

Protocol for the Examination of Resection Specimens from Patients with Primary Carcinoma of the Colon and / or Rectum

Version: 4.4.0.0

Protocol Posting Date: June 2025

CAP Laboratory Accreditation Program Protocol Required Use Date: March 2026

The changes included in this current protocol version affect accreditation requirements. The new deadline for implementing this protocol version is reflected in the above accreditation date.

For accreditation purposes, this protocol should be used for the following procedures AND tumor types:

Procedure	Description
Colectomy	Includes specimens designated total, partial, or segmental resection
Rectal Resection	Includes specimens designated low anterior resection or abdominoperineal resection
Tumor Type	Description
Carcinoma	Invasive carcinomas including small cell and large cell (poorly differentiated) neuroendocrine carcinoma

This protocol is NOT required for accreditation purposes for the following:

Procedure
Primary resection specimen with no residual cancer (e.g., following neoadjuvant therapy)
Cytologic specimens

The following should NOT be reported using this protocol:

Procedure
Excisional biopsy (polypectomy) (consider the Colon Excisional Biopsy protocol)
Endoscopic mucosal resection
Endoscopic mucosal dissection
Transanal disk excision
Tumor Type
Well-differentiated neuroendocrine tumors (consider the Colorectal NET protocol)
Lymphoma (consider the Precursor and Mature Lymphoid Malignancies protocol)
Sarcoma (consider the Soft Tissue protocol)

Version Contributors

Cancer Committee Authors: Dhanpat Jain, MD*, William V. Chopp, MD*, Rondell P. Graham, MBBS*, Yue Xue, MD, PhD*

* Denotes primary author.

For any questions or comments, contact: cancerprotocols@cap.org.

Glossary:

Author: Expert who is a current member of the Cancer Committee, or an expert designated by the chair of the Cancer Committee.

Expert Contributors: Includes members of other CAP committees or external subject matter experts who contribute to the current version of the protocol.

Accreditation Requirements

Synoptic reporting with core and conditional data elements for designated specimen types* is required for accreditation.

- Data elements designated as core must be reported.
- Data elements designated as conditional only need to be reported if applicable.
- Data elements designated as optional are identified with "+". Although not required for accreditation, they may be considered for reporting.

This protocol is not required for recurrent or metastatic tumors resected at a different time than the primary tumor. This protocol is also not required for pathology reviews performed at a second institution (i.e., second opinion and referrals to another institution).

Full accreditation requirements can be found on the CAP website under [Accreditation Checklists](#).

A list of core and conditional data elements can be found in the Summary of Required Elements under Resources on the CAP Cancer Protocols [website](#).

**Includes definitive primary cancer resection and pediatric biopsy tumor types.*

Synoptic Reporting

All core and conditionally required data elements outlined on the surgical case summary from this cancer protocol must be displayed in synoptic report format. Synoptic format is defined as:

- Data element: followed by its answer (response), outline format without the paired Data element: Response format is NOT considered synoptic.
- The data element should be represented in the report as it is listed in the case summary. The response for any data element may be modified from those listed in the case summary, including "Cannot be determined" if appropriate.
- Each diagnostic parameter pair (Data element: Response) is listed on a separate line or in a tabular format to achieve visual separation. The following exceptions are allowed to be listed on one line:
 - Anatomic site or specimen, laterality, and procedure
 - Pathologic Stage Classification (pTNM) elements
 - Negative margins, as long as all negative margins are specifically enumerated where applicable
- The synoptic portion of the report can appear in the diagnosis section of the pathology report, at the end of the report or in a separate section, but all Data element: Responses must be listed together in one location
- Organizations and pathologists may choose to list the required elements in any order, use additional methods in order to enhance or achieve visual separation, or add optional items within the synoptic report. The report may have required elements in a summary format elsewhere in the report IN ADDITION TO but not as replacement for the synoptic report i.e., all required elements must be in the synoptic portion of the report in the format defined above.

Summary of Changes

v 4.4.0.0

- Updates to Histologic Type question
- Rectal Tumor Location, Tumor Budding Score, and Distance from Invasive Carcinoma to Distal Margin questions updated from optional to conditionally reported
- Multiple Primary Sites question updated from required to conditionally reported

Reporting Template

Protocol Posting Date: June 2025

Select a single response unless otherwise indicated.

CASE SUMMARY: (COLON AND RECTUM: Resection)

Standard(s): AJCC 8

SPECIMEN

Procedure

- ☐ Right hemicolectomy
- ☐ Transverse colectomy
- ☐ Left hemicolectomy
- ☐ Sigmoidectomy
- ☐ Low anterior resection
- ☐ Total abdominal colectomy
- ☐ Abdominoperineal resection
- ☐ Other (specify): _____
- ☐ Not specified

Macroscopic Evaluation of Mesorectum (required only for rectal cancers) (Note [A](#))

- ☐ Not applicable
- ☐ Complete
- ☐ Near complete
- ☐ Incomplete
- ☐ Cannot be determined: _____

TUMOR

Tumor Site (Note [B](#)) (select all that apply)

- ☐ Cecum: _____
- ☐ Ileocecal valve: _____
- ☐ Ascending colon: _____
- ☐ Hepatic flexure: _____
- ☐ Transverse colon: _____
- ☐ Splenic flexure: _____
- ☐ Descending colon: _____
- ☐ Sigmoid colon: _____
- ☐ Rectosigmoid: _____
- ☐ Rectum: _____

Rectal Tumor Location (required for only rectal primaries) (Note [B](#))

- ☐ Not applicable
- ☐ Entirely above anterior peritoneal reflection
- ☐ Entirely below anterior peritoneal reflection
- ☐ Straddles anterior peritoneal reflection
- ☐ Not specified
- ☐ Colon, NOS: _____

___ Cannot be determined (explain): _____

Histologic Type (Note C)

- ___ Adenocarcinoma
- ___ Mucinous adenocarcinoma
- ___ Poorly cohesive carcinoma
- ___ Signet-ring cell carcinoma
- ___ Medullary carcinoma
- ___ Serrated adenocarcinoma
- ___ Micropapillary adenocarcinoma
- ___ Adenoma-like adenocarcinoma
- ___ Adenosquamous carcinoma
- ___ Undifferentiated carcinoma, NOS
- ___ Carcinoma with sarcomatoid component
- ___ Large cell neuroendocrine carcinoma
- ___ Small cell neuroendocrine carcinoma
- ___ Mixed neuroendocrine-non-neuroendocrine neoplasm (MiNEN) (specify components):

___ Other histologic type not listed (specify): _____

___ Carcinoma, type cannot be determined: _____

+Histologic Type Comment: _____

Histologic Grade (Note D)

- ___ G1, well-differentiated
- ___ G2, moderately differentiated
- ___ G3, poorly differentiated
- ___ G4, undifferentiated
- ___ Other (specify): _____
- ___ GX, cannot be assessed: _____
- ___ Not applicable: _____

Tumor Size

___ Greatest dimension in Centimeters (cm): _____ cm

+Additional Dimension in Centimeters (cm): ___ x ___ cm

___ Cannot be determined (explain): _____

Multiple Primary Sites (e.g., hepatic flexure and transverse colon) (required only if applicable)

___ Not applicable (no additional primary site(s) present)

___ Present: _____

Please complete a separate checklist for each primary site

Presence of multiple tumors can be challenging from staging perspective. In case of multiple tumors prefix "m" should be used similar to other sites and the final stage is based on the highest stage tumor. One situation is where separate resections are performed for each tumor, in which case each tumor is staged independently, and a synoptic summary is provided for each specimen. The other situation is where 2 or more tumors are identified in the same resection specimen. In this setting most often the regional lymph nodes for each tumor site cannot be separated and only one synoptic report is suggested where each tumor can be identified separately (e.g., T1, T2, etc.) with features specific to each tumor, recorded under each tumor (T stage, histologic type and grade, LVI, ancillary molecular tests when performed, etc.) and other common feature recorded in the synoptic summary as combined findings (e.g., specimen length, margin status, nodal status, etc.). The final stage should be based on the worst tumor.

