Protocol for the Examination of Specimens From Patients With Neuroendocrine Tumors (Carcinoid Tumors) of the Stomach

Protocol applies to well-differentiated neuroendocrine tumors of the stomach. Carcinomas with mixed endocrine/glandular differentiation, poorly differentiated carcinomas with neuroendocrine features, and small cell carcinomas are not included.

Based on AJCC/UICC TNM, 7th Edition
Protocol web posting date: June 2012

 Procedures
• Endoscopic Resection
• Gastrectomy (Partial or Complete)

 Authors
Laura H. Tang, MD, PhD, FCAP*
   Department of Pathology, Memorial Sloan-Kettering Cancer Center, New York, NY
Jordan Berlin, MD
   Department of Medicine, Vanderbilt University Medical Center, Nashville, TN
Philip Branton, MD, FCAP
   Department of Pathology, Inova Fairfax Hospital, Falls Church, VA
Lawrence J. Burgart, MD, FCAP
   Allina Laboratories, Abbott Northwestern Hospital, Minneapolis, MN
David K. Carter, MD, FCAP
   Department of Pathology, St. Mary’s/Duluth Clinic Health System, Duluth, MN
Carolyn C. Compton, MD, PhD, FCAP
   Critical Path Institute, Tucson, AZ
Patrick Fitzgibbons, MD, FCAP
   Department of Pathology, St. Jude Medical Center, Fullerton, CA
Wendy L. Frankel, MD, FCAP
   Department of Pathology, Ohio State University Medical Center, Columbus, OH
John Jessup, MD
   Division of Cancer Treatment and Diagnosis, National Cancer Institute, Bethesda, MD
Sanjay Kakar, MD, FCAP
   Department of Pathology, University of California San Francisco and the Veterans Affairs Medical Center, San Francisco, CA
Bruce Minsky, MD
   Department of Radiation Oncology, University of Chicago, Chicago, IL
Raouf Nakhleh, MD, FCAP
   Department of Pathology, Mayo Clinic, Jacksonville, FL
Kay Washington, MD, PhD, FCAP†
   Department of Pathology, Vanderbilt University Medical Center, Nashville, TN
For the Members of the Cancer Committee, College of American Pathologists

* Denotes primary author. † Denotes senior author. All other contributing authors are listed alphabetically.
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CAP Stomach NET Protocol Revision History

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

Version: StomachNET 3.2.0.0

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

Endoscopic Resection, Gastrectomy

Histologic Type; Alternate Histologic Classification; Histologic Grade
These three reporting elements were combined into one, and the word “checklist” was changed to "protocol" as follows:

Histologic Type and Grade

___ Not applicable
___ Well-differentiated neuroendocrine tumor; GX: Grade cannot be assessed
___ Well-differentiated neuroendocrine tumor; G1: Low grade (carcinoid)
___ Well-differentiated neuroendocrine tumor; G2: Intermediate grade (atypical carcinoid)
___ Other (specify): ____________________________

*For poorly differentiated (high-grade) neuroendocrine carcinomas (G3), the College of American Pathologists (CAP) protocol for carcinoma of the stomach should be used.

Margins
“Lateral Mucosal Margins” was changed to “Mucosal Margins.” Deep Margin and Mucosal Margins: added “required only if applicable,” changed “Uninvolved/Involved by tumor” to “Uninvolved/Involved by neuroendocrine tumor,” and deleted “Not applicable” data element. Other Margins: added “required only if applicable” and deleted “Not applicable” data element, as follows:

Deep Margin (a endoscopic resections) (required only if applicable)
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor

Mucosal Margins (endoscopic resections) (required only if applicable)
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor

Other Margin(s) (required only if applicable)
Specify margin(s): ____________________________
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor
+ ___ Involved by neuroendocrine cell hyperplasia/dysplasia

Ancillary Studies
Reporting on Ki-67 was updated, changing >2% to 3% and adding “specify,” as follows:
+ ___ Ki-67 labeling index (specify: _____)
  + ___ ≤2%
  + ___ 3% to 20%
  + ___ >20%
Explanatory Notes
"Neoplasm" was changed to "tumor."

