Protocol for the Examination of Specimens from Patients with Well-Differentiated Neuroendocrine Tumors (Carcinoid Tumors) of the Appendix

Version: 5.0.0.0
Protocol Posting Date: December 2023
CAP Laboratory Accreditation Program Protocol Required Use Date: September 2024

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:

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Excision</td>
<td>Includes specimens designated appendectomy, and right hemicolecotomy</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Tumor Type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well-differentiated neuroendocrine tumor</td>
<td></td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Biopsy</td>
<td>Primary resection specimen with no residual cancer (e.g., following neoadjuvant therapy)</td>
</tr>
<tr>
<td>Recurrent tumor</td>
<td></td>
</tr>
<tr>
<td>Cytologic specimens</td>
<td></td>
</tr>
</tbody>
</table>

The following tumor types should NOT be reported using this protocol:

<table>
<thead>
<tr>
<th>Tumor Type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poorly differentiated neuroendocrine carcinomas (including small cell and large cell neuroendocrine carcinomas) (consider the Appendix Carcinoma protocol)</td>
<td></td>
</tr>
<tr>
<td>Goblet cell adenocarcinoma (consider the Appendix Carcinoma protocol)</td>
<td></td>
</tr>
</tbody>
</table>

Authors
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With guidance from the CAP Cancer and CAP Pathology Electronic Reporting Committees.

* Denotes primary author.
Accreditation Requirements
This protocol can be utilized for a variety of procedures and tumor types for clinical care purposes. For accreditation purposes, only the definitive primary cancer resection specimen is required to have the core and conditional data elements reported in a synoptic format.

- Core data elements are required in reports to adequately describe appropriate malignancies. For accreditation purposes, essential data elements must be reported in all instances, even if the response is “not applicable” or “cannot be determined.”
- Conditional data elements are only required to be reported if applicable as delineated in the protocol. For instance, the total number of lymph nodes examined must be reported, but only if nodes are present in the specimen.
- Optional data elements are identified with “+” and although not required for CAP accreditation purposes, may be considered for reporting as determined by local practice standards.

The use of this protocol is not required for recurrent tumors or for metastatic tumors that are resected at a different time than the primary tumor. Use of this protocol is also not required for pathology reviews performed at a second institution (i.e., secondary consultation, second opinion, or review of outside case at second institution).

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 5.0.0.0

• Update to AJCC Version 9 pTNM Staging Classifications
• WHO 5th Edition update to content and explanatory notes
• “Lymphovascular Invasion” question updated to “Lymphatic and/or Vascular Invasion”
CASE SUMMARY: (APPENDIX NEUROENDOCRINE TUMOR)
Standard(s): AJCC-UICC 9

SPECIMEN

Procedure
___ Appendectomy
___ Right hemicolectomy
___ Other (specify): ____________________

TUMOR

Tumor Site (Note A)
___ Proximal half of appendix: _________________
___ Distal half of appendix: _________________
___ Diffusely involving appendix: _________________
___ Appendix, not otherwise specified: _________________
___ Other (specify): ____________________

Histologic Type and Grade# (Notes B,C)
# For poorly differentiated (high-grade) neuroendocrine carcinomas, the College of American Pathologists (CAP) checklist for carcinoma of the appendix should be used.
___ G1, well-differentiated neuroendocrine tumor
___ G2, well-differentiated neuroendocrine tumor
___ G3, well-differentiated neuroendocrine tumor
___ GX, grade cannot be assessed
___ Other (specify): ____________________
___ Not applicable: ____________________

†Histologic Type and Grade Comment: ____________________

Histologic Grade Determination (Note C)
Mitotic rate and / or Ki-67 labeling index is required to determine histologic grade

