tumor or recurrence.
 - In the rare instance that tumor tissue extends contiguously to an adjacent site and tissue from the primary site is not available, code grade from the contiguous site
 - b. If primary site is unknown, code grade to 9
 - c. If a range is given for a grade (e.g. 1-2 or 2-3), code the higher grade



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Autopsy Grading

New guideline “General Instruction for the Time Frames for Grade”

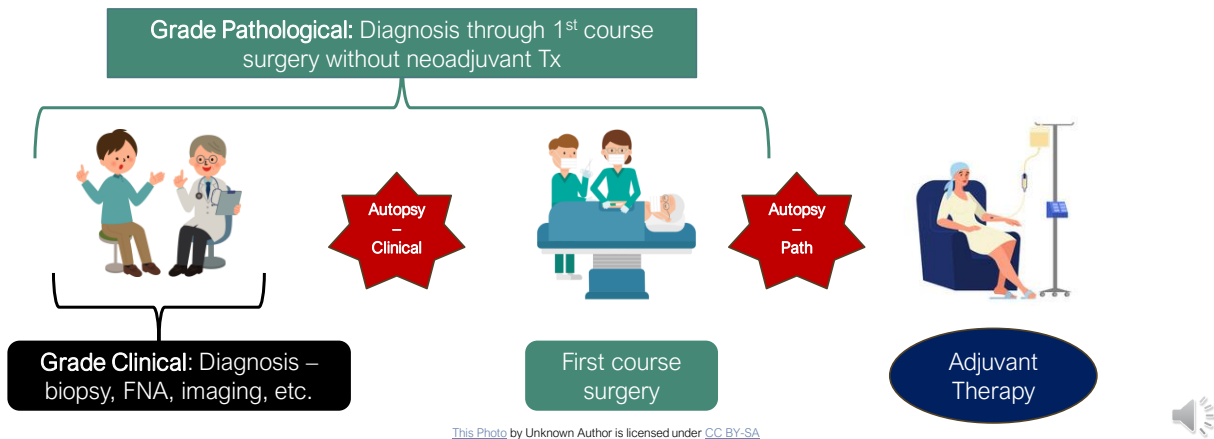
- If patient dies and has an **autopsy performed within the initial work up and treatment of their cancer** (without signs of progression), the grade from the autopsy can be used
- Include autopsy information from the appropriate timeframe: clinical, pathological, post therapy clin, post therapy path

Note: Do not automatically assign pathological grade based on the autopsy



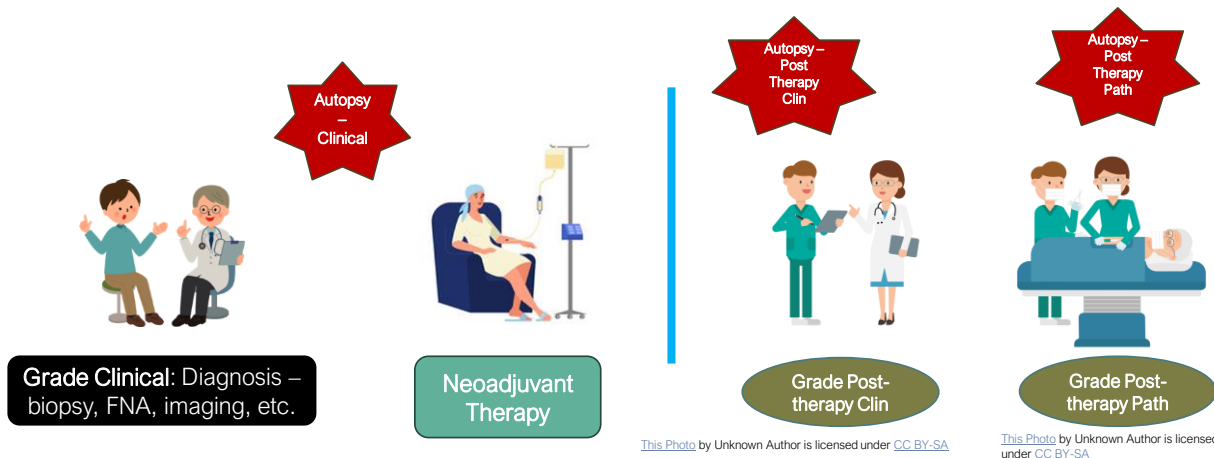
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Grade Coding Timeframe



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Post-Therapy Grade Coding Timeframe



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Autopsy Grade Coding

Example 1

Patient is diagnosed at autopsy.

