Introduction

**Note 1:** This section includes the following **primary sites:** Peripheral nerves C470-C479; cerebral meninges C700; spinal meninges C701; meninges NOS C709; brain C710-C719; spinal cord C720; cauda equina C721; olfactory nerve C722; optic nerve C723; acoustic nerve C724; cranial nerve NOS C725; overlapping lesion of brain and central nervous system C728; nervous system NOS C729; pituitary gland C751; cranioopharyngeal duct C752; pineal gland C753.

**Note 2:** Non-malignant intracranial and CNS tumors have a separate set of rules.

**Note 3:** 2007 MPH Rules and 2018 Solid Tumor Rules are used based on **date of diagnosis.**
- Tumors diagnosed 01/01/2007 through 12/31/2017: Use 2007 MPH Rules
- Tumors diagnosed 01/01/2018 and later: Use 2018 Solid Tumor Rules
- The original tumor diagnosed before 1/1/2018 and a subsequent tumor diagnosed 1/1/2018 or later in **the same primary site:** Use the 2018 Solid Tumor Rules.

**Note 4:** There must be a histologic, cytologic, radiographic, or clinical **diagnosis** of a **malignant** neoplasm /3.

**Note 5:** Tumors from a number of primary sites metastasize to the brain. Do not use these rules for tumors described as metastases; report metastatic tumors using the rules for that primary site.

**Note 6:** **Pilocytic astrocytoma/juvenile pilocytic astrocytoma** is reportable in North America as a **malignant** neoplasm 9421/3.

**Note 7:** Tables and rules refer to ICD-O rather than ICD-O-3. The specific version is not specified to allow for updates. Use the currently approved version of ICD-O and all updates.

**Note 8:** For those sites/histologies which have recognized **biomarkers,** the biomarkers may aid in the identification of histologic type. Currently, there are clinical trials being conducted to determine whether these biomarkers can be used to identify multiple primaries. Follow the Multiple Primary Rules; do not code multiple primaries based on biomarkers.

**Note 9:** See the Head and Neck Rules for coding paragangliomas.
Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

These terms can be used interchangeably:

- And; with
  Note: “And” and “with” are used as synonyms when describing multiple histologies within a single tumor.
- Cerebrospinal fluid; CSF
- Dura; meninges
- Extradural; not within the meninges; within the cranium; within the skull but not within the cerebral meninges
- Extramedullary; outside medulla oblongata C700
- Infratentorial; below the tentorium cerebelli; either cerebellum or brainstem
- Intracranial; within the skull, within the cranium
- Intradural; between layers of the cerebral meninges; C700
- Intradural-extradural; within the spinal canal but outside of the nerves
- Intraspinal; occurring within the spinal column especially the vertebral canal; spinal nerve roots
- Site; topography
- Supratentorial; above the tentorium cerebelli; cerebrum
  - Cerebrum contains frontal lobe, parietal lobe, occipital lobe, and temporal lobe
- Tentorium cerebelli; cerebellar tentorium
- Tumor; mass; lesion; neoplasm when
  - These terms are used ONLY to determine multiple primaries
  - Do not use these terms for casefinding or determining reportability
- Type; subtype; variant
- WHO Grade 3 and WHO Grade 4; malignant; invasive; /3
Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions
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Terms that are NOT Equivalent or Equal

This is a term that is not equivalent. There are no casefinding implications.

- **Component** is not equivalent to **subtype/type/variant**
  
  **Note:** Component is only coded when the pathologist specifies the component as a second carcinoma.

- **Phenotype** is not equivalent to **subtype/type/variant**

- **WHO Grade** is not equivalent to **tumor grade**

Changes from 2007 MPH Rules

These changes are effective with cases diagnosed 1/1/2018 and later.

1. 2016 CNS WHO presents major restructuring of the diffuse gliomas, medulloblastomas and other embryonal tumors, and incorporates new entities that are defined by both histology and molecular features, including glioblastoma, IDH-wildtype and glioblastoma, IDH-mutant; diffuse midline glioma, H3 K27M-mutant, RELA fusion-positive ependymoma, medulloblastoma, WNT-activated and medulloblastoma, SHH-activated, and embryonal tumor with multilayered rosettes, C19MC-altered. The 2016 edition has added newly recognized neoplasms and has referred to some entities, variants and patterns as “not recommended” (previously called obsolete).
   
   A. It has been determined that these “not recommended” terms no longer have diagnostic and/or biological relevance. For example, gliomatosis cerebri is a term which is no longer recommended. Gliomatosis cerebri is now termed a “growth pattern” rather than a histologic type.
   
   B. Terms which are not recommended are not included in the tables. When one of these terms are used, refer to the ICD-O and all updates for the correct histology code. For example, glioma NOS is an umbrella term for all gliomas and astrocytomas. Glioma NOS is not recommended because diagnostic methodology is able to determine a more specific diagnosis.

2. **Rule change:** The 2007 rules said a glioblastoma multiforme (GBM) following an astrocytic or glial tumor was a single primary (recurrence). In the 2018 Solid Tumor Rules, GBM subsequent to an astrocytic or glial tumor is a multiple primary. GBM is now being collected as a new primary so it is possible to analyze the frequency with which these tumors recur in a more aggressive form (GBM).
3. **Clarifications:**
   A. The following meningiomas are reportable: intraosseous, cavernous sinus and sphenoid wing.
   B. Multiple cerebral meningiomas are a single primary.
   C. Multiple brain tumors (same histology) are a single primary.
   D. Laterality is not used to determine multiple primaries.
   E. Timing is not used to determine multiple primaries.
   F. The brain (C710-C719) is a single primary site.
   G. Neurofibromatosis NOS, Neurofibromatosis 1 (NF1), Neurofibromatosis 2 (NF2), and schwannomatosis are genetic syndromes and not reportable neoplasms. People with this genetic syndrome do have a high risk of developing:
      i. Non-reportable non-malignant tumors occurring in skin and sites other than CNS AND
      ii. Reportable malignant tumors

4. The 2016 World Health Organization (WHO) Classification of Tumors of the Central Nervous System (CNS) contains newly recognized histologies and subtypes/variants. New codes/terms are identified by asterisks (*) in Table 3 in the Terms and Definitions.

---

**Reportability Criteria**

CNS neoplasms must meet all three of the conditions below to be reported as malignant:

1. The **behavior** must be malignant:
   A. Pathology designates the behavior as malignant/invasive, /3 OR
   B. The tumor is a WHO Grade 3 or 4 (See Section 1: Table 1)

   **Note 1:** WHO Grade 2 tumors may be non-malignant or malignant.
   **Note 2:** Always code the behavior as designated by the pathologist.
   **Note 3:** Never report a malignant behavior code for a meningioma based on tumor extension to brain, skin of scalp, or other regional organs/tissue. Non-malignant CNS tumors can extend to the regional tissue and bone.

2. The **primary site** must be reportable (See Section 2: Table 2) AND

3. The **histology** must be reportable (See Section 2: Table 3)
Information is presented in the general order in which a case is abstracted.

Section 1: Behavior Code
   A. Priority Order for Using Documentation to Assign Behavior
   B. Table 1: WHO Grades for Select CNS Neoplasms

Section 2: Reportable Primary Sites and Histologies
   A. Priorities for Coding Primary Site
   B. Reportable Primary Site Groups
   C. Table 2: Reportable Primary Sites
   D. Table 3: Specific Histologies, NOS, and Subtypes/Variants
   E. Table 4: Coding Primary Site for Malignant Tumors of Cranial and Peripheral Nerves

Section 3: Additional information to complete the abstract
   A. Conflicting information on Pathology report(s)
   B. Table 5: Paired Sites
   C. Table 6: Non-malignant CNS Tumors with the Potential of Transforming to Malignant Behavior
Section 1: Behavior Code

Note: Behavior determines which set of CNS rules should be used: malignant or non-malignant.

Instructions for using source documentation to determine behavior are in priority order. Start with Instruction 1 and STOP when you reach the instruction that applies to the case being abstracted. For example, when the resection pathology specifies behavior, use Instruction 1. Do not code the behavior using Instructions 2-5.

Priority Order for Using Documentation to Assign Behavior

1. Pathology: Tissue from resection
   A. Use the pathologist’s description of malignant/invasive behavior
   B. Cases are reportable as malignant when pathology states a WHO Grade 3 or 4
      i. WHO Grade 2 may be either non-malignant or malignant. Use the pathologist’s description of behavior (priority #1).
   C. Never change behavior described by pathologist

2. Pathology: Tissue from biopsy

3. Cytology (usually cerebrospinal fluid)

4. Physician’s documentation (no pathology report) in the following priority order:
   A. Tumor Board
   B. Documentation of original pathologic diagnosis and behavior
      Example: Pathology report is not in the medical record. Oncology consult states biopsy was done in a different facility and cites the original pathology diagnosis including the behavior.
   C. Documentation of behavior, no mention of original diagnosis
      Example: Pathology report is not in the medical record. Physician documents the behavior as malignant, or WHO Grade 3 or 4, but does not cite/mention original pathology report as source of behavior classification.

