Protocol for the Examination of Specimens From Patients With Retinoblastoma

Protocol applies to retinoblastoma only.

Based on AJCC/UICC TNM, 7th edition
Protocol web posting date: October 2013

Procedure
• Enucleation, Partial or Complete Exenteration

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CAP Retinoblastoma Protocol Revision History

Version Code
The definition of the version code can be found at www.cap.org/cancerprotocols.

Version: Retinoblastoma 3.1.0.0

Summary of Changes
The following changes have been made since the February 2011 release.

Enucleation, Partial or Complete Exenteration

Tumor Basal Area on Transillumination
The word “Size” was changed to “Area.”
The second dimension was deleted from “Anterior-posterior length” and “Transverse length” as follows:

Tumor Basal Area on Transillumination
___ Cannot be determined
Anterior-posterior length: ___ mm
Transverse length: ___ mm

Explanatory Notes

G. Resectioning the Globe
The following sentence and Figure 3 were added:
Each calotte should also be sampled. The calottes should be breadloafed and submitted in a cassette on edge for processing as shown in Figure 3.

References
Reference #19 was added.
Surgical Pathology Cancer Case Summary

Protocol web posting date: October 2013

RETINOBLASTOMA: Enucleation, Partial or Complete Exenteration (Notes A, B, C)

Select a single response unless otherwise indicated.

Procedure
___ Enucleation
___ Partial exenteration
___ Complete exenteration
___ Other (specify): ____________________________
___ Not specified

Specimen Size

For Enucleation
Anteroposterior diameter: ___ mm
Horizontal diameter: ___ mm
Vertical diameter: ___ mm
Length of optic nerve: ___ mm
Diameter of optic nerve: ___ mm
___ Cannot be determined (see Comment)

For Exenteration
Greatest dimension: ___ cm
+ Additional dimensions: ___ x ___ cm
___ Cannot be determined (see Comment)

Specimen Laterality
___ Right
___ Left
___ Not specified

Tumor Site (macroscopic examination/transillumination) (select all that apply) (Notes D, E)
___ Cannot be determined
___ Superotemporal quadrant of globe
___ Supronasal quadrant of globe
___ Inferotemporal quadrant of globe
___ Inferonasal quadrant of globe
___ Other (specify): ____________________________

Tumor Basal Area on Transillumination
___ Cannot be determined
Anterior-posterior length: ___ mm
Transverse length: ___ mm

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Tumor Size After Sectioning (Note F)
___ Cannot be determined
Base at cut edge: ___ mm
Height at cut edge: ___ mm
Greatest height: ___ mm

Tumor Location After Sectioning (Notes G, H)
___ Cannot be determined
Distance from anterior edge of tumor to limbus at cut edge: ___ mm
Distance of posterior margin of tumor base from edge of optic disc: ___ mm

Tumor Involvement of Other Ocular Structures (select all that apply) (Note I)
___ Cannot be determined
___ Cornea
___ Anterior chamber
___ Iris
___ Angle
___ Lens
___ Ciliary body
___ Vitreous
___ Retinal detachment
___ Optic disc
___ Choroid, minimal (solid tumor nest less than 3 mm in maximum diameter [width or thickness])
___ Choroid, massive (solid tumor nest 3 mm or more in maximum diameter [width or thickness])
___ Sclera
___ Vortex vein
___ Orbit

Histologic Features (select all that apply) (Notes J, K)
___ Cannot be determined
___ Undifferentiated
___ Differentiated
   + ___ Homer Wright rosettes
   + ___ Flexner-Wintersteiner rosettes
   + ___ Fleurettes
___ Necrotic

Growth Pattern (Note L)
___ Cannot be determined
___ Endophytic
___ Exophytic
___ Combined endophytic/exophytic
___ Diffuse

Extent of Optic Nerve Invasion
___ Cannot be determined
___ None
___ Anterior to lamina cribrosa
___ At lamina cribrosa
___ Posterior to lamina cribrosa but not to end of nerve
___ To cut end of optic nerve

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Histologic Grade
___ pGX: Grade cannot be assessed
___ pG1: Well differentiated
___ pG2: Moderately differentiated
___ pG3: Poorly differentiated
___ pG4: Undifferentiated

Margins (select all that apply)
___ Cannot be assessed
___ No tumor at margins
___ Tumor present at surgical margin of optic nerve
___ Extrascleral extension (for enucleation specimens)
___ Other margin(s) involved (specify): ________________________

Pathologic Staging (pTNM) (Note M)

