

MALIGNANT MELANOMA OF THE UVEA STAGING FORM

CLINICAL Extent of disease before any treatment	STAGE CATEGORY DEFINITIONS		PATHOLOGIC Extent of disease through completion of definitive surgery
<input type="checkbox"/> y clinical – staging completed after neoadjuvant therapy but before subsequent surgery	TUMOR SIZE: _____	LATERALITY: <input type="checkbox"/> left <input type="checkbox"/> right <input type="checkbox"/> bilateral	<input type="checkbox"/> y pathologic – staging completed after neoadjuvant therapy AND subsequent surgery
<p style="text-align: center;">PRIMARY TUMOR (T)</p> <p>All Uveal Melanomas</p> <p>Primary tumor cannot be assessed No evidence of primary tumor</p> <p>Iris*</p> <p><input type="checkbox"/> TX <input type="checkbox"/> T0</p> <p><input type="checkbox"/> T1 <input type="checkbox"/> T1a <input type="checkbox"/> T1b <input type="checkbox"/> T1c <input type="checkbox"/> T2 <input type="checkbox"/> T2a</p> <p><input type="checkbox"/> T3 <input type="checkbox"/> T3a</p> <p><input type="checkbox"/> T4 <input type="checkbox"/> T4a <input type="checkbox"/> T4b</p> <p>* Iris melanomas originate from, and are predominantly located in, this region of the uvea. If less than half of the tumor volume is located within the iris, the tumor may have originated in the ciliary body and consideration should be given to classifying it accordingly.</p> <p>Ciliary Body and Choroid (see Figure on p. 550)</p> <p>Primary ciliary body and choroidal melanomas are classified according to the four tumor size categories below:</p> <p><input type="checkbox"/> T1 <input type="checkbox"/> T1a</p> <p><input type="checkbox"/> T1b <input type="checkbox"/> T1c <input type="checkbox"/> T1d</p> <p><input type="checkbox"/> T2 <input type="checkbox"/> T2a <input type="checkbox"/> T2b <input type="checkbox"/> T2c <input type="checkbox"/> T2d</p> <p><input type="checkbox"/> T3 <input type="checkbox"/> T3a</p> <p><input type="checkbox"/> T1 <input type="checkbox"/> T1a</p> <p><input type="checkbox"/> T1b <input type="checkbox"/> T1c</p> <p><input type="checkbox"/> T1d</p> <p><input type="checkbox"/> T2 <input type="checkbox"/> T2a</p> <p><input type="checkbox"/> T2b <input type="checkbox"/> T2c</p> <p><input type="checkbox"/> T2d</p> <p><input type="checkbox"/> T3 <input type="checkbox"/> T3a</p>			