5. Scan, use behavior information from radiography in the following priority order:
   A. MRI
   B. CT
   C. PET
   D. Angiogram
6. When instructions 1-5 do not apply, use Table 1 to determine behavior.

### Table 1: WHO Grades for Select CNS Neoplasms

<table>
<thead>
<tr>
<th>WHO Grade</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>WHO Grade I</td>
<td>Circumscribed tumors of low proliferative potential associated with the possibility of cure following resection</td>
</tr>
<tr>
<td>WHO Grade II</td>
<td>Infiltrative tumors with low proliferative potential with increased risk of recurrence</td>
</tr>
<tr>
<td>WHO Grade III</td>
<td>Tumors with histologic evidence of malignancy, including nuclear atypia and mitotic activity, associated with an aggressive clinical course</td>
</tr>
<tr>
<td>WHO Grade IV</td>
<td>Tumors that are cytologically malignant, mitotically active, and associated with rapid clinical progression and potential for dissemination</td>
</tr>
</tbody>
</table>

#### Note 1:  
CNS WHO classifications use a grading scheme that is a "malignancy scale" ranging across a wide variety of neoplasms rather than a strict histologic grading system that can be applied equally to all tumor types.

#### Note 2:  
See the SEER and COC Manuals for instructions on coding grade for CNS tumors.

#### Note 3:  
The table does not contain all neoplasms that may occur in the CNS.

#### Note 4:  
WHO does not provide Grades for all CNS and peripheral nerve neoplasms.

#### WHO Grade Definitions

1. Use the malignant CNS rules for all WHO Grade 3 and 4.
2. Go to Section 1: Behavior Code to determine whether WHO Grade 2 neoplasms are non-malignant or malignant.
3. Use non-malignant CNS rules for all WHO Grade 1 (always non-malignant).

**Column 1** contains the histology term.  
**Column 2** contains the WHO Grade assigned based on the molecular features of the histology.
# Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

## Histology

<table>
<thead>
<tr>
<th>Histology</th>
<th>WHO Grade</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anaplastic (malignant) meningioma</td>
<td>3</td>
</tr>
<tr>
<td>Anaplastic astrocytoma, IDH-mutant</td>
<td>3</td>
</tr>
<tr>
<td>Anaplastic ependymoma</td>
<td>3</td>
</tr>
<tr>
<td>Anaplastic ganglioglioma</td>
<td>3</td>
</tr>
<tr>
<td>Anaplastic oligodendroglioma IDH-mutant and 1p/19q-codeleted</td>
<td>3</td>
</tr>
<tr>
<td>Anaplastic pleomorphic xanthroastrocytoma</td>
<td>3</td>
</tr>
<tr>
<td>Angiocentric glioma</td>
<td>1</td>
</tr>
<tr>
<td>Atypical choroid plexus papilloma</td>
<td>2</td>
</tr>
<tr>
<td>Atypical meningioma</td>
<td>2</td>
</tr>
<tr>
<td>Atypical teratoid/rhabdoid tumor</td>
<td>4</td>
</tr>
<tr>
<td>Central neurocytoma</td>
<td>2</td>
</tr>
<tr>
<td>Cerebellar liponeurocytoma</td>
<td>2</td>
</tr>
<tr>
<td>Chordoid glioma of third ventricle</td>
<td>2</td>
</tr>
<tr>
<td>Choroid plexus carcinoma</td>
<td>3</td>
</tr>
<tr>
<td>Choroid plexus papilloma</td>
<td>1</td>
</tr>
<tr>
<td>CNS embryonal tumor NOS</td>
<td>4</td>
</tr>
<tr>
<td>CNS embryonal tumor with rhabdoid features</td>
<td>4</td>
</tr>
<tr>
<td>Craniopharyngioma</td>
<td>1</td>
</tr>
<tr>
<td>Desmoplastic infantile astrocytoma and ganglioglioma</td>
<td>1</td>
</tr>
<tr>
<td>Diffuse astrocytoma, IDH-mutant</td>
<td>2</td>
</tr>
<tr>
<td>Diffuse midline glioma, H3K27M-mutant</td>
<td>4</td>
</tr>
<tr>
<td>Dysembryoplastic neuroepithelial tumor</td>
<td>1</td>
</tr>
<tr>
<td>Dysplastic gangliocytoma of cerebellum (Lhemitte-Duclos)</td>
<td>1</td>
</tr>
<tr>
<td>Ependymoma</td>
<td>2</td>
</tr>
</tbody>
</table>

Jump to [Multiple Primary Rules](#)
Jump to [Histology Coding Rules](#)
### Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

**C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753**  
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Histology</th>
<th>WHO Grade</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ependymoma, RELA fusion-positive</td>
<td>2 or 3</td>
</tr>
<tr>
<td><strong>Note:</strong> Tissue/pathology reports or CAP protocol/summary will specify whether this is WHO Grade 2 or 3</td>
<td></td>
</tr>
<tr>
<td>Extraventricular neurocytoma</td>
<td>2</td>
</tr>
<tr>
<td>Gangliocytoma</td>
<td>1</td>
</tr>
<tr>
<td>Glioblastoma, IDH-mutant</td>
<td>4</td>
</tr>
<tr>
<td>Glioblastoma, IDH-wildtype</td>
<td>4</td>
</tr>
<tr>
<td>Granular cell tumor</td>
<td>1</td>
</tr>
<tr>
<td>Hemangioblastoma</td>
<td>1</td>
</tr>
<tr>
<td>Malignant peripheral nerve sheath tumor (MPNST)</td>
<td>2, 3, or 4</td>
</tr>
<tr>
<td><strong>Note:</strong> Tissue/pathology reports or CAP protocol/summary will specify whether this is WHO Grade 2, 3, or 4</td>
<td></td>
</tr>
<tr>
<td>Medulloblastoma (including all subtypes)</td>
<td>4</td>
</tr>
<tr>
<td>Medulloepithelioma</td>
<td>4</td>
</tr>
<tr>
<td>Meningioma</td>
<td>1</td>
</tr>
<tr>
<td>Myxopapillary ependymoma</td>
<td>1</td>
</tr>
<tr>
<td>Neurofibroma</td>
<td>1</td>
</tr>
<tr>
<td>Oligodendroglioma IDH-mutant and 1p/19q deleted</td>
<td>2</td>
</tr>
<tr>
<td>Papillary gliioneuronal tumor</td>
<td>1</td>
</tr>
<tr>
<td>Papillary tumor of the pineal region</td>
<td>2 or 3</td>
</tr>
<tr>
<td><strong>Note:</strong> Tissue/pathology reports or CAP protocol/summary will specify whether it is WHO Grade 2 or 3</td>
<td></td>
</tr>
<tr>
<td>Perineuroma</td>
<td>1</td>
</tr>
<tr>
<td>Pilocytic astrocytoma</td>
<td>1</td>
</tr>
<tr>
<td><strong>Note:</strong> Collected as malignant /3 in North America</td>
<td></td>
</tr>
</tbody>
</table>
## Histology

<table>
<thead>
<tr>
<th>Histology</th>
<th>WHO Grade</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pineal parenchymal tumor of intermediate differentiation</td>
<td>2 or 3</td>
</tr>
<tr>
<td><strong>Note:</strong> Tissue/pathology reports or CAP protocol/summary will specify WHO Grade 2 or 3</td>
<td></td>
</tr>
<tr>
<td>Pineoblastoma</td>
<td>4</td>
</tr>
<tr>
<td>Pineocytoma</td>
<td>1</td>
</tr>
<tr>
<td>Pituicytoma</td>
<td>1</td>
</tr>
<tr>
<td>Pleomorphic xanthroastrocytoma</td>
<td>2</td>
</tr>
<tr>
<td>Rosette-forming glioneuronal tumor</td>
<td>1</td>
</tr>
<tr>
<td>Schwannoma</td>
<td>1</td>
</tr>
<tr>
<td>Solitary fibrous tumor/hemangiopericytoma</td>
<td>1, 2, or 3</td>
</tr>
<tr>
<td><strong>Note:</strong> Tissue/pathology reports or CAP protocol/summary will specify WHO Grade 1, 2, or 3</td>
<td></td>
</tr>
<tr>
<td>Spindle cell oncocytoma</td>
<td>1</td>
</tr>
<tr>
<td>Subependymal giant cell astrocytoma</td>
<td>1</td>
</tr>
<tr>
<td>Subependymoma</td>
<td>1</td>
</tr>
</tbody>
</table>
After determining behavior code, the next step is to confirm both the primary site and the histology are reportable.

