

COLON AND RECTUM STAGING FORM

CLINICAL <i>Extent of disease before any treatment</i>	STAGE CATEGORY DEFINITIONS	PATHOLOGIC <i>Extent of disease through completion of definitive surgery</i>																		
<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																		
<p style="text-align: center;">PRIMARY TUMOR (T)</p> <table style="margin-left: auto; margin-right: auto;"> <tr><td><input type="checkbox"/> TX</td><td>Primary tumor cannot be assessed</td></tr> <tr><td><input type="checkbox"/> T0</td><td>No evidence of primary tumor</td></tr> <tr><td><input type="checkbox"/> Tis</td><td>Carcinoma <i>in situ</i>: intraepithelial or invasion of lamina propria*</td></tr> <tr><td><input type="checkbox"/> T1</td><td>Tumor invades submucosa</td></tr> <tr><td><input type="checkbox"/> T2</td><td>Tumor invades muscularis propria</td></tr> <tr><td><input type="checkbox"/> T3</td><td>Tumor invades through the muscularis propria into pericolorectal tissues</td></tr> <tr><td><input type="checkbox"/> T4a</td><td>Tumor penetrates to the surface of the visceral peritoneum**</td></tr> <tr><td><input type="checkbox"/> T4b</td><td>Tumor directly invades or is adherent to other organs or structures**,***</td></tr> </table> <p>*Note: Tis includes cancer cells confined within the glandular basement membrane (intraepithelial) or mucosal lamina propria (intramucosal) with no extension through the muscularis mucosae into the submucosa.</p> <p>**Note: Direct invasion in T4 includes invasion of other organs or other segments of the colorectum as a result of direct extension through the serosa, as confirmed on microscopic examination (for example, invasion of the sigmoid colon by a carcinoma of the cecum) or, for cancers in a retro-peritoneal or subperitoneal location, direct invasion of other organs or structures by virtue of extension beyond the muscularis propria (i.e., respectively, a tumor on the posterior wall of the descending colon invading the left kidney or lateral abdominal wall; or a mid or distal rectal cancer with invasion of prostate, seminal vesicles, cervix or vagina).</p> <p>***Note: Tumor that is adherent to other organs or structures, grossly, is classified ct4b. However, if no tumor is present in the adhesion, microscopically, the classification should be pT1-4a depending on the anatomical depth of wall invasion. The V and L classifications should be used to identify the presence or absence of vascular or lymphatic invasion whereas the PN site-specific factor should be used for perineural invasion.</p>			<input type="checkbox"/> TX	Primary tumor cannot be assessed	<input type="checkbox"/> T0	No evidence of primary tumor	<input type="checkbox"/> Tis	Carcinoma <i>in situ</i> : intraepithelial or invasion of lamina propria*	<input type="checkbox"/> T1	Tumor invades submucosa	<input type="checkbox"/> T2	Tumor invades muscularis propria	<input type="checkbox"/> T3	Tumor invades through the muscularis propria into pericolorectal tissues	<input type="checkbox"/> T4a	Tumor penetrates to the surface of the visceral peritoneum**	<input type="checkbox"/> T4b	Tumor directly invades or is adherent to other organs or structures**,***		
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<input type="checkbox"/> NX <input type="checkbox"/> N0 <input type="checkbox"/> N1 <input type="checkbox"/> N1a <input type="checkbox"/> N1b <input type="checkbox"/> N1c <input type="checkbox"/> N2 <input type="checkbox"/> N2a <input type="checkbox"/> N2b	<p style="text-align: center;">REGIONAL LYMPH NODES (N)</p> <table style="margin-left: auto; margin-right: auto;"> <tr><td><input type="checkbox"/> NX</td><td>Regional lymph nodes cannot be assessed</td></tr> <tr><td><input type="checkbox"/> N0</td><td>No regional lymph node metastasis</td></tr> <tr><td><input type="checkbox"/> N1</td><td>Metastasis in 1 to 3 regional lymph nodes</td></tr> <tr><td><input type="checkbox"/> N1a</td><td>Metastasis in 1 regional lymph node</td></tr> <tr><td><input type="checkbox"/> N1b</td><td>Metastasis in 2-3 regional lymph nodes</td></tr> <tr><td><input type="checkbox"/> N1c</td><td>Tumor deposit(s) in the subserosa, mesentery, or non-peritonealized pericolic or perirectal tissues without regional nodal metastasis</td></tr> <tr><td><input type="checkbox"/> N2</td><td>Metastasis in 4 or more regional lymph nodes</td></tr> <tr><td><input type="checkbox"/> N2a</td><td>Metastasis in 4 to 6 regional lymph nodes</td></tr> <tr><td><input type="checkbox"/> N2b</td><td>Metastasis in 7 or more regional lymph nodes</td></tr> </table> <p>Note: A satellite peritumoral nodule in the pericolorectal adipose tissue of a primary carcinoma without histologic evidence of residual lymph node in the nodule may represent discontinuous spread, venous invasion with extravascular spread (V1/2) or a totally replaced lymph node (N1/2). Replaced nodes should be counted separately as positive nodes in the N category, whereas discontinuous spread or venous invasion should be classified and counted in the Site-Specific Factor category Tumor Deposits (TD).</p>	<input type="checkbox"/> NX	Regional lymph nodes cannot be assessed	<input type="checkbox"/> N0	No regional lymph node metastasis	<input type="checkbox"/> N1	Metastasis in 1 to 3 regional lymph nodes	<input type="checkbox"/> N1a	Metastasis in 1 regional lymph node	<input type="checkbox"/> N1b	Metastasis in 2-3 regional lymph nodes	<input type="checkbox"/> N1c	Tumor deposit(s) in the subserosa, mesentery, or non-peritonealized pericolic or perirectal tissues without regional nodal metastasis	<input type="checkbox"/> N2	Metastasis in 4 or more regional lymph nodes	<input type="checkbox"/> N2a	Metastasis in 4 to 6 regional lymph nodes	<input type="checkbox"/> N2b	Metastasis in 7 or more regional lymph nodes	<input type="checkbox"/> NX <input type="checkbox"/> N0 <input type="checkbox"/> N1 <input type="checkbox"/> N1a <input type="checkbox"/> N1b <input type="checkbox"/> N1c <input type="checkbox"/> N2 <input type="checkbox"/> N2a <input type="checkbox"/> N2b
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		DISTANT METASTASIS (M)		
<input type="checkbox"/>	M0	No distant metastasis (no pathologic M0; use clinical M to complete stage group)		
<input type="checkbox"/>	M1	Distant metastasis	<input type="checkbox"/>	M1
<input type="checkbox"/>	M1a	Metastasis confined to one organ or site (e.g., liver, lung, ovary, non-regional node).	<input type="checkbox"/>	M1a
<input type="checkbox"/>	M1b	Metastases in more than one organ/site or the peritoneum.	<input type="checkbox"/>	M1b