TNM Descriptors (required only if applicable) (select all that apply)
___ m (multiple primary tumors)
___ r (recurrent)
___ y (post-treatment)

Primary Tumor (pT)
___ pTX: Primary tumor cannot be assessed
___ pT0: No evidence of primary tumor
___ pT1: Tumor confined to the eye with no optic nerve or choroidal invasion
___ pT2: Tumor with minimal optic nerve and/or choroidal invasion:
   ___ pT2a: Tumor superficially invades optic nerve head but does not extend past lamina cribrosa or
             tumor exhibits focal choroidal invasion
   ___ pT2b: Tumor superficially invades optic nerve head but does not extend past lamina cribrosa and
             exhibits focal choroidal invasion
___ pT3: Tumor with significant optic nerve and/or choroidal invasion:
   ___ pT3a: Tumor invades optic nerve past lamina cribrosa but not to surgical resection line or tumor
             exhibits massive choroidal invasion
   ___ pT3b: Tumor invades optic nerve past lamina cribrosa but not to surgical resection line and exhibits
             massive choroidal invasion
___ pT4: Tumor invades optic nerve to resection line or exhibits extra-ocular extension elsewhere:
   ___ pT4a: Tumor invades optic nerve to resection line but no extra-ocular extension identified
   ___ pT4b: Tumor invades optic nerve to resection line and extra-ocular extension identified

Regional Lymph Nodes (pN)
___ pNX: Regional lymph nodes cannot be assessed
___ pN0: No regional lymph node involvement
___ pN1: Regional lymph node involvement (preauricular, cervical, submandibular)
___ pN2: Distant lymph node involvement

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Distant Metastasis (pM)
___ Not applicable
___ pM1: Metastasis to sites other than CNS
___ pM1a: Single lesion
___ pM1b: Multiple lesions
___ pM1c: CNS metastasis
___ pM1d: Discrete mass(es) without leptomeningeal and/or CSF involvement
___ pM1e: Leptomeningeal and/or CSF involvement

+ Additional Pathologic Findings (select all that apply)
+ ___ None identified
+ ___ Calcifications
+ ___ Mitotic rate: Number of mitoses per 40x objective with a field area of 0.152 mm² (specify): ___
+ ___ Apoptosis
+ ___ Basophilic vascular deposits
+ ___ Inflammatory cells
+ ___ Hemorrhage
+ ___ Neovascularization (specify site): ________________________
+ ___ Other (specify): __________________________

+ Comment(s)
Explanatory Notes

A. Cytology/Biopsy
Cytologic and biopsy specimens are rarely obtained from eyes with suspected retinoblastoma owing to the potential risk of tumor seeding. An anterior chamber paracentesis may be performed, if indicated by clinical findings, and is not associated with risk of tumor seeding.\textsuperscript{1,2}

B. Fixation
The minimum recommended fixation time for whole globes with intraocular tumors is 48 hours. The globe should be fixed in an adequate volume of fixative with a 10:1 ratio of fixative volume to specimen volume recommended. Incisions or windows in the globe are not necessary for adequate penetration of fixative and are not recommended. Injection of fixative into the globe is also not recommended.

C. Additional Studies
Genetic studies may be requested on neoplastic tissue and should be harvested prior to fixation.\textsuperscript{3} The surgical margin of the optic nerve should be obtained prior to opening the globe (Note E). Once tissue is harvested for genetic studies, the globe can be fixed prior to completing macroscopic examination. The appropriate materials/medium required by the laboratory performing genetic testing should be obtained prior to the procedure.

D. Processing With Tumor Sampling:
To collect the tumor specimen, the optic nerve should be removed before opening the globe to prevent the optic nerve from accidentally becoming contaminated with artifactual clumps of tumor cells (so-called “floaters”). The surgeon should first ink the surgical margin of the optic nerve, then cut the optic nerve stump off from the sclera with a sharp razor about 2 mm behind the globe. The optic nerve stump, which should be kept separate from the globe, should be placed into a jar of 10% buffered formaldehyde. Then, a sample of tumor should be obtained by opening a small sclero-choroidal window adjacent to the tumor near the equator with a 6- to 8-mm corneal trephine. Once the opening into the vitreous chamber is established, tumor tissue should be gently removed with forceps and scissors. It is best to leave a hinge on 1 side of the scleral flap so that it can be closed with 1 or 2 suture(s) following the removal of tumor sample. This is done in an attempt to maintain the overall spherical architecture of the specimen during fixation. The globe should be placed in a second jar of formalin (separate from the optic nerve stump) and be allowed to fix for at least 24 to 48 hours.