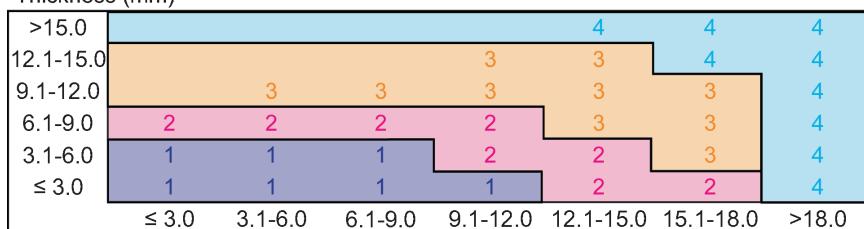
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<input type="checkbox"/> T3b <input type="checkbox"/> T3c <input type="checkbox"/> T3d <input type="checkbox"/> T4 <input type="checkbox"/> T4a <input type="checkbox"/> T4b <input type="checkbox"/> T4c <input type="checkbox"/> T4d <input type="checkbox"/> T4e	<p>Tumor size category 3 with ciliary body involvement Tumor size category 3 without ciliary body involvement but with extraocular extension less than or equal to 5 mm in diameter Tumor size category 3 with ciliary body involvement and extraocular extension less than or equal to 5 mm in diameter Tumor size category 4 Tumor size category 4 without ciliary body involvement and extraocular extension Tumor size category 4 with ciliary body involvement Tumor size category 4 without ciliary body involvement but with extraocular extension less than or equal to 5 mm in diameter Tumor size category 4 with ciliary body involvement and extraocular extension less than or equal to 5 mm in diameter Any tumor size category with extraocular extension more than 5 mm in diameter</p> <p>*Clinical: In clinical practice, the largest tumor basal diameter may be estimated in optic disc diameters (dd, average: 1 dd = 1.5 mm). Tumor thickness may be estimated in diopters (average: 2.5 diopters = 1 mm). However, techniques such as ultrasonography and fundus photography are used to provide more accurate measurements. Ciliary body involvement can be evaluated by the slit-lamp, ophthalmoscopy, gonioscopy and transillumination. However, high frequency ultrasonography (ultrasound biomicroscopy) is used for more accurate assessment. Extension through the sclera is evaluated visually before and during surgery, and with ultrasonography, computed tomography or magnetic resonance imaging.</p> <p>[†]Pathologic: When histopathologic measurements are recorded after fixation, tumor diameter and thickness may be underestimated because of tissue shrinkage.</p>	<input type="checkbox"/> T3b <input type="checkbox"/> T3c <input type="checkbox"/> T3d <input type="checkbox"/> T4 <input type="checkbox"/> T4a <input type="checkbox"/> T4b <input type="checkbox"/> T4c <input type="checkbox"/> T4d <input type="checkbox"/> T4e
<input type="checkbox"/> NX <input type="checkbox"/> N0 <input type="checkbox"/> N1	REGIONAL LYMPH NODES (N) Regional lymph nodes cannot be assessed No regional lymph node metastasis Regional lymph node metastasis	<input type="checkbox"/> NX <input type="checkbox"/> N0 <input type="checkbox"/> N1
<input type="checkbox"/> M0 <input type="checkbox"/> M1 <input type="checkbox"/> M1a <input type="checkbox"/> M1b <input type="checkbox"/> M1c	DISTANT METASTASIS (M) No distant metastasis (no pathologic M0; use clinical M to complete stage group) Distant metastasis Largest diameter of the largest metastasis ≤3 cm Largest diameter of the largest metastasis 3.1-8.0 cm Largest diameter of the largest metastasis 8.1 cm or more	<input type="checkbox"/> M1 <input type="checkbox"/> M1a <input type="checkbox"/> M1b <input type="checkbox"/> M1c

Thickness (mm)



Largest basal diameter (mm)

Classification for ciliary body and choroid uveal melanoma based on thickness and diameter.

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ANATOMIC STAGE • PROGNOSTIC GROUPING

CLINICAL				PATHOLOGIC			
GROUP	T	N	M	GROUP	T	N	M
<input type="checkbox"/> I	T1a	N0	M0	<input type="checkbox"/> I	T1a	N0	M0
<input type="checkbox"/> IIA	T1b-d	N0	M0	<input type="checkbox"/> IIA	T1b-d	N0	M0
	T2a	N0	M0		T2a	N0	M0
<input type="checkbox"/> IIB	T2b	N0	M0	<input type="checkbox"/> IIB	T2b	N0	M0
	T3a	N0	M0		T3a	N0	M0
<input type="checkbox"/> IIIA	T2c-d	N0	M0	<input type="checkbox"/> IIIA	T2c-d	N0	M0
	T3b-c	N0	M0		T3b-c	N0	M0
	T4a	N0	M0		T4a	N0	M0
<input type="checkbox"/> IIIB	T3d	N0	M0	<input type="checkbox"/> IIIB	T3d	N0	M0
	T4b-c	N0	M0		T4b-c	N0	M0
<input type="checkbox"/> IIIC	T4d-e	N0	M0	<input type="checkbox"/> IIIC	T4d-e	N0	M0
<input type="checkbox"/> IV	Any T	N1	M0	<input type="checkbox"/> IV	Any T	N1	M0
	Any T	Any N	M1a-c		Any T	Any N	M1a-c
<input type="checkbox"/> Stage unknown				<input type="checkbox"/> Stage unknown			
PROGNOSTIC FACTORS (SITE-SPECIFIC FACTORS)							
REQUIRED FOR STAGING: Tumor height and largest diameter _____							
CLINICALLY SIGNIFICANT:							
Measured thickness (depth) _____							
Chromosomal alterations _____							
Gene expression profile _____							
Positron emission tomography/computed tomography _____							
Confocal indocyanine green angiography _____							
Mitotic count per 40 high power fields (HPF) _____							
Mean diameter of the ten largest nucleoli (MLN) _____							
Presence of extravascular matrix patterns _____							
Microvascular density (MVD) _____							
Insulin-like growth factor 1 receptor (IGF1-R) _____							
Tumor-infiltrating lymphocytes _____							
Tumor-infiltrating macrophages _____							
HLA Class I expression _____							
Histologic Grade (G) (also known as overall grade)							
Grading system				Grade			
<input type="checkbox"/>	2 grade system	<input type="checkbox"/>	Grade I or 1				
<input type="checkbox"/>	3 grade system	<input type="checkbox"/>	Grade II or 2				
<input type="checkbox"/>	4 grade system	<input type="checkbox"/>	Grade III or 3				
<input type="checkbox"/>	No 2, 3, or 4 grade system is available	<input type="checkbox"/>	Grade IV or 4				

General Notes:
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.