ANATOMIC STAGE • PROGNOSTIC GROUPS

CLINICAL						PATHOLOGIC					
GROUP	T	N	M	Dukes*	MAC*	GROUP	T	N	M	Dukes*	MAC*
<input type="checkbox"/> 0	Tis	N0	M0	-	-	<input type="checkbox"/> 0	Tis	N0	M0	-	-
<input type="checkbox"/> I	T1	N0	M0	A	A	<input type="checkbox"/> I	T1	N0	M0	A	A
		T2	N0	M0	A	<input type="checkbox"/> T2	N0	M0	A	A	B1
<input type="checkbox"/> IIA	T3	N0	M0	B	B2	<input type="checkbox"/> IIA	T3	N0	M0	B	B2
<input type="checkbox"/> IIB	T4a	N0	M0	B	B2	<input type="checkbox"/> IIB	T4a	N0	M0	B	B2
<input type="checkbox"/> IIC	T4b	N0	M0	B	B3	<input type="checkbox"/> IIC	T4b	N0	M0	B	B3
<input type="checkbox"/> IIIA	T1-T2	N1/N1c	M0	C	C1	<input type="checkbox"/> IIIA	T1-T2	N1/N1c	M0	C	C1
		T1	N2a	M0	C	<input type="checkbox"/> T1	N2a	M0	C	C	C1
<input type="checkbox"/> IIIB	T3-T4a	N1/N1c	M0	C	C2	<input type="checkbox"/> IIIB	T3-T4a	N1/N1c	M0	C	C2
		T2-T3	N2a	M0	C	<input type="checkbox"/> T2-T3	N2a	M0	C	C	C1/C2
		T1-T2	N2b	M0	C	<input type="checkbox"/> T1-T2	N2b	M0	C	C	C1
<input type="checkbox"/> IIIC	T4a	N2a	M0	C	C2	<input type="checkbox"/> IIIC	T4a	N2a	M0	C	C2
		T3-T4a	N2b	M0	C	<input type="checkbox"/> T3-T4a	N2b	M0	C	C	C2
		T4b	N1-N2	M0	C	<input type="checkbox"/> T4b	N1-N2	M0	C	C	C3
<input type="checkbox"/> IVA	Any T	Any N	M1a	-	-	<input type="checkbox"/> IVA	Any T	Any N	M1a	-	-
<input type="checkbox"/> IVB	Any T	Any N	M1b	-	-	<input type="checkbox"/> IVB	Any T	Any N	M1b	-	-

*Dukes B is a composite of better (T3 N0 M0) and worse (T4 N0 M0) prognostic groups, as is Dukes C (Any TN1 M0 and Any T N2 M0). MAC is the modified Astler-Coller classification.