Mitotic Rate (required only when Ki-67 labeling index is not reported)#
# Mitotic rate should be reported as number of mitoses per 2 mm2, by evaluating at least 10 mm2 in the most mitotically active part of the tumor (e.g., if using a microscope with a field diameter of 0.55 mm, count 42 high power fields (10 mm2) and divide the resulting number of mitoses by 5 to determine the number of mitoses per 2 mm2 needed to assign tumor grade).
___ Not applicable (Ki-67 labeling index is reported)
___ Specify number of mitoses per 2 mm2: _________________ mitoses per 2 mm2
___ Less than 2 mitoses per 2 mm2
___ 2 to 20 mitoses per 2 mm2
___ Greater than 20 mitoses per 2 mm2
___ Cannot be determined (explain): ____________________
Ki-67 Labeling Index (required only when mitotic rate is not reported)
___ Not applicable (mitotic rate is reported)
___ Specify Ki-67 percentage: _________________ %
___ Less than 3%
___ 3% to 20%
___ Greater than 20%
___ Cannot be determined (explain): _________________

Tumor Size (Note D)
___ Greatest dimension in Centimeters (cm): _________________ cm
+Additional Dimension in Centimeters (cm): ____ x ____ cm
___ Cannot be determined (explain): _________________

Tumor Extent
___ Invades lamina propria
___ Invades submucosa
___ Invades muscularis propria
___ Invades subserosa or mesoappendix without involvement of visceral peritoneum
___ Perforates visceral peritoneum (serosa)
___ Directly invades other adjacent organ(s) or structure(s) (e.g., abdominal wall, skeletal muscle) excluding direct mural extension to adjacent subserosa of adjacent bowel (specify): _________________
___ Cannot be determined: _________________
___ No evidence of primary tumor

Lymphatic and / or Vascular Invasion
___ Not identified
___ Present
___ Cannot be determined: _________________

+Perineural Invasion
___ Not identified
___ Present
___ Cannot be determined: _________________

+Tumor Comment: _________________

MARGINS (Note E)

Margin Status
___ All margins negative for tumor
+Closest Margin(s) to Tumor (select all that apply)
___ Proximal: _________________
___ Distal: _________________
___ Radial: _________________
___ Mesenteric: _________________
___ Other (specify): _________________
___ Cannot be determined: ______________________

+Distance from Tumor 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: __________________
___ Tumor present at margin

Margin(s) Involved by Tumor (select all that apply)
___ Proximal: ______________________
___ Distal: ______________________
___ Radial: ______________________
___ Mesenteric: ______________________
___ 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): ______________________
+Regional Lymph Node Comment: ____________________

DISTANT METASTASIS

Distant Site(s) Involved, if applicable (select all that apply)
___ Not applicable
___ Liver: ___________________
___ Lung: ___________________
___ Ovary: ___________________
___ Nonregional lymph node(s): ___________________
___ Peritoneum: ___________________
___ Bone: ___________________
___ Other (specify): ___________________
___ Cannot be determined: ___________________

pTNM CLASSIFICATION (AJCC Version 9) (Note F)
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#
# Multiple tumors should be designated as such (the largest tumor should be used to assign T category). Use T(#); e.g., pT3(4) N0 M0, OR use the m suffix, T(m); e.g., pT3(m) N0 M0.
___ pT not assigned (cannot be determined based on available pathological information)
___ pT0: No evidence of primary tumor
___ pT1: Tumor less than or equal to 2 cm in greatest dimension
___ pT2: Tumor greater than 2 cm but less than or equal to 4 cm in greatest dimension
___ pT3: Tumor greater than 4 cm in greatest dimension, or with subserosal invasion, or involvement of the mesoappendix
___ pT4: Tumor perforates the peritoneum, or directly invades other adjacent organs or structures (excluding direct mural extension to adjacent subserosa of adjacent bowel), e.g., abdominal wall and skeletal muscle

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 tumor involvement of regional lymph node(s)
___ pN1: Tumor involvement of regional lymph node(s)
pM Category (required only if confirmed pathologically)

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

pM1: Microscopic confirmation of distant metastasis

___ pM1a: Microscopic confirmation of metastasis confined to liver

___ pM1b: Microscopic confirmation of metastasis in at least one extrahepatic site (e.g., lung, ovary, nonregional lymph node, peritoneum, bone)

___ pM1c: Microscopic confirmation of both hepatic and extrahepatic metastases

___ pM1 (subcategory cannot be determined)

ADDITIONAL FINDINGS (Note G)

+Additional Findings (select all that apply)