Tumor Extent

- ☐ No invasion (high-grade dysplasia)
- ☐ Invades lamina propria / muscularis mucosae (intramucosal carcinoma)
- ☐ Invades submucosa
- ☐ Invades into muscularis propria
- ☐ Invades through muscularis propria into the pericolic or perirectal tissue
- ☐ Invades visceral peritoneum (including tumor continuous with serosal surface through area of inflammation)
- ☐ Directly invades or adheres to adjacent structure(s) (specify): _____
- ☐ Cannot be determined: _____
- ☐ No evidence of primary tumor

Sub-mucosal Invasion (required only for pT1 tumors)

- ☐ Not applicable (not a pT1 tumor)
- ☐ Not identified
- ☐ Present

Depth of Sub-mucosal Invasion (Note [E](#))

- ☐ Less than 1 mm
- ☐ Greater than or equal to 1 mm and less than 2 mm
- ☐ Greater than 2 mm
- ☐ Exact depth in Millimeters (mm): _____ mm
- ☐ Cannot be determined (explain): _____

Extent of Sub-mucosal Invasion (Note [E](#))

- ☐ Tumor invades into upper one third of submucosa
- ☐ Tumor invades into middle one third of submucosa
- ☐ Tumor invades into lower one third of submucosa
- ☐ Cannot be determined (explain): _____

Macroscopic Tumor Perforation (Note [F](#))

- ☐ Not identified
- ☐ Present
- ☐ Cannot be determined: _____

Lymphatic and / or Vascular Invasion (Note [G](#)) (select all that apply)

- ☐ Not identified
- ☐ Small vessel: _____
- ☐ Large vessel (venous), intramural: _____
- ☐ Large vessel (venous), extramural: _____
- ☐ Present, NOS: _____
- ☐ Cannot be determined: _____

Perineural Invasion (Note [G](#))

- ☐ Not identified
- ☐ Present

___ Cannot be determined: _____

Tumor Budding Score (required only when applicable) (Note [H](#))

- ___ Not applicable
- ___ Low (0-4)
- ___ Intermediate (5-9)
- ___ High (10 or more)
- ___ Cannot be determined: _____

+Number of Tumor Buds (per 'hotspot' field) (Note [H](#))

- ___ Specify number in one 'hotspot' field (in an area = 0.785 mm²): _____ per 'hotspot' field
- ___ Other (specify): _____
- ___ Cannot be determined: _____

+Type of Polyp in which Invasive Carcinoma Arose (Note [I](#))

- ___ None identified
- ___ Tubular adenoma
- ___ Villous adenoma
- ___ Tubulovillous adenoma
- ___ Traditional serrated adenoma
- ___ Sessile serrated adenoma / sessile serrated polyp
- ___ Hamartomatous polyp
- ___ Other (specify): _____

Treatment Effect (Note [J](#))

- ___ No known presurgical therapy
- ___ Present, with no viable cancer cells (complete response, score 0)
- ___ Present, with single cells or rare small groups of cancer cells (near complete response, score 1)
- ___ Present, with residual cancer showing evident tumor regression, but more than single cells or rare small groups of cancer cells (partial response, score 2)
- ___ Present, NOS
- ___ Absent, with extensive residual cancer and no evident tumor regression (poor or no response, score 3)
- ___ Cannot be determined: _____

+Tumor Comment: _____

MARGINS (Note [K](#))

Margin Status for Invasive Carcinoma

- ___ All margins negative for invasive carcinoma

+Closest Margin(s) to Invasive Carcinoma (select all that apply)

- ___ Proximal: _____

___ Distal: _____
___ Radial (circumferential): _____
___ Mesenteric: _____
___ Deep: _____
___ Mucosal (specify location, if possible): _____
___ Other (specify): _____
___ Cannot be determined: _____

+Distance from Invasive Carcinoma to Closest Margin

Specify in Centimeters (cm)

___ Exact distance in cm: _____ cm
___ Greater than 1 cm

Specify in Millimeters (mm)

___ Exact distance in mm: _____ mm
___ Greater than 10 mm

Other

___ Other (specify): _____
___ Cannot be determined: _____

Distance from Invasive Carcinoma to Radial (Circumferential) Margin (required only for rectal tumors)

___ Not applicable (not a rectal tumor)
___ Distance already reported as closest margin: _____

Specify in Centimeters (cm)

___ Exact distance in cm: _____ cm
___ Greater than 1 cm

Specify in Millimeters (mm)

___ Exact distance in mm: _____ mm
___ Greater than 10 mm

Other

___ Other (specify): _____
___ Cannot be determined: _____

Distance from Invasive Carcinoma to Distal Margin (required only for rectal tumors)

___ Not applicable (not a rectal tumor)
___ Distance already reported as closest margin: _____

Specify in Centimeters (cm)

___ Exact distance in cm: _____ cm
___ Greater than 1 cm

Specify in Millimeters (mm)

___ Exact distance in mm: _____ mm
___ Greater than 10 mm

Other

___ Other (specify): _____
___ Cannot be determined: _____

___ Invasive carcinoma present at margin

Margin(s) Involved by Invasive Carcinoma (select all that apply)

___ Proximal: _____
___ Distal: _____
___ Radial (circumferential): _____
___ Mesenteric: _____

- ☐ Deep: _____
- ☐ Mucosal (specify location, if possible): _____
- ☐ Other (specify): _____
- ☐ Cannot be determined: _____
- ☐ Other (specify): _____
- ☐ Cannot be determined (explain): _____
- ☐ Not applicable

Margin Status for Non-Invasive Tumor (select all that apply)

- ☐ All margins negative for high-grade dysplasia / intramucosal carcinoma and low-grade dysplasia
- ☐ High-grade dysplasia / intramucosal carcinoma present at margin

Margin(s) Involved by High-Grade Dysplasia / Intramucosal Carcinoma (select all that apply)

- ☐ Proximal: _____
- ☐ Distal: _____
- ☐ Mucosal (specify location, if possible): _____
- ☐ Other (specify): _____
- ☐ Cannot be determined: _____
- ☐ Low-grade dysplasia present at margin

Margin(s) Involved by Low-Grade Dysplasia (select all that apply)

- ☐ Proximal: _____
- ☐ Distal: _____
- ☐ Mucosal (specify location, if possible): _____
- ☐ Other (specify): _____
- ☐ Cannot be determined: _____
- ☐ Other (specify): _____
- ☐ Cannot be determined (explain): _____
- ☐ Not applicable

+Margin Comment: _____

REGIONAL LYMPH NODES

Regional Lymph Node Status

- ☐ Not applicable (no regional lymph nodes submitted or found)
- ☐ Regional lymph nodes present
- ☐ All regional lymph nodes negative for tumor
- ☐ Tumor present in regional lymph node(s)

Number of Lymph Nodes with Tumor

- ☐ Exact number (specify): _____
- ☐ At least (specify): _____
- ☐ Other (specify): _____
- ☐ Cannot be determined (explain): _____
- ☐ Other (specify): _____
- ☐ Cannot be determined (explain): _____

Number of Lymph Nodes Examined

- ☐ Exact number (specify): _____

___ At least (specify): _____
___ Other (specify): _____
___ Cannot be determined (explain): _____

Tumor Deposits (Note [L](#))

___ Not identified
___ Present

Number of Tumor Deposits

___ Specify number: _____
___ Other (specify): _____
___ Cannot be determined (explain): _____
___ Cannot be determined: _____

+Regional Lymph Node Comment: _____

DISTANT METASTASIS

Distant Site(s) Involved, if applicable (select all that apply)

___ Not applicable
___ Non-regional lymph node(s): _____
___ Liver: _____
___ Other (specify): _____
___ Cannot be determined: _____

pTNM CLASSIFICATION (AJCC 8th Edition) (Note [E](#))

Reporting of pT, pN, and (when applicable) pM categories is based on information available to the pathologist at the time the report is issued. As per the AJCC (Chapter 1, 8th Ed.) it is the managing physician's responsibility to establish the final pathologic stage based upon all pertinent information, including but potentially not limited to this pathology report.