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Stage unknown

PROGNOSTIC FACTORS (SITE-SPECIFIC FACTORS)											
REQUIRED FOR STAGING: None											
CLINICALLY SIGNIFICANT:											
Pre-operative or pre-treatment carcinoembryonic antigen (CEA) ng/ml _____											
Tumor Deposits (TD) _____											
Circumferential Resection Margin (CRM) _____											
Perineural Invasion (PN) _____											
Microsatellite Instability (MSI) _____											
Tumor Regression Grade (with neoadjuvant therapy) _____											
KRAS gene analysis _____											
18q loss of heterozygosity (LOH) assay _____											
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"/> 3 grade system			<input type="checkbox"/> Grade III or 3			<input type="checkbox"/> 4 grade system			<input type="checkbox"/> Grade IV or 4		
<input type="checkbox"/> No 2, 3, or 4 grade system is available											

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.

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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):

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.

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

HOSPITAL NAME/ADDRESS

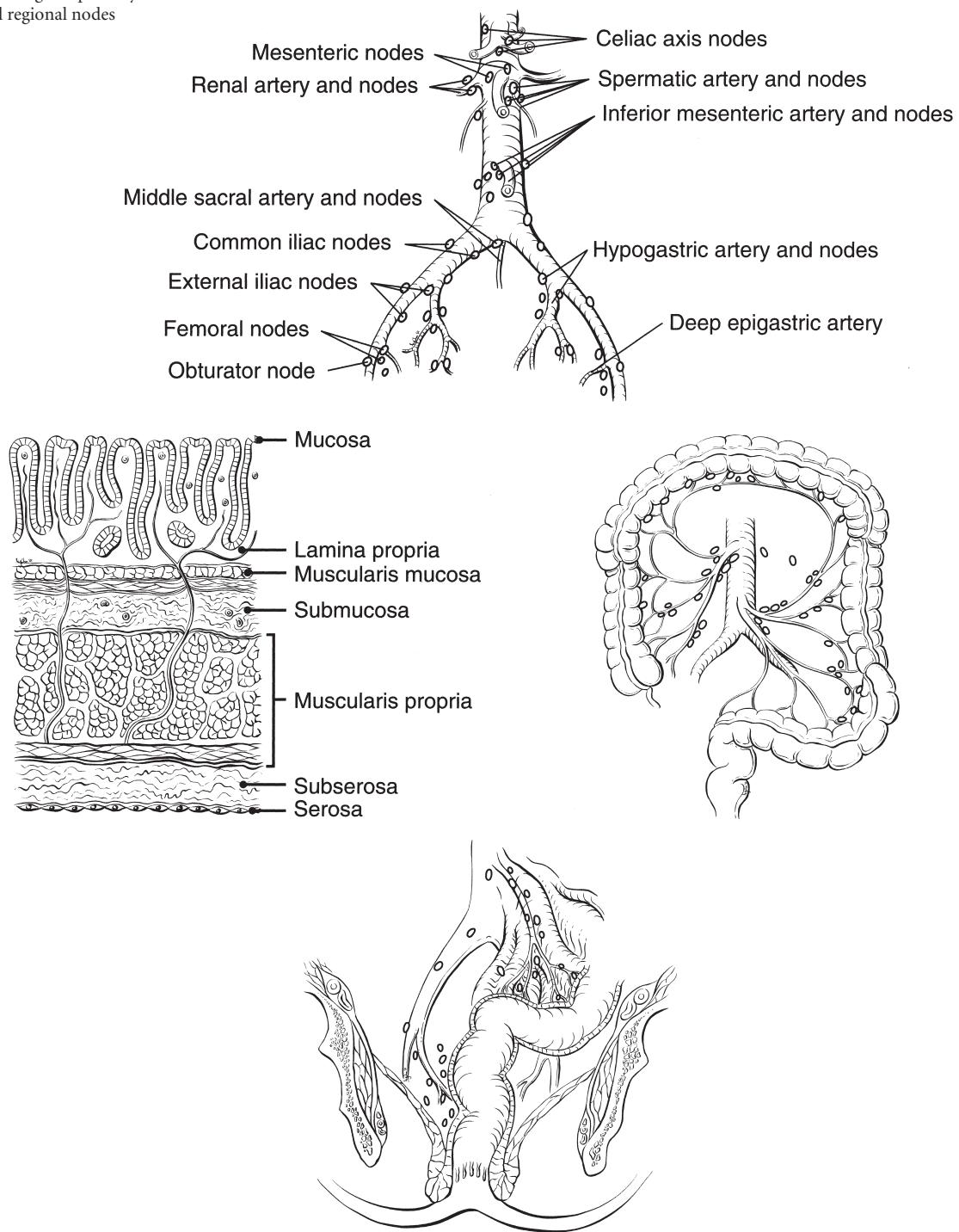
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Illustration

Indicate on diagram primary tumor and regional nodes involved.



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