**Note 1:** **Peripheral nerves** are included in the Malignant CNS and Peripheral Nerve rules because:
- **All malignant** tumors are reportable, including peripheral nerve tumors **AND**
- The Malignant CNS and Peripheral Nerve **rules** contain the **correct histologies** and coding **rules** for tumors of **peripheral nerves and meninges/dura.**

**Note 2:** **Peripheral nerves** are:
- **Extracranial**/outside the cranium **OR**
- **Extradural**/outside the spinal dura

**Note 3:** The following malignant **meningiomas** are reportable:
- **Intraosseous**
  **Note:** The **dura** layer of the meninges **contacts** the **endosteum** of the bones of the skull. The primary site for intraosseous meningioma is cranial meninges C700.
- **Sphenoid wing**
  **Note 1:** Sphenoid wing meningiomas arise in the **cranial meninges** C700 which covers the bony structure called the sphenoid wing.
  **Note 2:** The term “sphenoid wing meningioma” is used to identify the **location** of the meningioma because sphenoid wing meningiomas may be very **invasive**, spreading to the dura of the frontal, temporal and orbital regions.
- **Cavernous sinus**
  **Note 1:** Cavernous sinus is located between the endosteal and meningeal layers of the dura.
  **Note 2:** There is no ICD-O site code for cavernous sinus because tumors do not arise in the sinus itself; the primary sites are:
  - The **cranial nerves** passing through the sinus (trochlear, abducent C725) **OR**
  - The cerebral **meninges/dura** C700 covering the cranial nerve
Priorities for Coding Primary Site

Note 1: **Always** check the operative report(s) which will have information on whether the surgery or biopsy was **intracranial** (inside the cranium/skull) or **intraspinal** (within the dura/meninges covering the spinal cord).

Note 2: Code the specific primary site. Use an NOS site code only when a specific site is not known.

Use the list in hierarchical order:

1. **Resection**
   A. Operative report(s)
   B. Pathology report(s)

2. **Biopsy**
   A. Operative report(s)
   B. Pathology report(s)

3. Resection and/or biopsy performed, but operative report(s) and pathology are not available (minimal information)
   A. Tumor Board
   B. Code from physician’s documentation of original diagnosis from operative or pathology report OR
   C. Physician’s documentation of primary site in the medical record

   **Example:** The patient had a biopsy done at another facility. The operative and pathology reports are not in the medical record. The attending documents that the patient had a biopsy of a cerebral meningioma at a different facility. Code the primary site cerebral meninges C700.

4. For cases diagnosed by imaging (no pathology/resection or biopsy) use information from scans in the following priority order:
   A. MRI
   B. CT
   C. PET
   D. Angiogram

5. See **Table 2: Reportable Primary Sites** to confirm the primary site is reportable.

6. When the primary site is cranial nerve OR peripheral nerve, see **Table 4: Coding Primary Site for Malignant Tumors of Cranial and Peripheral Nerves** to determine whether the portion of the nerve is cranial or peripheral (different site codes).
The three major groups of reportable sites are:

1. **Intracranial** (within the skull/cranium) AND
2. **Spinal sites** (spinal meninges and sites within the spinal meninges)
3. **Peripheral nerves** (extracranial or extraspinal nerves).

Terms and sites for these broad categories are:

**Reportable Primary Sites and their ICD-O Codes**

1. Intracranial (intra means inside or within; intracranial is inside the skull).
   A. **Cerebral/cranial dura/meninges** C700. The cerebral meninges has three layers:
      i. **Dura** mater is the **superficial** layer of meninges
         - Tightly adherent to skull
         - Contains folds and **sinuses**
         - Contacts **endosteum** which lines the bones of the skull
      ii. **Arachnoid** mater forms the middle of the three layers of meninges
      iii. **Pia** mater is the delicate vascular fibrous membrane which is adherent to the brain.
   B. **Brain** C710-C719
   C. **Cranial nerves** C722-C729. See Table 4: Coding Primary Site for Malignant Tumors of Cranial and Peripheral Nerves
   D. **Intracranial glands** C751-C753
      i. Craniopharyngeal duct C752
      ii. Pineal gland C753
      iii. Pituitary gland C751

Continued on next page
2. **Spinal** sites: includes the **spinal meninges** and **all** structures **within** the meninges (Intradural, within/in the spinal meninges).
   A. **Spinal cord** C720
   B. **Spinal meninges** C701 covers/encloses the spinal nerve roots and the spinal cord.
   C. Spinal nerve roots:
      i. **Cervical** nerve, 8 pair, occipital nerve. Code to peripheral nerve of head, face, and neck C470
      ii. **Coccygeal** nerve, 1 pair. Code to cauda equina C721
      iii. **Lumbar** nerve, 5 pair. Code to cauda equina C721
      iv. **Sacral** nerve, 5 pair. Code to cauda equina C721
      v. **Thoracic** nerve, 12 pair. Code to peripheral nerves of thorax C473

3. **Peripheral nerves**
   i. **Cervical** nerve, 8 pair, occipital nerve. Code to peripheral nerve of head, face, and neck C470
   ii. **Thoracic** nerve, 12 pair. Code to peripheral nerves of thorax C473
   iii. **Lumbar** nerve, 5 pair. Code to cauda equina C721
   iv. **Sacral** nerve, 5 pair. Code to cauda equina C721
   v. **Coccygeal** nerve, 1 pair. Code to cauda equina C721
Use Table 2 to **determine** whether a primary site is **reportable**.

<table>
<thead>
<tr>
<th>Site Group</th>
<th>Reportable Subsite Terms and Code</th>
</tr>
</thead>
</table>
| Brain         | Brain NOS C719  
                Brain stem C717  
                Cerebellum NOS C716  
                Cerebrum C710  
                Frontal lobe C711  
                Occipital lobe C714  
                Overlapping lesion of brain C718  
                Parietal lobe C713  
                Temporal lobe C712  
                Ventricle NOS C715 |
| Cranial Nerves| Abducent (cranial nerve VI) C725  
                Accessory (cranial nerve XI) C725  
                Acoustic (cranial nerve VIII) C724  
                Cranial nerve NOS C725  
                Facial (cranial nerve VII) C725  
                Glossopharyngeal (cranial nerve IX) C725  
                Hypoglossal (cranial nerve XII) C725  
                Oculomotor (cranial nerve III) C725  
                Olfactory (cranial nerve I) C722  
                Optic (cranial nerve II) C723  
                Trigeminal (cranial nerve V) C725  
                Trochlear (cranial nerve IV) C725  
                Vagus (cranial nerve X) C725 |
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<table>
<thead>
<tr>
<th>Site Group</th>
<th>Reportable Subsite Terms and Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ill-Defined Sites Central Nervous System</td>
<td>Nervous system NOS C729</td>
</tr>
<tr>
<td></td>
<td>Overlapping lesion of brain and</td>
</tr>
<tr>
<td></td>
<td>central nervous system C728</td>
</tr>
<tr>
<td>Intracranial Duct and Glands</td>
<td>Craniopharyngeal duct C752</td>
</tr>
<tr>
<td></td>
<td>Pineal gland C753</td>
</tr>
<tr>
<td></td>
<td>Pituitary gland C751</td>
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<tr>
<td>Meninges</td>
<td>Cerebral meninges C700</td>
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<td></td>
<td>Meninges NOS C709</td>
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<tr>
<td></td>
<td>Spinal meninges C701</td>
</tr>
<tr>
<td>Peripheral Nerve and Autonomic Nervous</td>
<td>Abdomen C474</td>
</tr>
<tr>
<td>System</td>
<td>Autonomic nervous system NOS C479</td>
</tr>
<tr>
<td></td>
<td>Head, face and neck C470</td>
</tr>
<tr>
<td></td>
<td>Lower limb and hip C472</td>
</tr>
<tr>
<td></td>
<td>Overlapping lesion of peripheral</td>
</tr>
<tr>
<td></td>
<td>nerves and autonomic nervous</td>
</tr>
<tr>
<td></td>
<td>system C478</td>
</tr>
<tr>
<td></td>
<td>Thorax C473</td>
</tr>
<tr>
<td></td>
<td>Trunk NOS C476</td>
</tr>
<tr>
<td></td>
<td>Upper limbs and shoulder C471</td>
</tr>
<tr>
<td></td>
<td>Spinal Nerve NOS C479</td>
</tr>
<tr>
<td>Spinal Sites</td>
<td>Cauda equina/conus medullaris/filum</td>
</tr>
<tr>
<td></td>
<td>terminale C721</td>
</tr>
<tr>
<td></td>
<td>Meninges NOS C709</td>
</tr>
<tr>
<td></td>
<td>Spinal cord C720</td>
</tr>
<tr>
<td></td>
<td>Spinal meninges C701</td>
</tr>
</tbody>
</table>
Table 3 lists the more common histologies for CNS and Peripheral Nerves. For hematopoietic neoplasms such as lymphomas, myelomas, etc., see the Hematopoietic Database.

**Note:** Behavior codes are not listed because all histologies are malignant /3.

**Column 1** contains specific and NOS histology terms and codes.
- Specific histology terms do not have subtypes/variants
- NOS histology terms do have subtypes/variants

**Column 2** contains synonyms for the specific or NOS term. Synonyms have the same histology code as the specific or NOS term.