___ None identified

___ Tumor necrosis

___ Acute appendicitis

___ Other (specify): ____________________

COMMENTS

Comment(s): ____________________
Explanatory Notes

A. Application and Tumor Location
This protocol applies to well-differentiated neuroendocrine tumors (carcinoid tumors) of the appendix of any size. For poorly differentiated neuroendocrine carcinomas (including small cell carcinomas and large cell neuroendocrine carcinomas) and goblet cell adenocarcinoma (previously goblet cell carcinoid tumor), use the CAP cancer protocol for carcinoma of the appendix.1

The appendix is a common site of gastrointestinal neuroendocrine tumors, usually presenting as small solitary lesions incidentally discovered after appendectomy. Appendiceal neuroendocrine tumors are commonly diagnosed at a young age and arise in the deep mucosa or submucosa. Unlike other gastrointestinal neuroendocrine tumors or appendiceal adenocarcinomas, tumor size is considered more important than depth of invasion for appendiceal neuroendocrine tumor. Therefore, the staging system for appendiceal neuroendocrine tumors is different from those for other neuroendocrine tumors of the gut.2

References
2. AJCC Version 9 Neuroendocrine Tumors of the Appendix Cancer Staging System. Copyright 2023 American College of Surgeons.

B. Histologic Type
Most appendiceal NETs are low grade, with none to few mitoses and no necrosis, and have traditionally been classified as “carcinoids.” Although the term “carcinoid tumor” remains in widespread use, this term may cause confusion for clinicians, who might view a carcinoid tumor as a serotonin-producing tumor associated with functional manifestations of carcinoid syndrome. The use of the term “carcinoid” for neuroendocrine tumor reporting is therefore discouraged for these reasons.

Immunohistochemistry and other ancillary techniques are generally not required to diagnose well-differentiated neuroendocrine tumors. Specific markers that may be used to establish neuroendocrine differentiation include chromogranin A, synaptophysin, INSM1 and CD56.1,2,3 Because of their relative sensitivity and specificity, chromogranin A and synaptophysin are recommended.

Although specific histologic patterns in well-differentiated neuroendocrine tumors, such as trabecular, insular, and glandular, roughly correlate with tumor location,4 these patterns have not been clearly shown independently to predict response to therapy or risk of nodal metastasis and are rarely reported in clinical practice. Most appendiceal neuroendocrine tumors are derived from enterochromaffin cells. Rarely, L-cell neuroendocrine tumors of the appendix are encountered; because of their distinctive growth pattern of tear-drop-shaped tubules embedded in a fibrous stroma,5 these lesions are sometimes called tubular neuroendocrine tumors. It should be noted that these tumors are negative for chromogranin A but express enteroglucagon, peptide YY, and pancreatic polypeptide. Tubular neuroendocrine tumors are usually small lesions confined to the appendix and are found in female patients. These lesions exhibit benign behavior and should not be confused with adenocarcinoma.
References

C. Histologic Grade
Cytologic atypia in well differentiated neuroendocrine tumors has no impact on clinical behavior of these tumors. The WHO classification¹ and others² use mitotic rate and/or Ki-67 index as one of the criteria for potential for aggressive behavior. Mitotic rate should be reported as number of mitoses per 2 mm², by evaluating at least 10 mm² in the most mitotically active part of the tumor. Only clearly identifiable mitotic figures should be counted; hyperchromatic, karyorrhectic, or apoptotic nuclei are excluded. Because of variations in field size, the number of high-power fields (HPF) (at 40X magnification) for10 mm² (thereby 2 mm²) must be determined for each microscope (Table 1). For example, if using a microscope with a field diameter of 0.55 mm, count 42 HPF and divide the resulting number of mitoses by 5 to determine the number of mitoses per 2 mm² needed to assign tumor grade.