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

y prefix indicates those cases in which classification is performed during or following initial multimodality 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.

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

a prefix designates the stage determined at autopsy: aTNM.

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ADDITIONAL DESCRIPTORS

Lymphatic Vessel Invasion (L) and Venous Invasion (V) have been combined into Lymph-Vascular Invasion (LVI) for collection by cancer registrars. The College of American Pathologists' (CAP) Checklist should be used as the primary source. Other sources may be used in the absence of a Checklist. Priority is given to positive results.

- Lymph-Vascular Invasion Not Present (absent)/Not Identified
- Lymph-Vascular Invasion Present/Identified
- Not Applicable
- Unknown/Indeterminate

Residual Tumor (R)

The absence or presence of residual tumor after treatment. In some cases treated with surgery and/or with neoadjuvant therapy there will be residual tumor at the primary site after treatment because of incomplete resection or local and regional disease that extends beyond the limit of ability of resection.

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

General Notes (continued):

surgical margins is data field recorded by registrars describing the surgical margins of the resected primary site specimen as determined only by the pathology report.

neoadjuvant treatment is radiation therapy or systemic therapy (consisting of chemotherapy, hormone therapy, or immunotherapy) administered prior to a definitive surgical procedure. If the surgical procedure is not performed, the administered therapy no longer meets the definition of neoadjuvant therapy.

- Clinical stage was used in treatment planning (describe): _____
- National guidelines were used in treatment planning NCCN Other (describe): _____
-
-
-

Physician signature

Date/Time

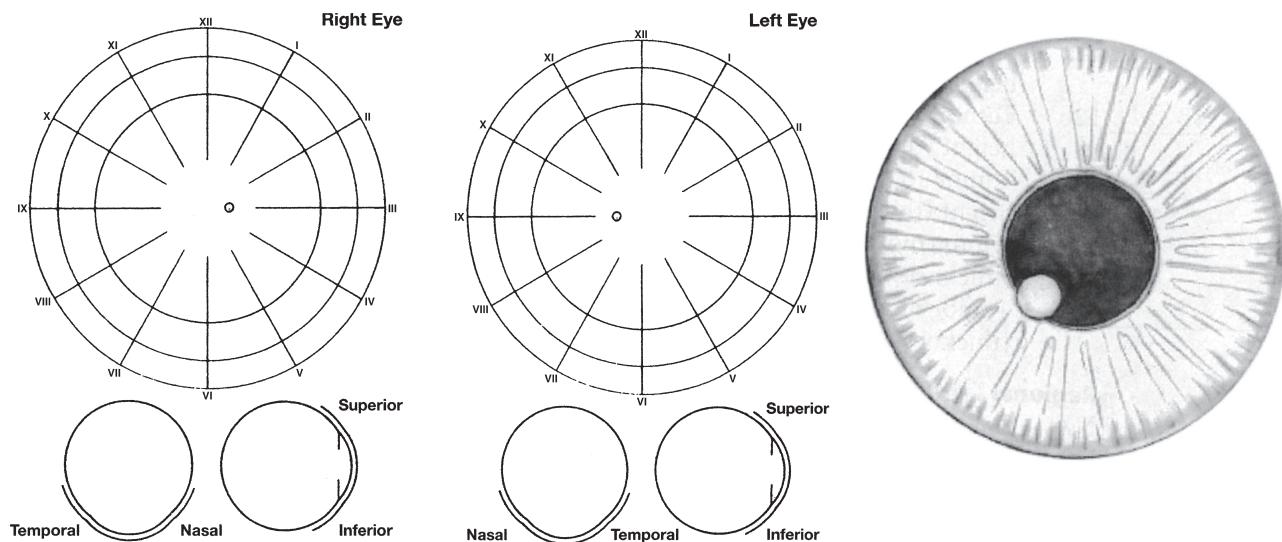
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Illustration

Indicate on diagram primary tumor and regional nodes involved.



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