**Column 3** contains subtypes/variants of the NOS histology. Subtypes/variants do not have the same histology code as the NOS term.

**Note:** All tumors are malignant/invasive /3.

Column 3 may contain NOS histologies which are part of a bigger histologic group. For example, sarcoma NOS 8800/3 (column 1) is a generic term that encompasses a number of soft tissue tumors, including chondrosarcoma 9220/3 (column 3). Chondrosarcoma is also a NOS because it has a subtype/variant, mesenchymal chondrosarcoma 9240. The subtype/variant is indented under the NOS (chondrosarcoma) in column 3. There is also a note in column 1 which calls attention to the fact that chondrosarcoma has a subtype/variant.

When using the Solid Tumor Rules, chondrosarcoma and mesenchymal chondrosarcoma are treated the same as all NOS and subtypes/variants.

**Table begins on next page**
## Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Specific and NOS Histology Codes</th>
<th>Synonyms</th>
<th>Subtypes/Variants</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anaplastic ganglioglioma 9505</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Astroblastoma 9430</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Astrocytoma NOS 9400</td>
<td>Diffuse astrocytoma IDH-mutant&lt;br&gt;Diffuse astrocytoma IDH-wildtype&lt;br&gt;Diffuse astrocytoma NOS</td>
<td>Anaplastic astrocytoma IDH-mutant/wildtype; anaplastic astrocytoma NOS 9401&lt;br&gt;Gemistocytic astrocytoma IDH-mutant 9411&lt;br&gt;Pleomorphic xanthoastrocytoma/ anaplastic pleomorphic xanthoastrocytoma 9424</td>
</tr>
<tr>
<td>Choriocarcinoma 9100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Choroid plexus carcinoma 9390</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CNS embryonal tumor with rhabdoid features 9508</td>
<td>Atypical teratoid/rhabdoid tumor&lt;br&gt;Embryonal tumor with rhabdoid features</td>
<td>CNS embryonal tumor 9473</td>
</tr>
<tr>
<td>CNS ganglioneuroblastoma 9490</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CNS neuroblastoma 9500</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diffuse midline glioma H3 K27M mutant 9385*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Embryonal carcinoma 9070</td>
<td></td>
<td>Yolk sac tumor 9071</td>
</tr>
<tr>
<td>Embryonal tumor with multilayered rosettes C19MC-altered 9478*</td>
<td>Embryonal tumor with multilayered rosettes, NOS&lt;br&gt;ETMR</td>
<td>Anaplastic ependymoma 9392&lt;br&gt;Ependymoma, RELA fusion-positive 9396*&lt;br&gt;Papillary ependymoma 9393</td>
</tr>
<tr>
<td>Ependymoma 9391</td>
<td>Clear cell ependymoma&lt;br&gt;Tanycytic ependymoma</td>
<td></td>
</tr>
<tr>
<td>Epithelioid hemangioendothelioma 9133</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Germinoma 9064</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

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Jump to [Histology Coding Rules](#)
## Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753

(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

### Specific and NOS Histology Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Synonyms</th>
<th>Subtypes/Variants</th>
</tr>
</thead>
<tbody>
<tr>
<td>9440</td>
<td>Glioblastoma NOS</td>
<td>Giant cell glioblastoma 9441</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Glioblastoma IDH-mutant 9445*</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Gliosarcoma 9442</td>
</tr>
<tr>
<td>9080</td>
<td>Immature teratoma</td>
<td>Mixed germ cell tumor 9085</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Teratoma with malignant transformation 9084</td>
</tr>
<tr>
<td>9530</td>
<td>Malignant meningioma</td>
<td>Papillary/rhabdoid meningioma 9538</td>
</tr>
<tr>
<td>9540</td>
<td>Malignant peripheral nerve sheath tumor</td>
<td>Anaplastic medulloblastoma 9474</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Medulloblastoma described as one of the following 9471</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Desmoplastic</td>
</tr>
<tr>
<td></td>
<td></td>
<td>SHH-activated and TP53-wildtype</td>
</tr>
<tr>
<td></td>
<td></td>
<td>With extensive nodularity</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nodular</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Medulloblastoma non-WNT/non-SHH; medulloblastoma group 4</td>
</tr>
<tr>
<td></td>
<td></td>
<td>9477*</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Medulloblastoma SHH-activated and TP53-mutant 9476*</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Medulloblastoma WNT-activated 9475*</td>
</tr>
<tr>
<td>9501</td>
<td>Medulloepithelioma</td>
<td></td>
</tr>
<tr>
<td>8720</td>
<td>Meningeal melanoma</td>
<td>Meningeal melanomatosis 8728</td>
</tr>
<tr>
<td>9382</td>
<td>Oligoastrocytoma NOS</td>
<td>Anaplastic oligoastrocytoma NOS</td>
</tr>
</tbody>
</table>

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### Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

**C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753**  
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Specific and NOS Histology Codes</th>
<th>Synonyms</th>
<th>Subtypes/Variants</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Oligodendroglioma NOS 9450</strong></td>
<td>Oligodendroglioma 1p/19q-codeleted</td>
<td>Anaplastic oligodendroglioma NOS 9451</td>
</tr>
<tr>
<td><em>Note:</em> Oligodendroglioma NOS is used when molecular markers cannot fully be determined</td>
<td>Oligodendroglioma IDH-mutant</td>
<td>IDH-mutant</td>
</tr>
<tr>
<td></td>
<td>Oligodendroglioma IDH-mutant and 1p/19q-codeleted</td>
<td>1p/19q-codeleted</td>
</tr>
<tr>
<td><strong>Peripheral primitive neuroectodermal tumor 9364</strong></td>
<td>Ewing sarcoma</td>
<td>Pilomyxoid astrocytoma 9425</td>
</tr>
<tr>
<td><strong>Pilocytic astrocytoma 9421</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Note:</em> ICD-O-3 lists a behavior code of /1. It is collected as /3 in North America.</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Pineal parenchymal tumor of intermediate differentiation 9362</strong></td>
<td>Pineoblastoma</td>
<td>Papillary tumor of the pineal region 9395</td>
</tr>
<tr>
<td><strong>Sarcoma NOS 8800</strong></td>
<td></td>
<td>Angiosarcoma 9120</td>
</tr>
<tr>
<td><em>Note 1: Chondrosarcoma 9220</em></td>
<td></td>
<td>Chondrosarcoma 9220</td>
</tr>
<tr>
<td>has the following subtype/variant:</td>
<td></td>
<td>Mesenchymal chondrosarcoma 9240</td>
</tr>
<tr>
<td>Mesenchymal chondrosarcoma 9240</td>
<td></td>
<td>Leiomyosarcoma/granular cell</td>
</tr>
<tr>
<td><em>Note 2: Leiomyosarcoma 8890</em></td>
<td></td>
<td>leiomyosarcoma/inflammatory</td>
</tr>
<tr>
<td>has the following subtypes/variants:</td>
<td></td>
<td>leiomyosarcoma 8890</td>
</tr>
<tr>
<td>Epithelioid leiomyosarcoma 8891</td>
<td></td>
<td>Epithelioid leiomyosarcoma 8891</td>
</tr>
<tr>
<td>Myxoid leiomyosarcoma 8896</td>
<td></td>
<td>Myxoid leiomyosarcoma 8896</td>
</tr>
<tr>
<td><strong>Solitary fibrous tumor grade 3 8815</strong></td>
<td>Hemangiopericytoma grade 3</td>
<td>Osteosarcoma 9180</td>
</tr>
<tr>
<td></td>
<td>Solitary fibrous tumor/Hemangiopericytoma grade 3 (CNS)</td>
<td>Undifferentiated pleomorphic sarcoma/malignant fibrous histiocytoma 8802</td>
</tr>
</tbody>
</table>

* These new codes were approved by the IARC/WHO Committee for ICD-O
Table 4: Coding Primary Site for Malignant Tumors of Cranial and Peripheral Nerves

**Note 1:** Neoplasms arising in a cranial or spinal nerve are coded to the specific nerve in which it arises.
**Note 2:** Neoplasms, commonly meningiomas, arising in the dura/meninges of an intracranial nerve (cranial nerve within the skull) are coded to cerebral meninges C700.
**Note 3:** Neoplasms, commonly meningiomas, arising in the dura/meninges of the spinal nerve roots are coded to the ICD-O site code spinal meninges C701.
**Note 4:** It is important to check the operative report to determine whether the surgery is intracranial or intradural.

Use Table 4 to determine whether a malignant cranial nerve tumor should be coded to a cranial nerve or to a peripheral nerve. The anatomic location of the tumor determines whether the primary site is a peripheral or a cranial nerve.