Table 1. Number of HPF Required for 10 mm² Using Microscopes With Different Field Diameter
<table>
<thead>
<tr>
<th>Field Diameter (mm)</th>
<th>Area (mm²)</th>
<th>Number of HPF for 10 mm²</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.40</td>
<td>0.125</td>
<td>80</td>
</tr>
<tr>
<td>0.41</td>
<td>0.132</td>
<td>75</td>
</tr>
<tr>
<td>0.42</td>
<td>0.139</td>
<td>70</td>
</tr>
<tr>
<td>0.43</td>
<td>0.145</td>
<td>69</td>
</tr>
<tr>
<td>0.44</td>
<td>0.152</td>
<td>65</td>
</tr>
<tr>
<td>0.45</td>
<td>0.159</td>
<td>63</td>
</tr>
<tr>
<td>0.46</td>
<td>0.166</td>
<td>60</td>
</tr>
<tr>
<td>0.47</td>
<td>0.173</td>
<td>58</td>
</tr>
<tr>
<td>0.48</td>
<td>0.181</td>
<td>55</td>
</tr>
<tr>
<td>0.49</td>
<td>0.189</td>
<td>53</td>
</tr>
<tr>
<td>0.50</td>
<td>0.196</td>
<td>50</td>
</tr>
<tr>
<td>0.51</td>
<td>0.204</td>
<td>49</td>
</tr>
<tr>
<td>0.52</td>
<td>0.212</td>
<td>47</td>
</tr>
<tr>
<td>0.53</td>
<td>0.221</td>
<td>45</td>
</tr>
<tr>
<td>0.54</td>
<td>0.229</td>
<td>44</td>
</tr>
<tr>
<td>0.55</td>
<td>0.238</td>
<td>42</td>
</tr>
<tr>
<td>0.56</td>
<td>0.246</td>
<td>41</td>
</tr>
<tr>
<td>0.57</td>
<td>0.255</td>
<td>39</td>
</tr>
<tr>
<td>0.58</td>
<td>0.264</td>
<td>38</td>
</tr>
</tbody>
</table>
Ki-67 index is reported as percent positive tumor cells in area of highest nuclear labeling (“hot spot”), although the precise method of assessment has not been standardized. A number of methods have used to assess Ki-67 index, including automatic counting and “eyeballing”. Automated counting is not widely available and requires careful modification of the software to circumvent the inaccuracies. Eye-balling can be used for most tumors; however, for tumors with Ki-67 index close to grade cut-offs, it is recommended to perform the manual count on the print of camera-captured image of the hot spot. It has been recommended that a minimum of 500 tumor cells be counted to determine the Ki-67 index, and a notation is made if less cells are available. Grade assigned based on Ki-67 index is typically higher than that based on mitotic count, and the case is assigned to the higher of the 2 if both methods are performed.

It is important to note that there are a small group of well-differentiated neuroendocrine tumors with a Ki-67 index >20% and a mitotic rate usually <20 per 10 HPF. In WHO-2010, these tumors were considered as G3 poorly differentiated neuroendocrine carcinomas. However, they have typical morphology of well-differentiated tumors.

Previous studies (most on pancreatic neuroendocrine tumors) have demonstrated that these tumors have a worse prognosis than grade 2 (Ki-67=3-20 % and mitosis <20/10 HPF) neuroendocrine tumors, but they are not as aggressive as poorly differentiated neuroendocrine carcinomas. In addition, these tumors do not have the genetic abnormalities seen in poorly differentiated neuroendocrine carcinomas. Furthermore, unlike poorly differentiated neuroendocrine carcinomas, they are less responsive to platinum-based chemotherapy. In the WHO-2019 blue book of digestive system tumors and the AJCC Version 9, those with typical morphology of well-differentiated tumors are classified as “well-differentiated neuroendocrine tumor” but as grade 3 (Table 2).

Table 2 Recommended Grading System for Well-Differentiated Gastroenteropancreatic Neuroendocrine Tumors

<table>
<thead>
<tr>
<th>Grade</th>
<th>Mitotic Rate (per 2mm²)</th>
<th>Ki-67 index (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well-differentiated neuroendocrine tumor, G1</td>
<td>&lt;2</td>
<td>&lt;3</td>
</tr>
<tr>
<td>Well-differentiated neuroendocrine tumor, G2</td>
<td>2-20</td>
<td>3-20</td>
</tr>
<tr>
<td>Well-differentiated neuroendocrine tumor, G3</td>
<td>&gt;20</td>
<td>&gt;20</td>
</tr>
</tbody>
</table>