Autopsy information can't be used to code grade

Example 2

Patient diagnosed prior to death/autopsy.
No treatment administered prior to death/autopsy.

Code Grade Clinical from autopsy findings



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Autopsy Grade Coding

Example 3

Patient diagnosed prior to death/autopsy.
Patient has surgical resection and dies soon afterwards. Autopsy performed.

Code Grade Pathological from surgery or autopsy, whichever is higher

Example 4

Patient diagnosed, underwent surgical resection of primary, and subsequently completed all first course treatment prior to death/autopsy. Patient has autopsy.

Autopsy grade can't be used as it was performed following diagnosis and first course treatment



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Grade Coding

Corpus Carcinoma and Carcinosarcoma – Table 13

- **Note 3:** For endometrioid carcinomas **only**
 - If “low grade” is documented **code 2** (FIGO Grade 2)
 - If “high grade” is documented **code 3** (FIGO Grade 3)



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Grade Coding – Table 07

Do **NOT** code grade based on the histology terms below:

- Neuroendocrine carcinoma, low grade (8240/3)
- Neuroendocrine carcinoma, well differentiated (8240/3)
- Neuroendocrine carcinoma, moderately differentiated (8249/3)
- Poorly differentiated neuroendocrine carcinoma (8246/3)
- Ki-67 and Mitotic rate are the preferred grading method
- See CAnswer Forum: <https://cancerbulletin.facs.org/forums/node/124336>
- Updated information on the 1/11/24 post



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Grade Coding – Table 07

Neuroendocrine carcinoma, Grade 1 (8240/3)

- Ki-67 less than 3 **AND/OR**
- Mitotic count less than 2

Note: many times, if Ki-67 is less than 3, mitotic count is not done

Neuroendocrine carcinoma, Grade 2 (8249/3)

- Ki-67 3-20 **OR**
- Mitotic count 2-20

Neuroendocrine carcinoma, Grade 3 (8249/3)

- Ki-67 >20 **OR**
- Mitotic count >20

If pathologist gives a stated specific grade and there is no Ki-67 or mitotic rate, then code the stated grade



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Definition	Code
G1: Mitotic less 2 AND Ki-67 less 3	1
G2: Mitotic = 2-20 AND Ki-67 = 3-20	2
G3: Mitotic greater 20 AND Ki-67 greater 20	3
Well Diff	A
Moderately Diff	B
Poorly Diff	C
Undifferentiated; Anaplastic	D
Unknown	9

Grade Coding – Table 07 Example

Neuroendocrine Tumor (NET)

- Appendix: well-differentiated neuroendocrine tumor
 - Histology = 8240/3
 - No Ki-67 or mitotic information
 - What is the correct grade?



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Grade Coding – Table 07 Examples

Well differentiated NET, WHO Grade 2, mitotic rate 1 per 2mm² and Ki-67 5%

Grade: 2

Neuroendocrine tumor, Grade 1-2, mitotic rate less than 2 per 2mm² and Ki-67 3%

Grade 2

Duodenum: Well differentiated NET (incidental); Ki67 labeling index: less than 3%

Grade: 1



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Grade Coding – Table 07 Examples

Neuroendocrine carcinoma, well differentiated; Ki-67 2%; no mitotic count information

Grade: 1

Neuroendocrine carcinoma, moderately differentiated; no Ki-67 or mitotic rate information

Grade 9

Poorly differentiated neuroendocrine carcinoma; Mitotic rate: 5 per 2mm²; Ki-67: 15%

Grade: 2



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Questions?

Submit questions to CAnswer Forum:

Site Specific Data Items (SSDI)/Grade 2018

<https://cancerbulletin.facs.org/forums/CancerForumHome>

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