Modified Classification (required only if applicable) (select all that apply)

___ Not applicable
___ y (post-neoadjuvant therapy)
___ r (recurrence)

pT Category

___ pT not assigned (cannot be determined based on available pathological information)
___ pT0: No evidence of primary tumor
___ pTis: Carcinoma in situ, intramucosal carcinoma (involvement of lamina propria with no extension through muscularis mucosae)
___ pT1: Tumor invades the submucosa (through the muscularis mucosa but not into the muscularis propria)
___ pT2: Tumor invades the muscularis propria
___ pT3: Tumor invades through the muscularis propria into pericorectal tissues
pT4: Tumor invades the visceral peritoneum or invades or adheres to adjacent organ or structure
___ pT4a: Tumor invades# through the visceral peritoneum (including gross perforation of the bowel through tumor and continuous invasion of tumor through areas of inflammation to the surface of the visceral peritoneum)

___ pT4b: Tumor directly invades# or adheres## to adjacent organs or structures

___ pT4 (subcategory cannot be determined)#, ##

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 retroperitoneal 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).

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 classification should be used to identify the presence or absence of vascular or lymphatic invasion whereas the PN prognostic factor should be used for perineural invasion.

T Suffix (required only if applicable)

___ Not applicable

___ (m) multiple primary synchronous tumors in a single organ

pN Category

___ pN not assigned (no nodes submitted or found)

___ pN not assigned (cannot be determined based on available pathological information)

___ pN0: No regional lymph node metastasis

pN1: One to three regional lymph nodes are positive (tumor in lymph nodes measuring greater than or equal to 0.2 mm), or any number of tumor deposits are present and all identifiable lymph nodes are negative

___ pN1a: One regional lymph node is positive

___ pN1b: Two or three regional lymph nodes are positive

___ pN1c: No regional lymph nodes are positive, but there are tumor deposits in the subserosa, mesentery, nonperitonealized pericolic or perirectal / mesorectal tissues

___ pN1 (subcategory cannot be determined)

pN2: Four or more regional nodes are positive

___ pN2a: Four to six regional lymph nodes are positive

___ pN2b: Seven or more regional lymph nodes are positive

___ pN2 (subcategory cannot be assessed)

pM Category (required only if confirmed pathologically)

___ Not applicable - pM cannot be determined from the submitted specimen(s)

pM1: Metastasis to one or more distant sites or organs or peritoneal metastasis is identified

___ pM1a: Metastasis to one site or organ is identified without peritoneal metastasis

___ pM1b: Metastasis to two or more sites or organs is identified without peritoneal metastasis

___ pM1c: Metastasis to the peritoneal surface is identified alone or with other site or organ metastases

___ pM1 (subcategory cannot be determined)

ADDITIONAL FINDINGS

+Additional Findings (select all that apply)

- ☐ None identified
- ☐ Adenoma(s)
- ☐ Ulcerative colitis
- ☐ Crohn disease
- ☐ Diverticulosis
- ☐ Dysplasia arising in inflammatory bowel disease
- ☐ Other (specify): _____

SPECIAL STUDIES (Note [M](#))

For reporting molecular testing and immunohistochemistry for mismatch repair proteins, and for other cancer biomarker testing results, the CAP Colorectal Biomarker Template should be used. Pending biomarker studies should be listed in the Comments section of this report.

COMMENTS

Comment(s): _____

Explanatory Notes

A. Mesorectal Envelope

The quality of the surgical technique is a key factor in the success of surgical treatment for rectal cancer, both in the prevention of local recurrence and in long-term survival. The procedures in which mesorectal evaluation is typically relevant include low anterior resection and abdominoperineal resection. Numerous studies have demonstrated that total mesorectal excision (TME) improves local recurrence rates and the corresponding survival by as much as 20%. This surgical technique entails precise sharp dissection within the areolar plane outside (lateral to) the visceral mesorectal fascia to remove the rectum. This plane encases the rectum, its mesentery, and all regional nodes and constitutes Waldeyer's fascia. High-quality TME surgery reduces local recurrence from 20% to 30%, to 8% to 10% or less, and increases 5-year survival from 48% to 68%.^{1,2} Adjuvant therapy in the presence of a high-quality TME may further reduce local recurrence (from 8% to 2.6%).³

Pathologic evaluation of the resection specimen has been shown to be a sensitive means of assessing the quality of rectal surgery. It is superior to indirect measures of surgical quality assessment, such as perioperative mortality, rates of complication, number of local recurrences, and 5-year survival. Macroscopic pathologic assessment of the completeness of the mesorectum, scored as complete, partially complete, or incomplete, accurately predicts both local recurrence and distant metastasis.³ Microscopic parameters, such as the status of the circumferential resection margin, the distance between the tumor and nearest circumferential margin (i.e., "surgical clearance"), and the distance between the tumor and the closest distal margin, are all important predictors of local recurrence and may be affected by surgical technique.

The nonperitonealized surface of the fresh specimen is examined circumferentially, and the completeness of the mesorectum is scored as described below.³ The entire specimen is scored according to the worst area.

Incomplete

- Little bulk to the mesorectum
- Defects in the mesorectum down to the muscularis propria
- After transverse sectioning, the circumferential margin appears very irregular

Nearly Complete

- Moderate bulk to the mesorectum
- Irregularity of the mesorectal surface with defects greater than 5 mm, but none extending to the muscularis propria
- No areas of visibility of the muscularis propria except at the insertion site of the levator ani muscles

Complete

- Intact bulky mesorectum with a smooth surface
- Only minor irregularities of the mesorectal surface
- No surface defects greater than 5 mm in depth
- No coning towards the distal margin of the specimen
- After transverse sectioning, the circumferential margin appears smooth

References

1. Arbmán G, Nilsson E, Hallbook O, Sjö Dahl R. Local recurrence following total mesorectal excision for rectal cancer. *Br J Surg*. 1996; 83(3):375-379.
2. Kapiteijn E, Marijnen CA, Nagtegaal ID, et al. Preoperative radiotherapy combined with total mesorectal excision for resectable rectal cancer. *N Engl J Med*. 2001;345(9):638-646. [summary for patients in *Med J Aust*. 2002 Nov 18;177(10):563-564; PMID: 12429007].
3. Nagtegaal ID, Marijnen CA, Kranenbarg EK, van de Velde CJ, van Krieken JH. Pathology Review Committee-Cooperative Clinical Investigators. Circumferential margin involvement is still an important predictor of local recurrence in rectal carcinoma: not one millimeter but two millimeters is the limit. *Am J Surg Pathol*. 2002; 26(3):350-357.

B. Anatomic Sites

The protocol applies to all carcinomas arising in the colon and rectum.¹ It excludes carcinomas of the vermiform appendix and low-grade neuroendocrine neoplasms (carcinoid tumors).

The colon is divided as shown in Figure 1. The right colon is subdivided into the cecum and the ascending colon.² The left colon is subdivided into the descending colon and sigmoid colon (see Table 1).¹

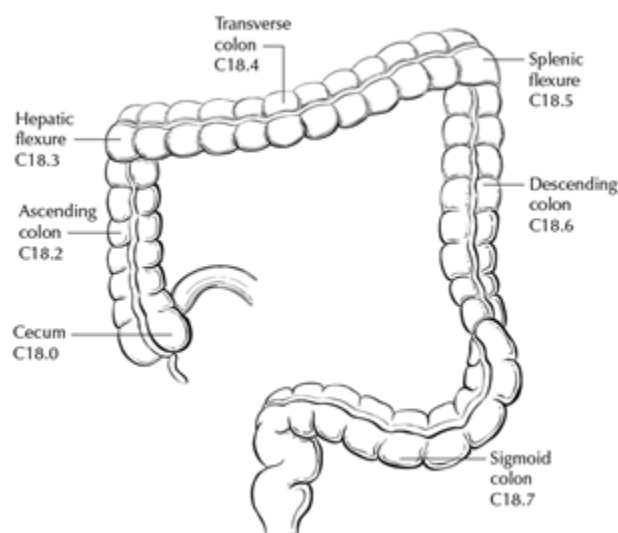


Figure 1. Anatomic subsites of the colon. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the AJCC Cancer Staging Atlas (2006), edited by Greene et al.² and published by Springer Science and Business Media, LLC, www.springerlink.com.