E. Orientation of Globe
The orientation of a globe may be determined by identifying extraocular muscle insertions, optic nerve and other landmarks as illustrated in Figure 1. The terms \textit{temporal} and \textit{nasal} are generally used in place of \textit{lateral} and \textit{medial} with reference to ocular anatomy.
Figure 1. Anatomic landmarks of the posterior aspect of the globe (right eye). The position of the inferior oblique muscle relative to the optic nerve is most helpful in orienting the globe. The inferior oblique muscle insertion is located temporal (lateral) to the optic nerve on the sclera, and its fibers travel inferonasally from its insertion. The long posterior ciliary artery is often seen as a blue-gray line in the sclera on either side of the optic nerve and marks the horizontal meridian of the globe. Reprinted with permission from WB Saunders Company.

F. Processing Without Tumor Sampling
If there is no need for fresh tissue sampling, the enucleated globe should simply be fixed in 10% buffered formaldehyde for at least 24 and preferably 48 hours. When the fixed globe is examined by the pathologist, if the optic nerve was not previously amputated in the operative room, that should be performed first as described above. The surgical margin of the nerve stump should be embedded face down in paraffin for sectioning (ie, thereby obtaining cross-sections of the nerve, starting at the surgical margin). Then, the eye itself is sectioned. First, a section should be made that extends from pupil through the optic nerve (the “P-O” section), which contains the center of the optic nerve with all the optic nerve structures (optic nerve head, lamina cribrosa, and postlaminar optic nerve). Preferably this plane should bisect the largest dimension of the tumor, previously identified by transillumination and during clinical examination. When possible, the plane should avoid the scleral opening if one was made for fresh tumor sampling. This section is critical for evaluation of the optic nerve for tumor invasion. The P-O section and minor calottes are then embedded in paraffin. The embedded P-O calotte is then sectioned every 100 to 150 microns (each section being about 5 microns thick), for a total of about 10 to 20 sections. Additional sections should also be made anterior-posteriorly in a bread-loaf fashion through the minor calottes. These segments should be submitted in 1 cassette per calotte on edge to evaluate the choroid for invasion. Three levels of this block are usually sufficient for examination. In total, 4 cassettes are submitted: the optic nerve stump, the P-O section, and the 2 minor calottes (unless 1 or both of these has no visible tumor).

G. Sectioning the Globe
The globe is generally sectioned in the horizontal or vertical plane, with care to include the pupil and optic nerve in the cassette to be submitted for microscopic examination. If the mass cannot be included with horizontal or vertical sectioning, the globe is sectioned obliquely to include tumor, pupil, and optic nerve (Figure 2). The surgical margin of the optic nerve should be sectioned and submitted
prior to sectioning the globe to ensure that intraocular malignant cells do not contaminate this important surgical margin. Retinoblastoma is an extremely friable tumor.

Figure 2. The most common methods of sectioning a globe. After transillumination, the tumor base is marked, if possible, and included in the pupil-optic (p-o) nerve section and submitted for processing. If tumor is found in either of the calottes, these may also be submitted for sectioning. The meridian in which the globe was sectioned should be included in the gross description of the pathology report. It is not uncommon to induce an artifactitious retinal detachment while sectioning the globe. This can be minimized by gentle handling and by avoiding a sawing motion with the blade. Reprinted with permission from WB Saunders Company.
Each calotte should also be sampled. The calottes should be breadloafed and submitted in a cassette on edge for processing as shown in Figure 3.

Figure 3. Calotte sampling. From Grossniklaus HE. Reproduced with kind permission of Springer Science+Business Media.

H. Sections Submitted for Microscopic Examination
Multiple sections should be examined, with special attention to sections containing optic nerve and tumor. The nerve should be sectioned along the various levels to determine tumor extension.

I. Rules for Classification

Choroidal Invasion: The presence and the extent (focal versus massive) of choroidal invasion by tumor should be stated. Differentiation should be made between true choroidal invasion and artifactual invasion due to seeding of fresh tumor cells during post-enucleation retrieval of tumor tissue and/or gross sectioning.

Artifactual invasion is identified when there are groups of tumor cells present in the open spaces between intraocular structures, extraocular tissues and/or subarachnoid space.

True invasion is defined as 1 or more solid nests of tumor cells that fills or replaces the choroid and has pushing borders. Note: Invasion of the sub-retinal pigment epithelium (RPE) space, where tumor cells are present under the RPE (but not beyond Bruch’s membrane into the choroid) is not choroidal invasion.

Focal choroidal invasion is defined as a solid nest of tumor that measures less than 3 mm in maximum diameter (width or thickness).