**Column 1:** The proper name for the cranial nerve (CN) and the cranial nerve number  
**Column 2:** The point at which the nerve exits the cranium  
**Column 3:** Portions of the nerve coded to cranial nerve  
**Column 4:** Portions of nerve coded to peripheral nerve

<table>
<thead>
<tr>
<th>Name and CN #</th>
<th>Exits Cranium Through</th>
<th>Site Code: Cranial Nerve</th>
<th>Site Code: Peripheral Nerve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cranial nerve NOS</td>
<td></td>
<td>Within cranium, unknown which nerve C725</td>
<td></td>
</tr>
<tr>
<td>Olfactory CN 1</td>
<td>Cribiform plate</td>
<td>Surface of the brain C722</td>
<td>Originates on the olfactory mucosa of nasal cavity, then travels through the cribiform plate of the ethmoid bone C470</td>
</tr>
<tr>
<td>Optic CN 2</td>
<td>Optic canal</td>
<td>All portions are covered by meninges/dura so are reportable as C723</td>
<td></td>
</tr>
<tr>
<td>Oculomotor CN 3</td>
<td>Superior orbital fissure</td>
<td>Originates in the midbrain C725</td>
<td>After exiting the superior orbital fissure, the nerve enters the orbit C470</td>
</tr>
</tbody>
</table>
### Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

*C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753*  
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Name and CN #</th>
<th>Exits Cranium Through</th>
<th>Site Code: Cranial Nerve</th>
<th>Site Code: Peripheral Nerve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Trochlear CN 4</td>
<td>Superior orbital fissure</td>
<td>Arises from the <strong>dorsal brain stem</strong>, loops around the brainstem and passes anteriorly within the <strong>subarachnoid space</strong>. It travels between the <strong>superior cerebellar and posterior cerebral arteries</strong> and through the <strong>dura</strong>, enters <strong>cavernous sinus C725</strong></td>
<td>Enters the orbital fissure <strong>C470</strong></td>
</tr>
</tbody>
</table>
| Trigeminal CN 5 | The ophthalmic, maxillary and mandibular branches leave the skull through three separate foramina.  
Superior orbital fissure.  
The foramen rotundum  
The foramen ovale. | CN5 originates in the **pons**. Upon leaving the pons it enters a **small fossa** posterior and inferolateral to the cavernous sinus called **Meckel's (trigeminal) cave C725**.  
• **Ophthalmic nerve branch** crosses the **pterygopalatine fossa**, inclines laterally on the back of the **maxilla**, and enters the **orbit** through the inferior orbital fissure where it is called the **infraorbital** nerve. It ends beneath the **quadatus labii superius**, and divides into multiple branches that spread to the side of the **nose**, the lower **eyelid**, and the upper **lip C470**  
• **Maxillary nerve** leaves foramen rotundum and traverses the **infraorbital groove** and **canal** in the floor of the **orbit**, and appears on the face at the infraorbital foramen **C470**  
• **Mandibular** nerve leaves via the foramen ovale travels along the **mandibular groove C470** | |

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### Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753

(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Name and CN #</th>
<th>Exits Cranium Through</th>
<th>Site Code: Cranial Nerve</th>
<th>Site Code: Peripheral Nerve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abducent CN 6</td>
<td>Cranial meninges</td>
<td>Exits brainstem at junction of <em>pons</em> and the <em>medulla</em>, enters the <em>subarachnoid</em> space and runs upward between the pons and the <em>clivus</em> entering the <em>cavernous sinus C725</em></td>
<td><em>Dorello's canal</em> and travels to the tip of the <em>temporal bone</em>. Enters <em>orbit C470</em></td>
</tr>
<tr>
<td>Facial CN 7</td>
<td>Internal acoustic meatus</td>
<td><em>CN7</em> originates in the <em>pons</em>, along the posterior cranial fossa (<em>posterior cranial fossa</em> is part of the intracranial cavity.) <em>C725</em></td>
<td>Enters the temple through the internal auditory meatus and runs through the facial canal. <em>C470</em></td>
</tr>
<tr>
<td>Acoustic or vestibulocochlear CN 8</td>
<td>Internal acoustic meatus</td>
<td>Originates in the <em>brain stem (medulla oblongata) between</em> the base of the brain (<em>pons</em>) and the <em>spinal cord C724</em></td>
<td>Both the <em>vestibular</em> branch and the <em>cochlear</em> branch are located in the <em>inner ear</em></td>
</tr>
<tr>
<td>Glossopharyngeal CN 9</td>
<td>Jugular foramin&lt;sup&gt;a&lt;/sup&gt;</td>
<td><em>Originates</em> in the anterior portion of the <em>medulla oblongata C725</em></td>
<td><em>Jugular foramen</em> Between the <em>internal jugular vein</em> and <em>internal carotid</em> artery Lies on the <em>stylopharyngeus</em> and middle <em>pharyngeal</em> constrictor muscle Passes under the <em>hypoglossus</em> muscle Palatine tonsil Extends to mucous <em>glands</em> of the <em>mouth</em>, and <em>base</em> of the <em>tongue C470</em></td>
</tr>
</tbody>
</table>

---

<sup>a</sup> Vertical line above the name indicates a branch of the nerve.
### Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions

**C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753**

(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

<table>
<thead>
<tr>
<th>Name and CN #</th>
<th>Exits Cranium Through</th>
<th>Site Code: Cranial Nerve</th>
<th>Site Code: Peripheral Nerve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vagus CN 10</td>
<td>Jugular foramen</td>
<td>The vagus nerve originates from the <strong>medulla of the brainstem</strong>. <strong>C725</strong></td>
<td>CN10 descends within the <strong>carotid sheath</strong> medial to the internal jugular vein at the root of the <strong>neck C470</strong>. The right vagus crosses in front of the subclavian artery and travels into the <strong>fat behind</strong> the blood vessels, reaching the <strong>thorax</strong>. It then inclines <strong>behind</strong> the <strong>hilum</strong> of the <strong>right lung</strong> and moves toward the esophagus. The nerve splits into the <strong>right</strong> and <strong>left</strong> vagus at the <strong>esophageal plexus C473</strong>. Forms the anterior and posterior <strong>gastric nerves C475</strong></td>
</tr>
<tr>
<td>Accessory CN 11</td>
<td>Jugular foramen</td>
<td>The spinal accessory nerve <strong>originates</strong> in the neurons of the upper spinal cord, specifically <strong>C1-C5/C6</strong> spinal nerve <strong>roots</strong>. The nerve enters the foramen magnum or lateral aspect of the <strong>medulla oblongata</strong>. The fibers of the spinal accessory nerve coalesce to form <strong>spinal rootlets, roots</strong>, and finally the spinal accessory nerve itself <strong>C725</strong></td>
<td>The nerve <strong>exits</strong> the skull through the jugular foramen. It then runs along the internal <strong>carotid artery</strong> within the <strong>neck C470</strong>. Reaches the sternocleidomastoid muscle and the trapezius <strong>C476</strong></td>
</tr>
<tr>
<td>Hypoglossal CN 12</td>
<td>Hypoglossal canal</td>
<td><strong>CN12</strong> starts in the <strong>hypoglossal nucleus of the brainstem, C725</strong></td>
<td><strong>CN12</strong> exits the hypoglossal canal, traveling <strong>between</strong> the <strong>carotid artery</strong> and <strong>jugular vein</strong>, ending under the <strong>tongue C470</strong></td>
</tr>
</tbody>
</table>

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**Conflicting Information on Pathology Report(s)**

The *classification* of brain tumors is a *subjective* matter because definitive criteria have not been established/accepted. Pathologists may *disagree* on the *histology* or *behavior*.

Follow the steps below for coding the histology and/or behavior of tumor when there are discrepancies:

- When possible, get advice from the *pathologist*
- When possible, contact *attending* physician
- When possible, consult with *registry advisor*
- If none of those *options* are *available*, code the histology and behavior from the *most dependable source* (see *Priority List for Coding Histology*).

The following are examples of how conflicting information occurs in single and in multiple pathology reports:

- **Single pathology report:**
  - *Multiple pathologists* within the institution review the slides
  - Slides are sent for *outside review* and the information from the *consulting* lab *conflicts* with the *original* pathology report

- **Multiple pathology reports:** The *first report* is from a biopsy and the *second report* is from a resection. Code the histology and/or behavior from the resection.
Table 5: Paired Sites

Use Table 5 to identify sites for which laterality **must** be coded. Do **not** use this table to determine multiple primaries.

<table>
<thead>
<tr>
<th>Paired Sites and Codes</th>
<th>Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acoustic nerve</td>
<td>C724</td>
</tr>
<tr>
<td>Cerebral meninges</td>
<td>C700</td>
</tr>
<tr>
<td>Cerebrum</td>
<td>C710</td>
</tr>
<tr>
<td>Cranial nerves</td>
<td>C725</td>
</tr>
<tr>
<td>Frontal lobe</td>
<td>C711</td>
</tr>
<tr>
<td>Occipital lobe</td>
<td>C714</td>
</tr>
<tr>
<td>Olfactory nerve</td>
<td>C722</td>
</tr>
<tr>
<td>Optic nerve</td>
<td>C723</td>
</tr>
<tr>
<td>Parietal lobe</td>
<td>C713</td>
</tr>
<tr>
<td>Temporal lobe</td>
<td>C712</td>
</tr>
</tbody>
</table>

**Note 1:** Midline tumors are common for glioblastoma multiform and meningiomas.

**Note 2:** SEER allows laterality to be coded for sites other than those in the table.
Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Table 6: Non-malignant CNS Tumors with the Potential of Transforming to Malignant Behavior

The word “transformation” as used in this table means that:
- Residual tumor becomes more aggressive /3 OR
- The tumor recurs as a more aggressive /3 histology

The table identifies non-malignant tumors that have the potential of transforming to a malignant tumor (new primary).