References
Appendiceal neuroendocrine tumors smaller than 1.0 cm do not recur or metastasize, whereas those between 1.0 and 2.0 cm rarely do. A recent multi-center retrospective study on a large cohort of appendiceal NET between 1-2cm suggest that right hemicolectomy is not needed and further medical imaging or assessment of histopathologic risk factors provides no added advantage. Tumor size >2.0 cm and mesoappendiceal invasion have been correlated with nodal metastasis, but not with poor outcome. For these reasons, appendectomy is usually sufficient for tumors 1.0 cm or smaller, as well as many tumors between 1.0 and 2.0 cm. More extensive procedures (e.g., right hemicolectomy) are usually reserved for patients with tumors larger than 2.0 cm or with invasion beyond the muscularis propria. Most appendiceal neuroendocrine tumors are identified incidentally, usually for the first time on histology but sometimes during gross examination. When initially identified on histology, the entire appendix including the surgical resection margin should be submitted for histology. In this scenario, the size is estimated by number of slides involved by tumor and approximate thickness of tissue slice in each block. While the thickness of the tissue slices tends to vary between labs, prosectors and cases, these typically range between 2-5mm and an average thickness of 3mm can be used to estimate the tumor size. Between gross examination and histologic evaluation, the largest size estimated should be ascribed to a given case for staging.

References
E. Margins
Margins in a simple appendectomy specimen include the proximal and circumferential (radial) margin. It is recommended that the proximal margin on a simple appendectomy specimen should be taken en face in order to evaluate the entire appendiceal mucosa and muscularis circumferentially. In the vast majority of cases, the appendix is entirely peritonealized (Figure, A), and the closest distance between the tumor and the mesoappendiceal resection margin represents the radial margin and should be measured. Even retrocecal appendices are usually invested by peritoneum but have adhered to the posterior cecum, either because of inflammation or tumor. Exceptionally, a retrocecal appendix may be retroperitoneal, in which case the distance between the tumor and the nonperitonealized radial resection margin is the “surgical clearance” and should be measured.

In general, the circumferential (radial) margin must be assessed for any segment of gastrointestinal tract either incompletely encased or unencased by peritoneum (Figure, B and C). The posterior surface of the ascending colon portion of a right hemicolectomy specimen lacks a peritoneal covering and thus constitutes a circumferential margin, which in rare cases may be relevant in right hemicolectomy specimens resected for treatment of appendiceal neuroendocrine tumors. The circumferential (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. The distance between the tumor and circumferential (radial) margin should be reported, if applicable. The circumferential (radial) margin is considered positive if the tumor is present at the inked nonperitonealized surface. This assessment includes tumor within a lymph node as well as direct tumor extension, but if circumferential (radial) margin positivity is based solely on intranodal tumor, this should be so stated.

The mesenteric resection margin is the only relevant circumferential margin in segments completely encased by peritoneum (e.g., cecum; mesoappendiceal resection margin for the appendix) (Figure, A). Involvement of this margin should be reported even if tumor does not penetrate the serosal surface.
A: Mesenteric margin in viscus completely encased by peritoneum (dotted line). B: Circumferential (radial) margin (dotted line) in viscus incompletely encased by peritoneum. C: Circumferential (radial) margin (dotted line) in viscus completely unencased by peritoneum.

F. pTNM Classification
The TNM staging system for appendiceal NETs of the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC) is recommended.1

pTNM Pathologic Classification
The pT, pN, and pM categories correspond to the T, N, and M categories except that pM0 (no distant metastasis) does not exist as a category.

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 (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.

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 (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.

The “a” prefix designates the stage determined at autopsy: aTNM.
T Category Considerations
Tumor that is adherent to other organs or structures, macroscopically, is classified cT4. However, if no tumor is present in the adhesion and there is no serosal penetration microscopically, the classification should be pT1 to pT3.

N Category Considerations
The regional lymph nodes for the appendix are the ileocolic lymph nodes. Histologic examination of a regional lymphadenectomy specimen will ordinarily include 12 or more lymph nodes. If the lymph node results are negative, but the number ordinarily examined is not met, classify as pN0.

References
1. AJCC Version 9 Neuroendocrine Tumors of the Appendix Cancer Staging System. Copyright 2023 American College of Surgeons.

G. Additional Findings
Coagulative tumor necrosis, usually punctate, may indicate more aggressive behavior and should be reported. Appendiceal NETs are often an incidental finding in specimens removed for acute appendicitis.

References