Table 1. Anatomic Subsites of the Colon and Rectum

Site	Relationship to Peritoneum (see Note J)	Dimensions (approximate)
Cecum	Entirely covered by peritoneum	6-9 cm
Ascending colon	Retroperitoneal; posterior surface lacks peritoneal covering; lateral and anterior surfaces covered by visceral peritoneum (serosa)	15-20 cm
Transverse colon	Intraperitoneal; has mesentery	Variable

Descending colon	Retroperitoneal; posterior surface lacks peritoneal covering; lateral and anterior surfaces covered by visceral peritoneum (serosa)	10-15 cm
Sigmoid colon	Intraperitoneal; has mesentery	Variable
Rectum	Upper third covered by peritoneum on anterior and lateral surfaces; middle third covered by peritoneum only on anterior surface; lower third has no peritoneal covering	16-20 cm

The transition from sigmoid to rectum is marked by the fusion of the tenia coli of the sigmoid to form the circumferential longitudinal muscle of the rectal wall approximately 16 to 20 cm from the dentate line. The rectum is defined clinically as the distal large intestine commencing opposite the sacral promontory and ending at the anorectal ring, which corresponds to the proximal border of the puborectalis muscle palpable on digital rectal examination (Figure 2).¹ When measuring below with a rigid sigmoidoscope, it extends 16 cm from the anal verge.

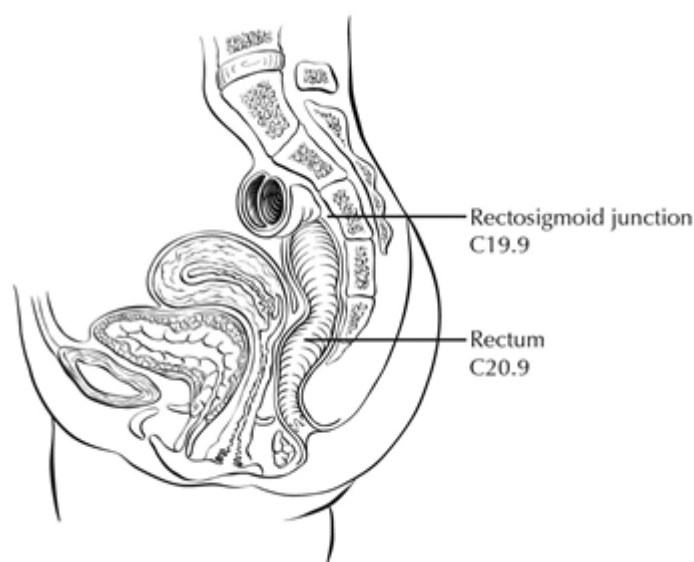


Figure 2. Anatomic subsites of the rectum. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the *AJCC Cancer Staging Atlas* (2006), edited by Greene et al.² and published by Springer Science and Business Media, LLC, www.springerlink.com.

Tumors located at the border between two subsites of the colon (e.g., cecum and ascending colon) are registered as tumors of the subsite that is more involved. If two subsites are involved to the same extent, the tumor is classified as an 'overlapping' lesion.

A tumor is classified as rectal if its inferior margin lies less than 16 cm from the anal verge or if any part of the tumor is located at least partly within the supply of the superior rectal artery.³ The rectum commences at the sacral promontory, and the junction of sigmoid colon and rectum is anatomically marked by fusion of tenia coli to form the circumferential longitudinal muscle of the rectal wall. Intraoperatively, the rectosigmoid junction corresponds to the sacral promontory. A tumor is classified as rectosigmoid when differentiation between rectum and sigmoid according to the previously mentioned guidelines is not possible.⁴

Anteriorly, the peritoneal reflection is located at the junction of middle and lower third of the rectum, while laterally, it is located at the junction of upper and middle third of the rectum. Posteriorly, the reflection is located higher and most of the posterior rectum does not have a serosal covering.

(a) Whether an adenocarcinoma located in the rectum has a radial (circumferential) resection margin or a peritoneal (serosal) surface depends on its location in relation to the peritoneal reflections. Tumors below the anterior peritoneal reflection will have a 360-degree radial margin while those above it may have a radial margin or a peritoneal (serosal) surface, or both, depending on the precise location.

(b) Neoadjuvant therapy and mesorectal excision are considered standard of care for rectal adenocarcinomas “below the anterior peritoneal reflection”, while the opinions about use of these modalities vary for rectal adenocarcinomas located above the anterior peritoneal reflection. Conservative options like transanal disc excisions are often considered for location “below the anterior peritoneal reflection”. In these contexts, the peritoneal reflection refers to the junction of upper and middle third of the rectum; there is ongoing debate in the surgical literature about the concept of peritoneal reflection.⁵ If information about tumor location with respect to the peritoneal reflection is included in the report, the aspect of rectum in question (posterior, lateral, anterior) should also be noted.

References

1. Amin MB, Edge SB, Greene FL, et al., eds. *AJCC Cancer Staging Manual*. 8th ed. New York, NY: Springer; 2017.
2. Greene FL, Compton CC, Fritz AG, Shah J, Winchester DP, eds. *AJCC Cancer Staging Atlas*. New York, NY: Springer; 2006.
3. Fielding LP, Arsenaault PA, Chapuis PH, et al. Clinicopathological staging for colorectal cancer: an International Documentation System (IDS) and an International Comprehensive Anatomical Terminology (ICAT). *J Gastroenterol Hepatol*. 1991; 6(4):325-344.
4. Wittekind C, Henson DE, Hutter RVP, Sobin LH, eds. *TNM Supplement: A Commentary on Uniform Use*. 2nd ed. New York, NY: Wiley-Liss; 2001.
5. Kenig J, Richter P. Definition of the rectum and level of the peritoneal reflection - still a matter of debate? *Wideochir Inne Tech Maloinwazyjne*. 2013; 8:183-186.

C. Histologic Types

For consistency in reporting, the histologic classification proposed by the World Health Organization (WHO) is recommended.¹

The histologic types of colorectal carcinoma that have been shown to have adverse prognostic significance independent of stage are signet-ring cell carcinoma² and poorly differentiated neuroendocrine carcinomas (large cell and small cell subtypes).³

Medullary carcinoma is a distinctive histologic type strongly associated with high levels of microsatellite instability (MSI-H), indicative of defects in DNA repair gene function. Medullary carcinoma may occur either sporadically or in association with Lynch syndrome.^{4,5,6} This tumor type is characterized by solid growth in nested, organoid, or trabecular patterns, with no immunohistochemical evidence of neuroendocrine differentiation. Medullary carcinomas are also characterized by numerous tumor infiltrating lymphocytes and a better prognosis.

Micropapillary carcinoma is characterized by small, tight clusters of tumor cells in cleft-like spaces and is often present in association with conventional adenocarcinoma. This variant is strongly associated with lymphovascular invasion and lymph node metastasis.⁷

Serrated adenocarcinomas are characterized by neoplastic glands showing prominent serrations, tumor cells with basal nuclei and eosinophilic cytoplasm, and no or minimal luminal necrosis. These tumors are thought to be related to traditional serrated adenomas and may have a more aggressive course than conventional adenocarcinoma.⁸

References

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D. Histologic Grade

A number of grading systems for colorectal cancer have been suggested, but a single widely accepted and uniformly used standard for grading is lacking. Most systems stratify tumors into 3 or 4 grades as follows:

- | | |
|---------|---|
| Grade 1 | Well-differentiated (>95% gland formation) |
| Grade 2 | Moderately differentiated (50-95% gland formation) |
| Grade 3 | Poorly differentiated (<50% gland formation) |
| Grade 4 | Undifferentiated (no gland formation or mucin; no squamous or neuroendocrine differentiation) |

Despite a significant degree of interobserver variability,¹ histologic grade has been shown to be an important prognostic factor in many studies,^{2,3} with strong correlation between poor differentiation and adverse outcome.⁴ While some studies have stratified grade into a two-tiered low- and high-grade system, a three- or four-tier system is more commonly used for gastrointestinal carcinomas. The AJCC

has specified use of a four-tiered grading system for colorectal cancer for the 8th edition of the TNM manual.⁵ Pathologists should use the four-tier histologic grading scheme as specified above to prevent errors in data recording. As per WHO, the grading scheme applies to adenocarcinoma, not otherwise specified, and not to histologic variants. For example, medullary carcinomas behave as low-grade tumors even though they may appear poorly differentiated. This grading scheme is also not applicable to poorly differentiated neuroendocrine carcinomas.