Use Table 6 when directed to by the Multiple Primary Rules.

Column 1 is the non-malignant ICD-O histology term and code.
Column 2 is the malignant /3 ICD-O histology term and code to which the non-malignant tumor can transform.

<table>
<thead>
<tr>
<th>Original Histology and Code</th>
<th>Transformed Histology and Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chondroma 9220/0</td>
<td>Chondrosarcoma 9220/3</td>
</tr>
<tr>
<td>Ganglioglioma 9505/1</td>
<td>Anaplastic ganglioglioma 9505/3</td>
</tr>
<tr>
<td>Hemangioma 9120/0</td>
<td>Angiosarcoma 9120/3</td>
</tr>
<tr>
<td>Hemangiopericytoma 9150/1</td>
<td>Anaplastic hemangiopericytoma 9150/3</td>
</tr>
<tr>
<td>Leiomyoma 8890/0</td>
<td>Leiomyosarcoma 8890/3</td>
</tr>
<tr>
<td>Lipoma 8850/0</td>
<td>Liposarcoma 8850/3</td>
</tr>
<tr>
<td>Osteoma 9180/0</td>
<td>Osteosarcoma 9180/3</td>
</tr>
<tr>
<td>Perineurioma 9571/0</td>
<td>Malignant perineurioma 9571/3</td>
</tr>
<tr>
<td>Rhabdomyoma 8900/0</td>
<td>Rhabdomyosarcoma 8900/3</td>
</tr>
<tr>
<td>Teratoma 9080/1</td>
<td>Immature teratoma 9080/3</td>
</tr>
<tr>
<td>Teratoma, mature 9080/0</td>
<td>Immature teratoma 9080/3</td>
</tr>
</tbody>
</table>
Malignant CNS and Peripheral Nerves Equivalent Terms and Definitions
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
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Malignant CNS and Peripheral Nerves Multiple Primary Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Note 1: Non-malignant intracranial and CNS tumors have a separate set of rules.
Note 2: Laterality is not used to determine multiple primaries for malignant CNS tumors.
Note 3: Timing is not used to determine multiple primaries for malignant CNS tumors.
Note 4: GBM following an astrocytic or glial tumor is a multiple primary.
Note 5: These rules are NOT used for tumor(s) described as metastases.
Note 6: 2007 MPH Rules and 2018 Solid Tumor Rules are used based on date of diagnosis.
  • Tumors diagnosed 01/01/2007 through 12/31/2017: Use 2007 MPH Rules
  • Tumors diagnosed 01/01/2018 and later: Use 2018 Solid Tumor Rules
  • The original tumor diagnosed before 1/1/2018 and a subsequent tumor diagnosed 1/1/2018 or later in the same primary site: Use the 2018 Solid Tumor Rules.

Unknown If Single or Multiple Tumors

Rule M1 Abstract a single primary\(^1\) when it is not possible to determine if there is a single tumor or multiple tumors.

Note 1: Use this rule only after all information sources have been exhausted
Note 2: Examples of cases with minimal information include
  • Death certificate only (DCO)
  • Cases for which information is limited to pathology report only
  • Outpatient biopsy with no follow-up information available
  • Multiple pathology reports which do not specify whether a single tumor or multiple tumors have been biopsied and/or resected

This is the end of instructions for Unknown if Single or Multiple Tumors.

\(^1\) Prepare one abstract. Use the histology rules to assign the appropriate histology code.
Malignant CNS and Peripheral Nerves Multiple Primary Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Single Tumor

**IMPORTANT:** The **major difference** between M4 and M5 is:
M4: No resection as first course of treatment AND when the same tumor is subsequently resected, pathology proves malignant behavior
M5: Tumor resected as first course of treatment. Subsequent tumor (recurrence or de novo) is malignant

**Rule M2**
Abstract a single primary when there is a single tumor.

*Note 1:* A single tumor is always a single primary.
*Note 2:* The tumor may overlap onto or extend into adjacent/contiguous site or subsites.
*Note 3:* The tumor may have two or more histologic components.

**Rule M3**
Code a single primary when a neoplasm is originally diagnosed as an oligodendroglioma and subsequently recurs in residual tumor tissue with different features such as a densely cellular tumor with pseudo palisading necrosis.

*Note 1:* The pathology may state that the recurrence “looks like” or “has the appearance of” a glioblastoma multiforme (GBM). This is not a true GBM.
*Note 2:* Record as a recurrence for those registrars who collect recurrence data.

**Rule M4**
Abstract a single primary (the malignant) when a single tumor meets the following two criteria:

1. The original diagnosis was non-malignant /0 or /1 AND
   - First course treatment was active surveillance (no tumor resection). Diagnosis was:
     - Clinical
     - Radiographic
     - Stereotactic biopsy

2. Subsequent resection pathology is malignant /3

*Note 1:* This is a new rule which clarifies that a single tumor is always a single primary and the malignant behavior is reported.
*Note 2:* The resection pathology is more accurate than clinical, radiographic or stereotactic biopsy information. While stereotactic biopsy provides a pathologic specimen, it is small and may not have included the malignant portion of tumor.
*Note 3:* There is no time requirement from initial diagnosis to resection.
*Note 4:* Edit the original abstract as follows:
• Do not change date of diagnosis.
• For cases which have been abstracted, change behavior code on original abstract from /0 or /1 to /3.
• Report all data changes for cases which have been submitted to the central registry.
• See the COC and SEER manuals for instructions on coding other data items such as Accession Year, Treatment and Sequence Number.

Note 5: The physician may have staged the non-malignant tumor at the time of original diagnosis (benign). The physician will also stage the malignant tumor. Staging and determining multiple primaries are done for different reasons. Staging determines which treatment would be most effective. Determining multiple primaries is done to stabilize the data for the study of epidemiology (long-term studies done on incidence, mortality, and causation of a disease with the goal of reducing or eliminating that disease).

Example 1: A patient is diagnosed by MRI with ganglioglioma 9505/1 in July 2018. After months of active surveillance (watchful waiting), the patient becomes symptomatic. A resection is done in April 2019; the resection pathology is anaplastic ganglioglioma 9505/3. Change behavior code on the original abstract from /1 to /3. Do not change date of diagnosis.

Example 2: A November 15, 2017 MRI and subsequent stereotactic biopsy of lateral ventricle of brain show a mature teratoma 9080/1. The patient becomes symptomatic in 2018 and the neoplasm is resected on October 31, 2018. The resection pathology diagnoses immature teratoma 9080/3. Change behavior code on the original abstract. Do not change date of diagnosis.

This is the end of instructions for Single Tumor.

Prepare one abstract. Use the histology coding rules to assign the appropriate histology code.
Malignant CNS and Peripheral Nerves Multiple Primary Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Multiple Tumors

**Note 1:** Multiple tumors may be a single primary or multiple primaries.

**Note 2:** Separate, non-contiguous tumors are always multiple primaries when:

- In the CNS (see Table 2) AND in a site other than the CNS
  
  **Example:** Patient has a malignant nerve sheath tumor of the right arm C471 and non-metastatic adenocarcinoma of the lung. The lung is not a CNS site. Abstract two primaries.

- In different CNS sites (see Rule M8)

**IMPORTANT:** The major difference between M4 and M5 is:

M4: No resection as first course of treatment AND when the same tumor is subsequently resected, pathology proves malignant behavior

M5: Tumor resected as first course of treatment. Subsequent tumor (recurrence or de novo) is malignant

**Rule M5** Abstract multiple primaries\(^4\) when there are multiple CNS tumors, one of which is malignant /3 and the other is non-malignant /0 or /1.

- Original non-malignant tumor followed by malignant tumor
  - Patient had a resection of the non-malignant tumor (not the same tumor) OR
  - It is unknown/not documented if the patient had a resection

- Simultaneous non-malignant and malignant tumors
  - Abstract both the malignant and the non-malignant tumors

**Note 1:** The rules are hierarchical. Only use when previous rules do not apply.

**Note 2:** See Table 2 in the Equivalent Terms and Definitions for a listing of CNS sites.

**Note 3:** A non-malignant CNS tumor and a malignant CNS tumor are always multiple primaries (timing and primary sites are irrelevant). Prepare two abstracts; one for the non-malignant and another for the malignant tumor.
Rule M6  Abstract multiple primaries\(^4\) when a patient has a glial tumor and is subsequently diagnosed with a glioblastoma multiforme 9440 (GBM).