Massive choroidal invasion is defined as a solid tumor nest 3 mm or more in maximum diameter (width or thickness) in contact with the underlying sclera.

J. Histologic Features
Typical histologic features include cells with large, basophilic nuclei and scant cytoplasm. Mitoses are generally frequent. Calcification and necrosis are common with sleeves of viable cells typically surrounding blood vessels (pseudorosettes). Apoptotic cells may be seen. The extent of differentiation may be judged based on the presence and type of rosettes. Homer-Wright rosettes similar to those seen in neuroblastoma or medulloblastoma may be seen and are not a sign of significant differentiation. Flexner-Wintersteiner rosettes are evidence of higher differentiation. Fleurettes are considered the most differentiated form of rosette found in the tumor. A benign variant of retinoblastoma termed retinocytoma or retinoma has been described. This tumor consists entirely of benign, well-differentiated...
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cells often with associated calcification. The cells have smaller, less hyperchromatic nuclei and abundant cytoplasm. Necrosis is typically absent and mitotic figures are rare. Retinoblastomas may arise in multicentric foci.

K. Histologic Features of Additional Prognostic Significance
Histologic features with prognostic significance for survival include the following: invasion of optic nerve, particularly if tumor is present at the surgical margin (most important feature); invasion of sclera; invasion of choroid; tumor size; basophilic staining of tumor vessels; seeding of vitreous; degree of differentiation; involvement of anterior segment; and growth pattern. This list should not be confused with the Reese-Ellsworth classification, which is intended as a predictor for visual outcome, not survival.

L. Growth Pattern
Endophytic growth pattern indicates growth from the inner retinal surface into the vitreous cavity. Exophytic tumors grow primarily from the outer surface of the retina into the subretinal space toward the choroid. Mixed growth pattern exhibits features of both endophytic and exophytic growth. Diffuse infiltrating tumors grow laterally within the retina without significant thickening.

M. TNM and Stage Groupings
The American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC) TNM staging system for retinoblastoma is shown below.

By AJCC/UICC convention, the designation “T” refers to a primary tumor that has not been previously treated. The symbol “p” refers to the pathologic classification of the TNM, as opposed to the clinical classification, and is based on gross and microscopic examination. pT entails a resection of the primary tumor or biopsy adequate to evaluate the highest pT category, pN entails removal of nodes adequate to validate lymph node metastasis, and pM implies microscopic examination of distant lesions. Clinical classification (cTNM) is usually carried out by the referring physician before treatment during initial evaluation of the patient or when pathologic classification is not possible.

Pathologic staging is usually performed after surgical resection of the primary tumor. Pathologic staging depends on pathologic documentation of the anatomic extent of disease, whether or not the primary tumor has been completely removed. If a biopsied tumor is not resected for any reason (eg, when technically unfeasible) and if the highest T and N categories or the M1 category of the tumor can be confirmed microscopically, the criteria for pathologic classification and staging have been satisfied without total removal of the primary cancer.

Clinical TNM Classifications:
Primary Tumor (T)

| TX | Primary tumor cannot be assessed. |
| T0 | No evidence of primary tumor. |
| T1 | Tumors no more than 2/3 the volume of the eye with no vitreous or subretinal seeding. |
| T1a | No tumor in either eye is greater than 3 mm in largest dimension or located closer than 1.5 mm to the optic nerve or fovea. |
| T1b | At least one tumor is greater than 3 mm in largest dimension or located closer than 1.5 mm to the optic nerve or fovea. No retinal detachment or subretinal fluid beyond 5 mm from the base of the tumor. |
| T1c | At least one tumor is greater than 3 mm in largest dimension or located closer than 1.5 mm to the optic nerve or fovea. With retinal detachment or subretinal fluid beyond 5 mm from the base of the tumor. |
| T2 | Tumors no more than 2/3 the volume of the eye with vitreous or subretinal seeding. Can have retinal detachment. |
T2a  Focal vitreous and/or subretinal seeding of fine aggregates of tumor cells is present, but no large clumps or “snowballs” of tumor cells.
T2b  Massive vitreous and/or subretinal seeding is present, defined as diffuse clumps or “snowballs” of tumor cells.
T3  Severe intraocular disease
T3a  Tumor fills more than 2/3 of the eye
T3b  One or more complications present, which may include tumor-associated neovascular or angle closure glaucoma, tumor extension into the anterior segment, hyphema, vitreous hemorrhage, or orbital cellulitis.
T4  Extraocular disease detected by imaging studies.
T4a  Invasion of optic nerve.
T4b  Invasion into the orbit.
T4c  Intracranial extension not past chiasm.
T4d  Intracranial extension past chiasm.