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E. pTNM Classification

Surgical resection remains the most effective therapy for colorectal carcinoma, and the best estimation of prognosis is derived from the pathologic findings on the resection specimen. The anatomic extent of disease is by far the most important prognostic factor in colorectal cancer.

The protocol recommends the TNM staging system of the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC)¹ but does not preclude the use of other staging systems.

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 or biopsy 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.

TNM Descriptors:

For identification of special cases of TNM or pTNM classifications, the “m” suffix and “y” and “r” 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 (i.e., 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 (i.e., 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.

T Category Considerations (Figures 4-6):

pTis: For colorectal carcinomas, carcinoma in situ (*pTis*) as a staging term refers to tumors involving the lamina propria and/or muscularis mucosae, but not extending through it (intramucosal carcinoma). Tumor extension through the muscularis mucosae into the submucosa is classified as T1 (Figure 5). A synoptic report is required only for invasive tumors, but not Tis.

pT1: Assessment of the depth of submucosal invasion in early colon carcinoma has also been shown to be important in predicting adverse outcomes, but the methodology has varied and evolved with time.^{2,3,4,5,6} In pedunculated polyps the depth of invasion was evaluated as four levels (head, neck, stalk and beyond stalk) called Haggitt levels. Submucosal involvement has been also been divided into superficial, mid, and deep levels (Kikuchi levels sm1, sm2, and sm3), and the risk of nodal metastasis increases with increasing depth of submucosal invasion.^{2,3,4,5,6} Other systems use actual measurement of depth of invasion and show the risk of nodal metastasis is none to minimal when the depth of invasion is <1mm and becomes significant with invasion of ≥2mm.^{5,6} However, it can be difficult to accurately assess the depth or extent of submucosal involvement due to improper orientation and when the muscularis mucosae is completely destroyed due tumor invasion and extensive desmoplasia. In such a setting, it has been suggested to draw a line joining preserved muscularis mucosae at the edges of the tumor to indicate the likely level of muscularis mucosae or when that is not possible measure the depth of invasion from the surface of the tumor. Each method for evaluating submucosal involvement has certain limitations, and while one is not clearly superior to the other, the Haggitt levels and Kikuchi levels can only be applied to a subset of colorectal carcinoma specimens and hence measurement of the actual depth appears most practical for the sake of consistency and the recommended method for malignant polyps.

pT4: Tumors that involve the serosal surface (visceral peritoneum) or directly invade adjacent organs or structures are assigned to the T4 category (Figures 4 and 6).

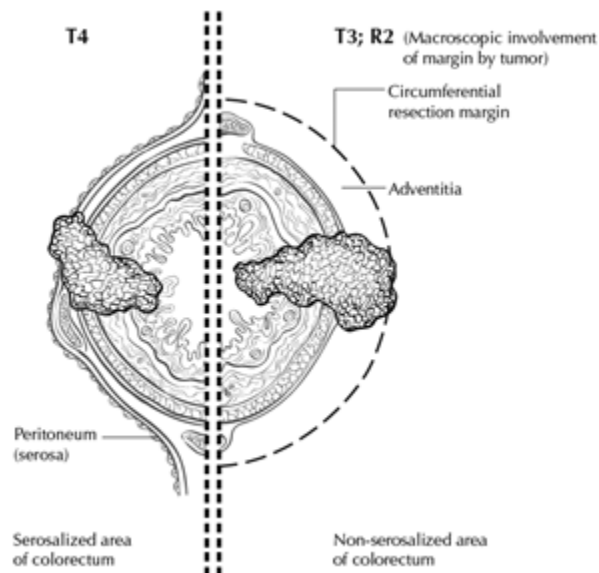
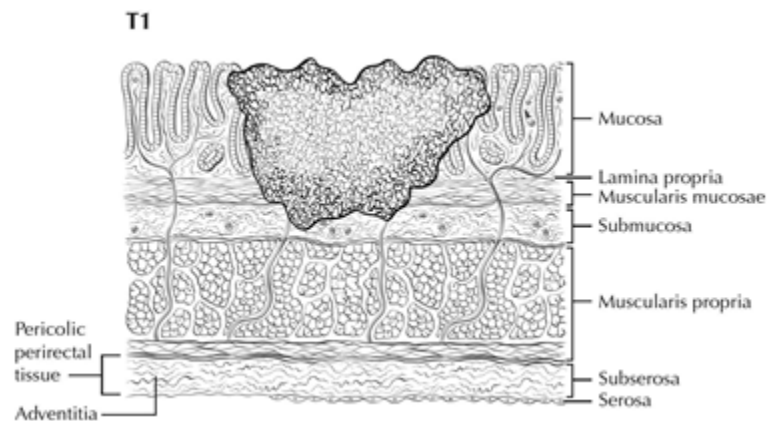


Figure 4. T4a (left side) with involvement of serosa (visceral peritoneum) by tumor cells in a segment of colorectum with a serosal covering. In contrast, the right side of the diagram shows T3 with macroscopically positive circumferential margin (designated R2 in AJCC staging system), corresponding to gross disease remaining after surgical excision. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the *AJCC Cancer Staging Atlas* (2006), edited by Greene et al.⁷ and published by Springer Science and Business Media, LLC, www.springerlink.com.



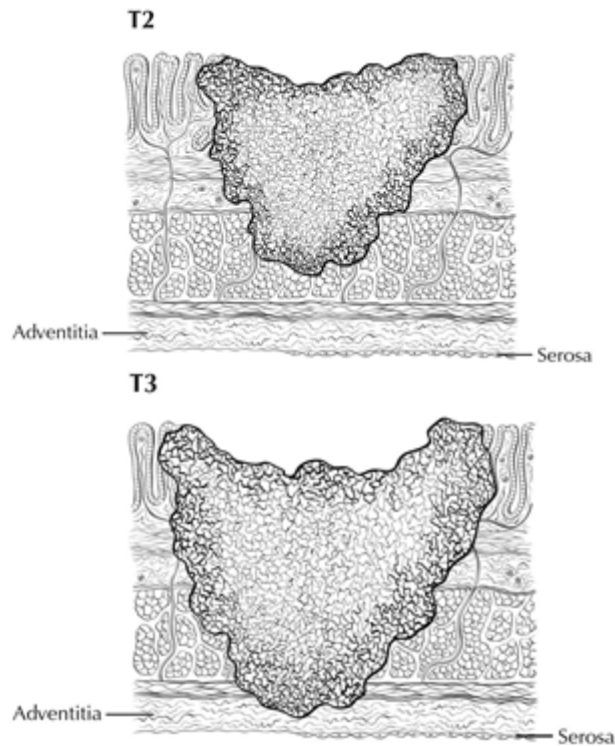


Figure 5. T1 tumor invades submucosa; T2 tumor invades muscularis propria; T3 tumor invades through the muscularis propria into the subserosa or into nonperitonealized pericolic or perirectal tissues (adventitia). Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the *AJCC Cancer Staging Atlas* (2006) edited by Greene et al.⁷ and published by Springer Science and Business Media, LLC, www.springerlink.com.