Note 1: Definition of a glial tumor: Any tumor arising from the CNS glial cells. The following is simply a list of all tumors which would be classified as glial.
- Astroblastoma 9430
- Astrocytomas 9400 and all subtypes
  - Anaplastic astrocytoma IDH-mutant/wildtype; anaplastic astrocytoma NOS 9401
  - Gemistocytic astrocytoma IDH-mutant 9411
- Diffuse midline glioma H3 K27M Mutant 9385
- Ependymoma 9391 and all subtypes
  - Anaplastic ependymoma 9392
  - Ependymoma, RELA fusion-positive 9396
  - Papillary ependymoma 9393
- Glioblastoma 9440 and all subtypes (this is a glial tumor; however do not apply this rule to a GBM followed by a GBM)
  - Giant cell glioblastoma 9441
  - Glioblastoma IDH-mutant 9445
  - Gliosarcoma 9442
- Oligodendrogliaoma and all subtypes 9450
  - Anaplastic oligodendroglialoma; IDH-mutant; 1p/19q-codeleted; IDH-mutant and 1p/19q-codeleted 9451
- Pleomorphic xanthroastrocytoma 9424

Note 2: This is a change from the 2007 Rules.

Note 3: Abstracting GBM as a new primary will allow analysis of:
- The number of tumors that recur as a more aggressive histology (GBM)
- The time interval between occurrence of the glial or astrocytic tumors and a GBM
- Which histologies are more likely to recur as a GBM
Rule M7  Abstract a **single primary** when there are separate, non-contiguous tumors in the brain (multicentric/multifocal) with the same histology. Tumors may be any of the following combinations:

- In the same lobe; for example, two tumors in right temporal lobe C712 (same site code)
- Different lateralties of the same lobe; for example, left and right frontal lobes C711 (same site code)
- In different lobes; for example, parietal lobe C713 and occipital lobe C714 (different site codes)

**Example:** The patient had a resection of an anaplastic astrocytoma 9401 in the right parietal lobe. Three months later the patient is diagnosed with a de novo anaplastic astrocytoma in the left parietal lobe. This is one primary because neither laterality nor timing are used to determine multiple primary status.

**Note 1:** Multiple sites/subsites and/or different lateralities imply either metastatic or multifocal/multicentric disease. Glioblastoma multiforme commonly exhibits multiple tumors which are described as multifocal/multicentric.

**Note 2:** Metastases are never used to determine multiple primaries. Seeding metastasis is often noted for the following tumors:

- Glioblastoma multiforme
- pNET-medulloblastoma
- Oligodendroglioma

**Note 3:** Hereditary syndromes frequently exhibit multiple tumors including the following:

- Neurofibromatosis type 1 (NF1)
  - Malignant peripheral nerve sheath tumors (MPNST)
- Neurofibromatosis type 2 (NF2)
  - Anaplastic ependymomas
  - Meningiomas

**Note 4:** Most malignant neoplasms are **single tumors** with the exception of those listed in this rule.

**Note 5:** This is a change from/clarification to previous rules.
Rule M8  Abstract multiple primaries when multiple tumors are present in any of the following sites or subsites:

- Any lobe of the brain C710-C719 AND any other part of CNS
- Cauda equina C721 AND any other part of CNS
- Cerebral meninges C700 AND spinal meninges C701
- Cerebral meninges C700 AND any other part of CNS
- Any one of the cranial nerves C722-C725 AND any other part of the CNS
- Any two or more of the cranial nerves
  - C722 Olfactory, C723 Optic, C724 Acoustic, C725 Cranial nerves NOS
- Meninges of cranial or peripheral nerves C709 AND any other part of the CNS
- Spinal cord C720 AND any other part of CNS
- Spinal meninges C701 AND any other part of CNS

Rule M9  Abstract multiple primaries when separate, non-contiguous tumors are two or more different subtypes/variants in Column 3, Table 3 in the Equivalent Terms and Definitions. Timing is irrelevant.

Note: The tumors may be subtypes/variants of the same or different NOS histologies.

- Same NOS: Anaplastic astrocytoma IDH-mutant 9401 and gemistocytic astrocytoma IDH-mutant 9411 are both subtypes of astrocytoma NOS 9400/3 but are distinctly different histologies. Abstract multiple primaries.
- Different NOS: Papillary ependymoma 9393 is a subtype of ependymoma NOS 9391; gliosarcoma 9442 is a subtype of glioblastoma NOS 9440. They are distinctly different histologies. Abstract multiple primaries.

Rule M10  Abstract a single primary when separate, non-contiguous tumors are on the same row in Table 3 in the Equivalent Terms and Definitions. Timing is irrelevant.

Note: The same row means the tumors are:

- The same histology (same four-digit ICD-O code) OR
- One is the preferred term (column 1) and the other is a synonym for the preferred term (column 2) OR
- A NOS (column 1/column 2) and the other is a subtype/variant of that NOS (column 3)
Malignant CNS and Peripheral Nerves Multiple Primary Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
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Rule M11  Abstract multiple primaries\(^i\) when separate, non-contiguous tumors are on different rows in Table 3 in the Equivalent Terms and Definitions. Timing is irrelevant.

*Note:* Each row in the table is a distinctly different histology.

Rule M12  Abstract a single primary\(^i\) when multiple tumors do not meet any of the above criteria.

*Note:* Use this rule as a last resort. Please confirm that you have not overlooked an applicable rule.

This is the end of instructions for Multiple Tumors.

\(^i\) Prepare one abstract. Use the histology coding rules to assign the appropriate histology code. For registries collecting recurrence data: When a subsequent tumor is “single primary,” record that subsequent tumor as a recurrence.

\(^i\) Prepare two or more abstracts. Use the histology rules to assign the appropriate histology code to each case abstracted.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Note: Non-malignant CNS tumors have a separate set of rules.

Priority Order for Using Documentation to Identify Histology

IMPORTANT NOTES

   Note 1: Histology changes do occur following immunotherapy, chemotherapy and radiation therapy.
   Note 2: Neoadjuvant treatment is any tumor-related treatment given prior to surgical removal of the malignancy.

2. Code the histology assigned by the physician. Do not change histology in order to make the case applicable to staging.

The priority list is used for single primaries (including multiple tumors abstracted as a single primary)

This is a hierarchical list of source documentation.

1. Pathology/tissue from resection of primary tumor
   A. Biomarkers
      Note 1: Biomarkers are emerging as an important part of cancer diagnosis and treatment. Some biomarkers are used to determine treatment rather than histologic type. The efficacy of identification of histologic type using biomarkers differs from primary site to primary site. When a histologic type is identified using a biomarker, code the identified histology. Biomarkers do not identify all histologic types.
      Note 2: Biomarkers are not listed because they change rapidly.
      Example: Biomarkers are required to diagnose medulloblastoma SHH-activated and TP53-wildtype. Biomarkers include genetic testing, FISH, MYCN.
   B. The addendum(s) and/or comment(s)
   C. Final diagnosis / synoptic report as required by CAP
   D. CAP protocol

2. Pathology/tissue from biopsy of primary tumor
   A. Biomarkers
      Note 1: Biomarkers are emerging as an important part of cancer diagnosis and treatment. Some biomarkers are used to determine treatment rather than histologic type. The efficacy of identification of histologic type using biomarkers differs from primary site to primary site. When a histologic type is identified using a biomarker, code the identified histology. Biomarkers do not identify all histologic types.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

**Note 2:** Biomarkers are not listed because they change rapidly.

*Example:* Biomarkers are required to diagnose medulloblastoma SHH-activated and TP53-wildtype. Biomarkers include genetic testing, FISH, MYCN.

B. The addendum and/or comments
C. Final diagnosis / synoptic report as required by CAP
D. CAP protocol

**Note 1:** Addendums and comments are given the second priority because they often contain information about molecular testing, genetic testing, and/or special stains which give a more specific diagnosis.

**Note 2:** The pathologist’s diagnosis is always the most reliable, so the final diagnosis is the third priority.

**Note 3:** Do not use the microscopic or gross section of the pathology report for coding.

**Note 4:** The CAP protocol is a checklist which
- Provides guidelines for collecting the essential data elements for complete reporting of tumors and optimal patient care.
- Allows physicians to check multiple histologies

3. **Cytology** (most frequently cerebrospinal fluid)

4. Tissue/pathology from a metastatic site

**Note 1:** Code the behavior /3

**Note 2:** The tissue from a metastatic site often shows variations from the primary tumor. When it is the only tissue available, it is more accurate than a scan.