Regional Lymph Nodes (N)
NX  Regional lymph nodes cannot be assessed
N0  No regional lymph node involvement
N1  Regional lymph node involvement (preauricular, cervical, submandibular)
N2  Distant lymph node involvement

Metastasis (M)
M0  No metastasis
M1  Systemic metastasis
M1a  Single lesion to sites other than CNS
M1b  Multiple lesions to sites other than CNS
M1c  Prechiasmatic CNS lesion(s)
M1d  Postchiasmatic CNS lesion(s)
M1e  Leptomeningeal and/or CSF involvement

Pathologic Classification (pTNM):
Primary Tumor (pT)
pTX  Primary tumor cannot be assessed.
pT0  No evidence of primary tumor.
pT1  Tumor confined to eye with no optic nerve or choroidal invasion.
pT2  Tumor with minimal optic nerve and/or choroidal invasion:
pT2a  Tumor superficially invades optic nerve head but does not extend past lamina cribrosa or tumor exhibits focal choroidal invasion.
pT2b  Tumor superficially invades optic nerve head but does not extend past lamina cribrosa and exhibits focal choroidal invasion.
pT3  Tumor with significant optic nerve and/or choroidal invasion:
pT3a  Tumor invades optic nerve past lamina cribrosa but not to surgical resection line or tumor exhibits massive choroidal invasion.
pT3b  Tumor invades optic nerve past lamina cribrosa but not to surgical resection line and exhibits massive choroidal invasion.
pT4  Tumor invades optic nerve to resection line or exhibits extra-ocular extension elsewhere.
pT4a  Tumor invades optic nerve to resection line but no extra-ocular extension identified.
pT4b  Tumor invades optic nerve to resection line and extra-ocular extension identified.

Regional Lymph Nodes (pN)
pNX  Regional lymph nodes cannot be assessed
pN0  No regional lymph node involvement
pN1  Regional lymph node involvement (preauricular, cervical)
pN2  Distant lymph node involvement

**Metastasis (pM)**
- pM0  No metastasis
- pM1  Metastasis to sites other than CNS
  - pM1a  Single lesion
  - pM1b  Multiple lesions
  - pM1c  CNS metastasis
  - pM1d  Discrete mass(es) without leptomeningeal and/or CSF involvement
  - pM1e  Leptomeningeal and/or CSF involvement

**TNM Stage Groupings**
No stage grouping applies.

**TNM Descriptors**
For identification of special cases of TNM or pTNM classifications, the “m” suffix and “y,” “r,” and “a” prefixes are used. Although they do not affect the stage grouping, they indicate cases needing separate analysis.

The “m” suffix indicates the presence of multiple primary tumors in a single site and is recorded in parentheses: pT(m)NM.

The “y” prefix indicates those cases in which classification is performed during or following initial multimodality therapy (ie, neoadjuvant chemotherapy, radiation therapy, or both chemotherapy and radiation therapy). The cTNM or pTNM category is identified by a “y” prefix. The ycTNM or ypTNM categorizes the extent of tumor actually present at the time of that examination. The “y” categorization is not an estimate of tumor prior to multimodality therapy (ie, before initiation of neoadjuvant therapy).

The “r” prefix indicates a recurrent tumor when staged after a documented disease-free interval, and is identified by the “r” prefix: rTNM.

The “a” prefix designates the stage determined at autopsy: aTNM.

**Additional Descriptors**

**Residual Tumor (R)**
Tumor remaining in a patient after therapy with curative intent (eg, surgical resection for cure) is categorized by a system known as R classification, shown below.

- RX  Presence of residual tumor cannot be assessed
- R0  No residual tumor
- R1  Microscopic residual tumor
- R2  Macroscopic residual tumor

For the surgeon, the R classification may be useful to indicate the known or assumed status of the completeness of a surgical excision. For the pathologist, the R classification is relevant to the status of the margins of a surgical resection specimen. That is, tumor involving the resection margin on pathologic examination may be assumed to correspond to residual tumor in the patient and may be classified as macroscopic or microscopic according to the findings at the specimen margin(s).
Lymph-Vascular Invasion (LVI)
LVI indicates whether microscopic lymph-vascular invasion is identified in the pathology report. LVI includes lymphatic invasion, vascular invasion, or lymph-vascular invasion. By AJCC/UICC convention, LVI does not affect the T category indicating local extent of tumor unless specifically included in the definition of a T category.

References

Bibliography