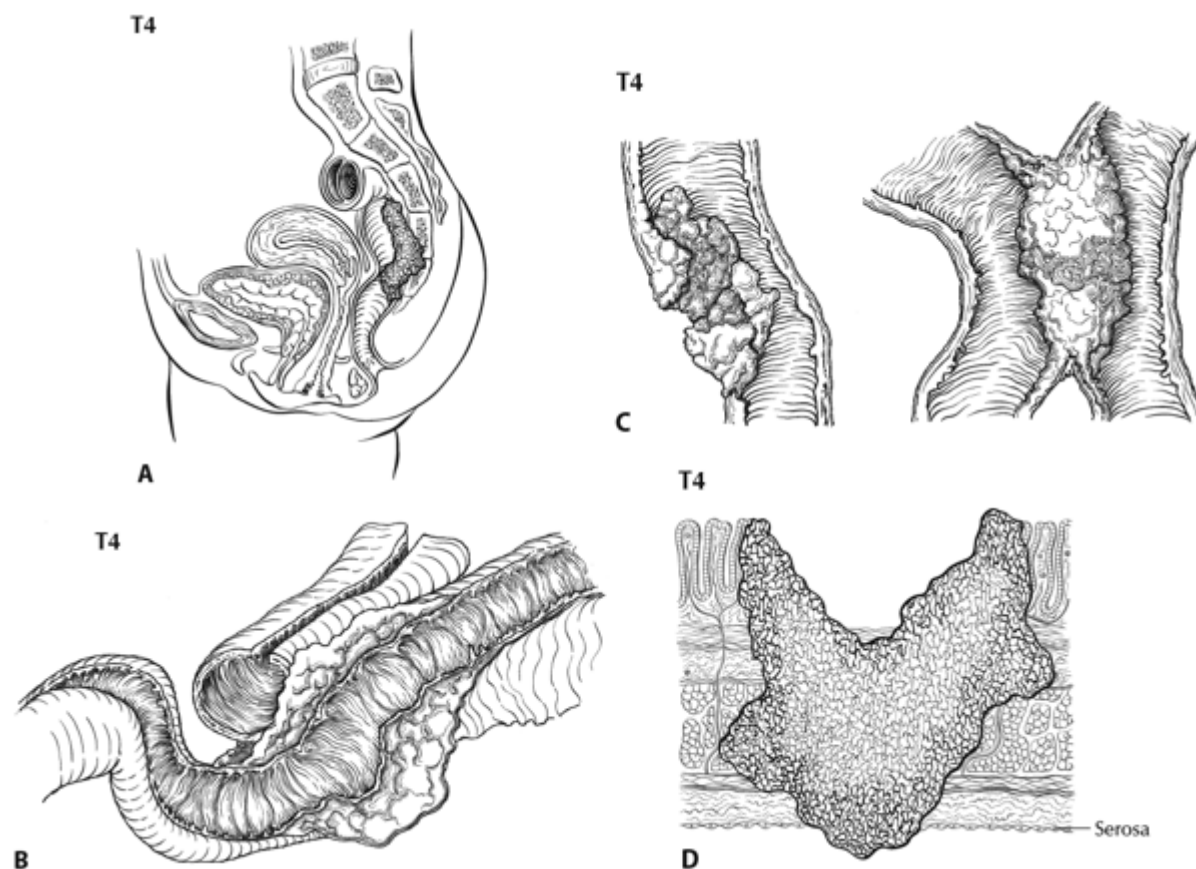


Figure 6. A, T4b tumor showing direct invasion of coccyx. B, T4b tumor directly invading adjacent loop of small bowel. C, T4a tumor showing gross perforation of bowel through tumor (left). The right-hand panel shows T4b tumor directly invading adjacent bowel. D, T4a tumor with involvement of serosa (visceral peritoneum) by tumor cells. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the *AJCC Cancer Staging Atlas* (2006), edited by Greene et al.⁷ and published by Springer Science and Business Media, LLC, www.springerlink.com.

Tumor that is adherent to other organs or structures macroscopically is classified clinically as cT4. However, if no tumor is found within the adhesion microscopically, the tumor should be assigned pT3.¹

For rectal tumors, invasion of the external sphincter and/or levator ani muscle(s) is classified as T4b.

Tumor in veins or lymphatics does not affect the pT classification.

Subdivision of T4 into T4a and T4b: Serosal (visceral peritoneal) involvement by tumor cells (pT4a) has been demonstrated by multivariate analysis to have a negative impact on prognosis,^{8,9} as does direct invasion of adjacent organs (pT4b). Visceral peritoneal involvement can be missed without thorough sampling and/or sectioning, and malignant cells have been identified in serosal scrapings in as many as 26% of specimens categorized as pT3 by histologic examination alone.^{10,11} Although the absence of standard guidelines for assessing peritoneal involvement may contribute to underdiagnosis, the following findings are considered to represent serosal involvement by tumor:

- Tumor present at the serosal surface
- Free tumor cells on the serosal surface (on the visceral peritoneum) with underlying erosion/ulceration of mesothelial lining, mesothelial hyperplasia and/or inflammatory reaction^{9,10}
- Perforation in which the tumor cells are continuous with the serosal surface through inflammation

The significance of tumors that are <1 mm from the serosal surface and accompanied by serosal reaction is unclear, with some¹⁰ but not all studies⁹ indicating a higher risk of peritoneal recurrence. Multiple level sections and/or additional sections of the tumor should be examined in these cases. If the serosal involvement is not present after additional evaluation, the tumor should be assigned to the pT3 category. The use of elastic stains has been advocated for identification of T4a tumors by demonstrating tumor involvement of the subperitoneal elastic lamina. The elastic stain can be difficult to interpret in this region and the elastic lamina is not uniformly present even in normal colon. Hence routine use of this stain is not considered standard practice. In portions of the colorectum that are not peritonealized (e.g., posterior aspects of ascending and descending colon, lower portion of rectum), the T4a category is not applicable.

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 retroperitoneal 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).

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 prognostic factor should be used for perineural invasion.

Intramural extension of tumor from one subsite (segment) of the large intestine into an adjacent subsite or into the ileum (e.g., for a cecal carcinoma) or anal canal (e.g., for a rectal carcinoma) does not affect the pT classification. Transmural extension into another organ or site is necessary for T4b designation. Both types of peritoneal involvement are associated with decreased survival. Although small studies suggested that serosal involvement was associated with worse outcome than invasion of adjacent organs, data from a large cohort of more than 100,000 colon cancer cases¹¹ indicate that penetration of the visceral peritoneum carries a 10% to 20% better 5-year survival than locally invasive carcinomas for the same pN category.

N Category Considerations:

The regional lymph nodes for the anatomical subsites of the large intestine (Figure 7) are as follows:

Cecum:

Pericolic, ileocolic, right colic

Ascending colon:

Pericolic, ileocolic, right colic, right branch of middle colic

Hepatic flexure:

Pericolic, ileocolic, middle colic, right colic

Transverse colon:

Pericolic, middle colic

Splenic flexure:

Pericolic, middle colic, left colic

Descending colon:

Pericolic, left colic, inferior mesenteric, sigmoid

Sigmoid colon:

Pericolic, sigmoid, inferior mesenteric, superior rectal (hemorrhoidal)

Rectosigmoid: Pericolic, sigmoid, superior rectal (hemorrhoidal)

Rectum:

Mesorectal, superior rectal (hemorrhoidal), inferior mesenteric, internal iliac, inferior rectal (hemorrhoidal)

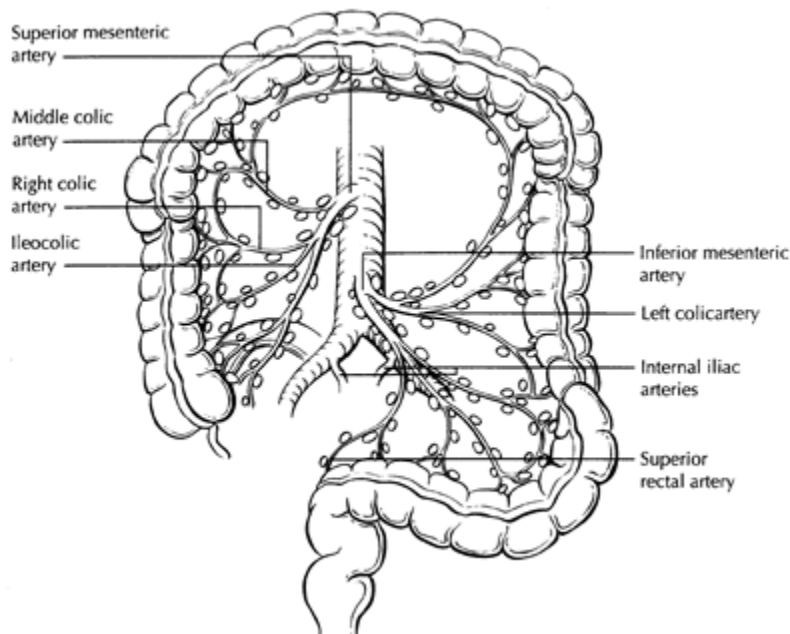


Figure 7. The regional lymph nodes of the colon and rectum. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Ill. The original source for this material is the *AJCC Cancer Staging Atlas* (2006), edited by Greene et al.⁷ and published by Springer Science and Business Media, LLC, www.springerlink.com.