5. **Scan:** The following list is in priority order.
   A. MRI
   B. CT
   C. PET
   D. Angiogram

6. Code the histology documented by the physician when none of the above are available. Use the documentation in the following priority order
   A. Treatment plan
   B. Documentation from Tumor Board
   C. Documentation in the medical record that refers to original pathology, cytology, or scan(s)
   D. Physician’s reference to type of cancer (histology) in the medical record

**Note 1:** Code the specific histology when documented.

**Note 2:** Code the histology to malignant neoplasm NOS 8000 or as stated by the physician when nothing more specific is documented.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Coding Histology

**Note 1:** The priority is to code the most specific histology. **DO NOT USE BREAST HISTOLOGY CODING RULES FOR THIS SITE.**

**Note 2:** Only use this section for one or more histologies within a single tumor.

**Note 3:** Do not use this section in place of the Histology Rules.

1. **Code the most specific histology or subtype/variant, regardless of whether it is described as:**
   - **A.** The majority or predominant part of tumor
   - **B.** The minority of tumor
   - **C.** A component

   **Example 1:** Diagnosis for a single tumor is astrocytoma 9400 with the majority or predominant part of tumor being anaplastic astrocytoma IDH-mutant 9401. Code the subtype/variant: anaplastic astrocytoma IDH-mutant 9401.

   **Example 2:** Diagnosis for a single tumor is CNS ganglioneuroblastoma 9490 with minority of tumor being CNS embryonal tumor 9473. Code the subtype/variant: CNS embryonal tumor 9473.

   **Example 3:** Diagnosis for a single tumor is ependymoma 9391 with a component of anaplastic ependymoma 9392. Code the subtype/variant: anaplastic ependymoma 9392.

   **Note:** When the most specific histology is described as differentiation or features, see #2.

2. **Code the histology described as differentiation or features/features of ONLY when there is a specific ICD-O code for the “NOS with ____ features” or “NOS with ____ differentiation.”**

   **Note:** Do not code differentiation or features when there is no specific ICD-O code.

3. **Code the specific histology described by ambiguous terminology (list follows) ONLY when A or B is true:**
   - **A.** The only diagnosis available is one histology term described by ambiguous terminology
     - CoC and SEER require reporting of cases diagnosed only by ambiguous terminology
     - Case is accessioned (added to your database) based on ambiguous terminology and no other histology information is available/document.

   **Example:** Outpatient biopsy says probably desmoplastic medulloblastoma. The case is accessioned (entered into the database) as required by both SEER and COC. No further information is available. Code the histology desmoplastic medulloblastoma. The case meets the criteria in #3A.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
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B. There is a NOS histology and a more specific (subtype/variant) described by ambiguous terminology
   • Specific histology is clinically confirmed by a physician (attending, pathologist, oncologist, etc.) OR
   • Patient is receiving treatment based on the specific histology described by ambiguous term

   Example 1: The pathology diagnosis is astrocytoma consistent with gemistocytic astrocytoma IDH-mutant. The oncology consult says the patient has gemistocytic astrocytoma IDH-mutant in the occipital lobe. This is clinical confirmation of the diagnosis, code gemistocytic astrocytoma IDH-mutant. The case meets the criteria in bullet 1.

   Example 2: The pathology diagnosis is ependymoma consistent with papillary ependymoma. The treatment plan says the patient will receive the following treatment for papillary ependymoma. Treatment plan confirms papillary ependymoma; code papillary ependymoma. The case meets the criteria in bullet 2.

   If the specific histology does not meet the criteria in #3B, then code the NOS histology.

   List of Ambiguous Terminology

   Apparently Most likely
   Appears Presumed
   Comparable with Probable
   Compatible with Suspect(ed)
   Consistent with Suspicious (for)
   Favor(s) Typical (of)
   Malignant appearing

4. **Do not code** histology when described as:
   • Architecture
   • Foci; focus; focal
   • Pattern
**Single Tumor**

**Rule H1** Code the reportable CNS tumor (Table 3 in the Equivalent Terms and Definitions) when a patient has any of the following:
- Neurofibromatosis type 1 (NF1)
- Neurofibromatosis type 2 (NF2)
- Schwannomatosis

*Note 1:* Do not code NF1 or NF2 as neurofibromatosis. NF1, NF2, and schwannomatosis are genetic syndromes which have a high risk of developing reportable and non-reportable tumors. ONLY abstract reportable tumors such as malignant peripheral nerve sheath tumors.

*Note 2:* Tumors are reportable when they meet the behavior (/3) and histology requirements (see Reportability Criteria).

*Note 3:* Schwannomatosis is a newer term for a distinct subtype/variant of the genetic diseases NF1 and NF2.

*Example:* Patient presents with vestibular schwannoma (acoustic neuroma). Genetic testing proves the patient has NF2. Report the acoustic nerve neuroma.

**Rule H2** Code malignant meningioma 9530 when the diagnosis specifically states malignant/invasive.

*Note:* Code the behavior designated in the pathology report. Do not code malignant /3 based on invasion of brain or skull. Atypical meningioma (non-malignant) may invade contiguous structures.

**Rule H3** Code the histology when only one histology is present.

*Note 1:* Use Table 3 to code histology. New codes, terms, and synonyms are included in Table 3 and coding errors may occur if the table is not used.

*Note 2:* When the histology is not listed in Table 3, use the ICD-O and all updates.

*Note 3:* Submit a question to Ask a SEER Registrar when the histology code is not found in Table 3, ICD-O or all updates.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

Rule H4  Code the **subtype/variant** when there is a NOS and a **single subtype/variant** of that NOS such as the following:

*Note:* All tumors are malignant/invasive /3.

- Astrocytoma 9400 and a subtype/variant of astrocytoma
- CNS ganglioneuroblastoma 9490 and a subtype/variant of CNS ganglioneuroblastoma
- Embryonal carcinoma 9070 and a subtype/variant of embryonal carcinoma
- Ependymoma 9391 and a subtype/variant of ependymoma
- Glioblastoma 9440 and a subtype/variant of glioblastoma
- Immature teratoma 9080 and a subtype/variant of immature teratoma
- Malignant meningioma 9530 and a subtype/variant of malignant meningioma
- Malignant peripheral nerve sheath tumor 9540 and a subtype/variant of malignant peripheral nerve sheath tumor
- Medulloblastoma 9470 and a subtype/variant of medulloblastoma
- Meningeal melanoma 8720 and a subtype/variant of meningeal melanoma
- Oligodendroglioma 9450 and a subtype/variant of oligodendroglioma
- Pilocytic astrocytoma 9421 and a subtype/variant of pilocytic astrocytoma
- Pineal parenchymal tumor of intermediate differentiation 9362 and a subtype/variant of pineal parenchymal tumor of intermediate differentiation
- Sarcoma 8800 and a subtype/variant of sarcoma

*Note:* See Table 3 in the Equivalent Terms and Definitions to find NOS and subtypes/variants.

This is the end of instructions for Single Tumor

Multiple Tumors Abstracted as a Single Primary

Rule H5  Code malignant meningioma 9530 when the diagnosis specifically states malignant/invasive.

*Note:* Code the behavior designated in the pathology report. Do not code malignant /3 based on invasion of brain or skull. Atypical meningioma (non-malignant) may invade contiguous structures.
Malignant CNS and Peripheral Nerves Histology Rules
C470-C479, C700, C701, C709, C710-C719, C720-C725, C728, C729, C751-C753
(Excludes lymphoma and leukemia M9590 – M9992 and Kaposi sarcoma M9140)

**Rule H6**
Code the histology when only one histology is present in all tumors.

*Note 1:* Use Table 3 to code histology. New codes, terms, and synonyms are included in Table 3 and coding errors may occur if the table is not used.

*Note 2:* When the histology is not listed in Table 3, use the ICD-O and all updates.

*Note 3:* Submit a question to Ask a SEER Registrar when the histology code is not found in Table 3, ICD-O or all updates.

**Rule H7**
Code the subtype/variant when all tumors are a NOS and a single subtype/variant of that NOS such as the following:

*Note:* All tumors are malignant/invasive /3.

- Astrocytoma 9400 and a subtype/variant of astrocytoma
- CNS ganglioneuroblastoma 9490 and a subtype/variant of CNS ganglioneuroblastoma
- Embryonal carcinoma 9070 and a subtype/variant of embryonal carcinoma
- Ependymoma 9391 and a subtype/variant of ependymoma
- Glioblastoma 9440 and a subtype/variant of glioblastoma
- Immature teratoma 9080 and a subtype/variant of immature teratoma
- Malignant meningioma 9530 and a subtype/variant of malignant meningioma
- Malignant peripheral nerve sheath tumor 9540 and a subtype/variant of malignant peripheral nerve sheath tumor
- Medulloblastoma 9470 and a subtype/variant of medulloblastoma
- Meningeal melanoma 8720 and a subtype/variant of meningeal melanoma
- Oligodendroglioma 9450 and a subtype/variant of oligodendroglioma
- Pilocytic astrocytoma 9421 and a subtype/variant of pilocytic astrocytoma
- Pineal parenchymal tumor of intermediate differentiation 9362 and a subtype/variant of pineal parenchymal tumor of intermediate differentiation
- Sarcoma 8800 and a subtype/variant of sarcoma

*Note:* See Table 3 in the Equivalent Terms and Definitions to find NOS and subtypes/variants.

This is the end of instructions for Multiple Tumors Abstracted as a Single Primary

Code the histology using the rule that fits the case.