For rectal cancers, metastasis in the external iliac or common iliac nodes is classified as distant metastasis.¹

Submission of Lymph Nodes for Microscopic Examination. All grossly negative or equivocal lymph nodes should be submitted entirely. Grossly positive lymph nodes may be partially submitted for microscopic confirmation of metastasis.

The accuracy and predictive value of stage II assignment are directly proportional to the thoroughness of the surgical technique in removing all regional nodes and the pathologic examination of the resection specimen in identifying and harvesting all regional lymph nodes for microscopic assessment. The National Quality Forum lists the presence of at least 12 lymph nodes in a surgical resection among the key quality measures for colon cancer care in the United States (see <http://www.facs.org/cancer/qualitymeasures.html>).

The likelihood of detecting metastasis increases with the number of lymph nodes examined; hence 12 lymph nodes should be considered the minimum target, but all possible lymph nodes should be retrieved and examined.^{12,13}

The clinical outcome is linked to lymph node harvest in stage II disease,¹⁴ indicating a positive effect of optimal mesenteric resection by the surgeon, optimal lymph node harvest from the resection specimen by the pathologist, or both.

The number of lymph nodes recovered from a resection specimen is dependent on several factors. Surgical technique, surgery volume, and patient factors (e.g., age and anatomic variation) alter the actual number of nodes in a resection specimen, but the diligence and skill of the pathologist in identifying and harvesting lymph nodes in the resection specimen also are major factors. Lymph nodes may be more difficult to identify in specimens from patients who are obese¹⁵ or elderly, or after neoadjuvant therapy.¹⁶ Because it has been shown that nodal metastasis in colorectal cancer is often found in small lymph nodes (<5 mm in diameter), diligent search for lymph nodes is required on gross examination of resection specimens. If fewer than 12 lymph nodes are found, re-examining the specimen for additional lymph nodes, with or without visual enhancement techniques, should be considered. The pathology report should clearly state the total number of lymph nodes examined and the total number involved by metastases. Data are insufficient to recommend routine use of tissue levels or special/ancillary techniques.

Nonregional Lymph Nodes. For microscopic examination of lymph nodes in large resection specimens, lymph nodes must be designated as regional versus nonregional, according to the anatomic location of the tumor. Metastasis to nonregional lymph nodes is classified as distant metastasis and designated as M1.

Lymph Nodes Replaced by Tumor. A tumor nodule in the pericolic/perirectal fat without histologic evidence of residual lymph node tissue is classified as a tumor deposit (peritumoral deposit or satellite nodule) and is not considered a positive lymph node. In the absence of unequivocal lymph node metastases, tumor deposits are recorded as N1c.¹

Isolated Tumor Cells. Isolated tumor cells (ITCs) are defined as single tumor cells or small clusters of tumor cells measuring less than 0.2 mm, usually found by special techniques such as immunohistochemical staining, and are classified as N0.¹ Because the biologic significance of ITCs (either a single focus in a single node, multiple foci within a single or multiple nodes) remains unproven, N0 is considered justified.¹⁷ The number of lymph nodes involved by ITCs should be clearly stated in a comment section or elsewhere in the report. Metastatic deposits 0.2 mm-2.0 mm have been referred to as micrometastasis. These nodes should be considered as involved by cancer.

Routine assessment of regional lymph nodes is limited to conventional pathologic techniques (gross assessment and histologic examination), and data are currently insufficient to recommend special measures to detect ITCs. Thus, neither multiple levels of paraffin blocks nor the use of special/ancillary techniques such as immunohistochemistry are recommended for routine examination of regional lymph nodes.

TNM Anatomic Stage/Prognostic Groupings

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 (e.g., 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.

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F. Perforation

Tumor perforation is an uncommon complication of colorectal cancer, but one that is associated with a poor outcome, including high in-hospital mortality and morbidity.¹ Perforation of the uninvolved colon proximal to an obstructing tumor is also associated with high mortality because of generalized peritonitis and sepsis. Reported perforation rates range from 2.6% to 9%. Perforation is more likely to occur in older patients.

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G. Lymphatic and/or Vascular Invasion and Perineural Invasion

It is recommended that small vessel vascular invasion should be reported separately from venous (large vessel) invasion. Small vessel invasion indicates tumor involvement of thin-walled structures lined by endothelium, without an identifiable smooth muscle layer or elastic lamina. Small vessels include lymphatics, capillaries, and postcapillary venules. Differentiation of lymphatics from other vascular channels may require application of lymphatic specific endothelial markers like D2-40.¹ Small vessel invasion is associated with lymph node metastasis and has been shown to be independent indicator of adverse outcome in several studies.^{2,3} The higher prognostic significance of extramural small vessel

invasion has been suggested,⁴ but the importance of anatomic location in small vessel invasion (extramural or intramural) is not well defined.

Tumor involving endothelium-lined spaces with an identifiable smooth muscle layer or elastic lamina is considered venous (large vessel) invasion. Circumscribed tumor nodules surrounded by an elastic lamina on hematoxylin-eosin (H&E) or elastic stain are also considered venous invasion. Venous invasion can be extramural (beyond muscularis propria) or intramural (submucosa or muscularis propria). Extramural venous invasion has been demonstrated by multivariate analysis to be an independent adverse prognostic factor in multiple studies and is a risk factor for liver metastasis.⁴ The significance of intramural venous invasion is less clear. Histologic features like tumor deposits adjacent to arteries (“orphan artery” sign) and elongated tumor nodules extending into pericolic fat from the muscularis propria (“protruding tongue” sign) can raise the suspicion for venous invasion.⁵ Elastic stain can lead to 2- to 3-fold increase in the detection of venous invasion and may be used to improve assessment of this feature.⁶

Perineural invasion has been shown to be independent indicator of poor prognosis.^{7,8,9} While some series did not find perineural invasion to be a significant predictive factor in stage II disease,^{10,11} many studies have confirmed its adverse effect on survival in stage II disease.^{3,12} Extramural perineural invasion may have a greater adverse prognostic effect,⁸ but the distinction between intramural and extramural perineural invasion has not been well studied.

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H. Tumor Budding

The presence of single cells or small clusters of less than five cells at the advancing front of the tumor is considered as peritumoral tumor budding. Numerous studies have shown that high tumor budding in adenocarcinoma arising in a polyp is a significant risk factor for nodal involvement,^{1,2,3,4,5,6} with tumor budding being the most significant factor in some studies.³ Similarly, the adverse prognosis of high tumor budding has been shown in stage II patients and its inclusion as a high risk factor for making chemotherapy decisions for stage II patients has been advocated.^{4,6} Different criteria for evaluating and reporting tumor budding have been followed in the literature. An international tumor budding consensus conference (ITBCC) in 2016 recommended the following criteria for evaluating tumor budding:⁷

1. Tumor budding counts should be done on H&E sections. In cases of obscuring factors like inflammation, immunohistochemistry for keratin can be obtained to assess the advancing edge for tumor buds, but the scoring should be done on H&E sections.
2. Tumor budding should be reported by selecting a “hotspot” chosen after review of all available slides with invasive tumor. The total number of buds should be reported in an area measuring 0.785 mm², which corresponds to 20x field in some microscopes (use appropriate conversion for other microscopes, see table below).
3. Both total number of buds and a three-tier score (based on 0.785 mm² field area) should be reported: low (0-4 buds), intermediate (5-9 buds), and high (10 or more buds).

This is now a required element for conventional adenocarcinoma (well or moderately differentiated). Tumor budding cannot be assessed or is not applicable for other specific variants of colorectal adenocarcinoma (e.g., micropapillary carcinoma, signet ring carcinoma or mucinous carcinoma) or poorly/undifferentiated carcinoma, unless there is also an associated component of well or moderately differentiated conventional adenocarcinoma.

Objective Magnification:		20		
Eyepiece FN Diameter	Eyepiece FN Radius	Specimen FN Radius	Specimen Area	Normalization Factor
(mm)	(mm)	(mm)	(mm ²)	
18	9.0	0.450	0.636	0.810
19	9.5	0.475	0.709	0.903
20	10.0	0.500	0.785	1.000
21	10.5	0.525	0.866	1.103
22	11.0	0.550	0.950	1.210
23	11.5	0.575	1.039	1.323
24	12.0	0.600	1.131	1.440
25	12.5	0.625	1.227	1.563
26	13.0	0.650	1.327	1.690

Table. ITBCC Normalization Table for Reporting Tumor Budding According to Microscope.

To obtain tumor bud count for your field of view, divide by the normalization number.

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I. Polyps

The adenocarcinoma can arise in adenomatous (tubular, tubulovillous, or villous) or serrated (sessile serrated adenoma/polyp or traditional serrated adenoma) polyp. Sessile serrated adenoma often develops cytologic dysplasia resembling tubular adenoma during neoplastic progression. These are presumed to be the precursors of right-sided adenocarcinomas with high levels of microsatellite instability (MSI-H).¹

References

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J. Treatment Effect

Neoadjuvant chemoradiation therapy in rectal cancer is associated with significant tumor response and downstaging.¹ Because eradication of the tumor, as detected by pathologic examination of the resected specimen, is associated with a significantly better prognosis,² specimens from patients receiving neoadjuvant chemoradiation should be thoroughly sectioned, with careful examination of the tumor site. Minimal residual disease has been shown to have a better prognosis than gross residual disease.³ A modified Ryan scheme is suggested for scoring of tumor response, and has been shown to provide good interobserver reproducibility of prognostic significance.⁴ Several other systems have been studied and can be chosen to report the tumor regression score.

Modified Ryan Scheme for Tumor Regression Score

Description	Tumor Regression Score
No viable cancer cells (complete response)	0
Single cells or rare small groups of cancer cells (near complete response)	1
Residual cancer with evident tumor regression, but more than single cells or rare small groups of cancer cells (partial response)	2
Extensive residual cancer with no evident tumor regression (poor or no response)	3

Tumor regression should be assessed only in the primary tumor; lymph node metastases should not be included in the assessment.

Acellular pools of mucin in specimens following neoadjuvant therapy are considered to represent completely eradicated tumor and are not used to assign pT stage or counted as positive lymph nodes.

References

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K. Margins

It may be helpful to mark the margin(s) closest to the tumor with ink following close examination of the serosal surface for puckering and other signs of tumor involvement. Margins marked by ink should be designated in the macroscopic description of the surgical pathology report. The serosal surface (visceral peritoneum) does not constitute a surgical margin.

In addition to addressing the proximal and distal margins, the radial margin (Figure 3A-3C) must be assessed for any segment either unencased (Figure 3C) or incompletely encased by peritoneum (Figure 3B) (see Note A). The radial margin represents the adventitial soft tissue margin closest to the deepest penetration of tumor and is created surgically by blunt or sharp dissection of the retroperitoneal or subperitoneal aspect, respectively. Since the lower rectum is entirely extraperitoneal, the radial margin extends circumferentially and has been referred to as the circumferential radial margin. Multivariate analysis has suggested that tumor involvement of the radial margin is the most critical factor in predicting local recurrence in rectal cancer.¹ A positive radial margin in rectal cancer increases the risk of recurrence by 3.5-fold and doubles the risk of death from disease. For this reason, the radial margin should be assessed in all rectal carcinomas as well as colonic segments with nonperitonealized surfaces. The radial margin is considered negative if the tumor is more than 1 mm from the inked nonperitonealized surface but should be recorded as positive if tumor is located 1 mm or less from the nonperitonealized surface, because local recurrence rates are similar with clearances of 0 to 1 mm. There is limited outcome data for cases with intranodal or intravascular tumor within 1 mm of radial resection margin, but follow-up based

on a small number of patients suggests that local recurrence in these tumors may be similar to those with negative margin.^{1,2}

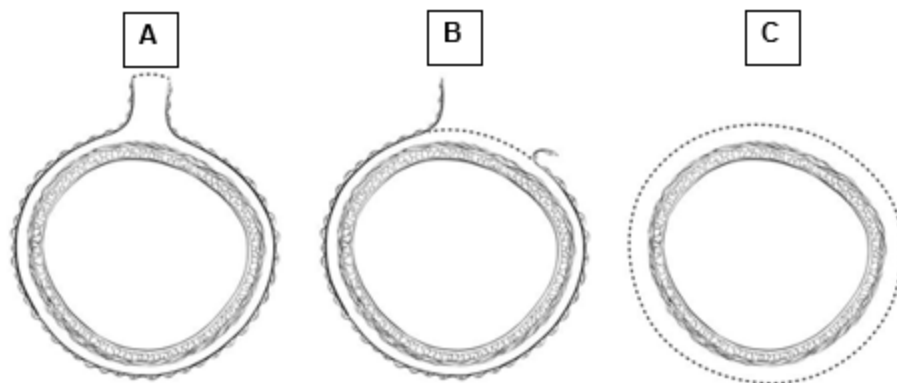


Figure 3. A, Mesenteric margin in portion of colon completely encased by peritoneum (dotted line). B, Radial margin (dotted line) in portion of colon incompletely encased by peritoneum. C, radial margin (dotted line) in rectum, completely unencased by peritoneum.

Sections to evaluate the proximal and distal resection margins can be obtained either by longitudinal sections perpendicular to the margin or by en face sections parallel to the margin. The distance from the tumor edge to the closest resection margin(s) may also be important, particularly for low anterior resections. For these cases, a distal resection margin of 2 cm is considered adequate; for T1 and T2 tumors, 1 cm may be sufficient distal clearance. Anastomotic recurrences are rare when the distance to the closest margin is 5 cm or greater.

In cases of carcinoma arising in a background of inflammatory bowel disease, proximal and distal resection margins should be evaluated for dysplasia and active inflammation. Proximal, distal, and radial/mesenteric resection margins should be reported in all resection specimens. Deep margin and mucosal margins should be reported in all transanal disk excisions.

References

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L. Tumor Deposits

A tumor focus in the pericolic/perirectal fat or in adjacent mesentery (mesocolic or rectal fat) within the lymph drainage area of the primary tumor, but without identifiable lymph node tissue or vascular structure. If the vessel wall or its remnant is identified (H&E, elastic, or any other stain), it should be classified as vascular (venous) invasion, and not as tumor deposit. Similarly, a tumor focus is present in or around a

large nerve, should be classified as perineural invasion and not as tumor deposit. Size and shape of the tumor focus are not relevant for classification as a tumor deposit.

The presence of tumor deposits in the absence of any regional node involvement is categorized as N1c, irrespective of T category. Tumor deposits are an adverse prognostic factor^{1,2} and adjuvant therapy is generally warranted in cases that are categorized as N1c regardless of T classification. If there are positive lymph nodes in addition to the tumor deposits, appropriate N1 (N1a or N1b) category should be used and not N1c, although the presence and number of tumor deposits should be noted in the synoptic summary (see note E).

In the setting of preoperative or neoadjuvant therapy, the designation of tumor deposit should be used with caution as the tumor foci may represent residual primary tumor with incomplete response.

References

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M. Ancillary Studies

Universal testing for microsatellite instability and/or status of DNA mismatch repair enzymes by immunohistochemistry is recommended by the EGAPP guidelines.^{1,2} MSI-high cancers are associated with right-sided location, tumor infiltrating lymphocytes, Crohn-like infiltrate, pushing borders, mucinous/signet ring/medullary subtypes, intratumoral heterogeneity (mixed conventional, mucinous, and poorly differentiated carcinoma), high-grade histology, and lack of dirty necrosis.^{3,4} In view of recommendations for universal testing, chance of missing cases of Lynch syndrome with testing based on Bethesda guidelines and implications for treatment with immune check-point inhibitors,⁴ evaluation of histologic features associated with MSI is redundant and is no longer included in the synoptic comment.

Further details about mismatch repair enzyme immunohistochemistry and PCR for MSI testing, as well as other ancillary molecular testing in colorectal cancer (such as *KRAS*, *BRAF*, *Her2*, etc.) can be found in the CAP Colon and Rectum Biomarkers protocol.

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