D. Histologic Type
The first sentence was changed to the following:
The World Health Organization (WHO) classifies neuroendocrine neoplasms as well-differentiated neuroendocrine tumors (either the primary tumor or metastasis) and poorly differentiated neuroendocrine carcinomas.⁵⁻⁸

Alternative Classification Based Upon WHO Classification: Neuroendocrine Tumors of the Appendix
This section was deleted.

E. Histologic Grade
"Mitotic count" was changed to "mitotic rate."
Ki-67 Index (%): >2 was changed to 3.
The last sentence of the first paragraph was changed to:
- The following grading system is recommended by both the European Neuroendocrine Tumor Society (ENETS) and the WHO⁸,⁹:

References
Reference #8 was updated.
Surgical Pathology Cancer Case Summary

Protocol web posting date: June 2012

STOMACH: Endoscopic Resection, Gastrectomy (Note A)

Select a single response unless otherwise indicated.

Specimen (select all that apply)
___ Stomach
___ Portion of stomach
   ___ Gastric body
   ___ Gastric antrum
   ___ Not specified
___ Distal esophagus
___ Proximal duodenum
___ Other (specify): ____________________________
___ Not specified

Procedure
___ Endoscopic resection
___ Partial gastrectomy, proximal
___ Partial gastrectomy, distal
___ Partial gastrectomy, other (specify): ____________________________
___ Total gastrectomy
___ Other (specify): ____________________________
___ Not specified

+ Specimen Size (if applicable)
+ Specify: ___ (length) x ___ x ___ cm

Tumor Site (select all that apply) (Note B)
___ Gastric cardia
___ Gastric fundus
___ Gastric body
___ Gastric antrum
___ Other (specify): ____________________________
___ Not specified

Tumor Size (Note C)
Greatest dimension: ___ cm (specify size of largest tumor if multiple tumors are present)
+ Additional dimensions: ___ x ___ cm
___ Cannot be determined (see “Comment”)

Tumor Focality
___ Unifocal
___ Multifocal (specify number of tumors: _____)
___ Cannot be determined

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Histologic Type and Grade (Notes D and E)*

___ Not applicable
___ Well-differentiated neuroendocrine tumor; GX: Grade cannot be assessed
___ Well-differentiated neuroendocrine tumor; G1: Low grade (carcinoid)
___ Well-differentiated neuroendocrine tumor; G2: Intermediate grade (atypical carcinoid)
___ Other (specify): ____________________________

* For poorly differentiated (high-grade) neuroendocrine carcinomas (G3), the College of American Pathologists (CAP) protocol for carcinoma of the stomach should be used.

Mitotic Rate (Note E)
Specify: ___/10 high-power fields (HPF)
___ Cannot be determined

Microscopic Tumor Extension
___ Cannot be assessed
___ No evidence of primary tumor
___ Tumor invades lamina propria
___ Tumor invades into but not through muscularis mucosae
___ Tumor invades submucosa
___ Tumor invades muscularis propria
___ Tumor invades subserosal tissue without involvement of visceral peritoneum
___ Tumor penetrates serosa (visceral peritoneum)
___ Tumor directly invades adjacent structures (specify: _______________)
___ Tumor penetrates to the surface of the visceral peritoneum (serosa) and directly invades adjacent structures (specify: _______________)

Margins (select all that apply)
If all margins uninvolved by neuroendocrine tumor:
   Distance of tumor from closest margin: ___ mm or ___ cm
   Specify margin: _________________________

Proximal Margin
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor
+ ___ Involved by neuroendocrine cell hyperplasia/dysplasia

Distal Margin
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor
+ ___ Involved by neuroendocrine cell hyperplasia/dysplasia

Omental (Radial) Margin (Note F)
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Deep Margin (endoscopic resections) (required only if applicable)
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor

Mucosal Margins (endoscopic resections) (required only if applicable)
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor

Other Margin(s) (required only if applicable)
Specify margin(s): ______________________
___ Cannot be assessed
___ Uninvolved by neuroendocrine tumor
___ Involved by neuroendocrine tumor
+ ___ Involved by neuroendocrine cell hyperplasia/dysplasia

Lymph-Vascular Invasion
___ Not identified
___ Present
___ Indeterminate

+ Perineural Invasion
+ ___ Not identified
+ ___ Present
+ ___ Indeterminate

Pathologic Staging (pTNM) (Note G)

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

Primary Tumor (pT)
___ pTX: Primary tumor cannot be assessed
___ pT0: No evidence of primary tumor
___ pTis: Carcinoma in situ/dysplasia (tumor size less than 0.5 mm), confined to mucosa
___ pT1: Tumor invades lamina propria or submucosa and 1 cm or less in size
___ pT2: Tumor invades muscularis propria or more than 1 cm in size
___ pT3: Tumor penetrates subserosa
___ pT4: Tumor invades visceral peritoneum (serosal) or other organs or adjacent structures

Regional Lymph Nodes (pN)
___ Cannot be assessed
___ pN0: No regional lymph node metastasis
___ pN1: Metastasis in regional lymph nodes
___ No nodes submitted or found

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
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StomachNET 3.2.0.0

*Number of Lymph Nodes Examined*
Specify: ___
___ Number cannot be determined (explain): ______________________

*Number of Lymph Nodes Involved*
Specify: ___
___ Number cannot be determined (explain): ______________________

*Distant Metastasis (pM)*
___ Not applicable
___ pM1: Distant metastasis
   + Specify site(s), if known: __________________________

+ **Ancillary Studies (select all that apply) (Notes E and H)**
  + ___ Ki-67 labeling index (specify: _____)
    + ___ ≤2%
    + ___ 3% to 20%
    + ___ >20%
  + ___ Other (specify): __________________________
  + ___ Not performed

+ **Additional Pathologic Findings (select all that apply) (Note I)**
  + ___ Atrophic gastritis
  + ___ Intestinal metaplasia of gastric mucosa
  + ___ Glandular dysplasia of gastric mucosa
  + ___ Endocrine cell hyperplasia
  + ___ Absence of parietal cells
  + ___ Tumor necrosis
  + ___ Other, specify: __________________________

+ **Comment(s)**

* Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
A. Application and Tumor Location
This protocol applies to well-differentiated neuroendocrine tumors (carcinoid tumors) of the stomach. Poorly differentiated neuroendocrine carcinomas, small cell carcinomas, and tumors with mixed glandular/neuroendocrine differentiation are not included.

Because of site-specific similarities in histology, immunohistochemistry, and histochemistry, neuroendocrine tumors of the digestive tract have traditionally been subdivided into those of foregut, midgut, and hindgut origin (Table 1). In general, the distribution pattern along the gastrointestinal (GI) tract parallels that of the progenitor cell type, and the anatomic site of origin of GI neuroendocrine tumors is an important predictor of clinical behavior.²

Table 1. Site of Origin of Gastrointestinal Neuroendocrine Tumors

<table>
<thead>
<tr>
<th>Site</th>
<th>Foregut Tumors</th>
<th>Midgut Tumors</th>
<th>Hindgut Tumors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Site</td>
<td>Stomach, Proximal</td>
<td>Jejunum, Ileum,</td>
<td>Distal Colon, Rectum</td>
</tr>
<tr>
<td></td>
<td>Duodenum</td>
<td>Appendix, Proximal</td>
<td></td>
</tr>
<tr>
<td>Immunohistochemistry</td>
<td>86%-100% + ⁹</td>
<td>82%-92% +</td>
<td>40%-58% +</td>
</tr>
<tr>
<td>Chromogranin A</td>
<td>90%-100% +</td>
<td>95%-100% +</td>
<td>80%-87% +</td>
</tr>
<tr>
<td>Neuron-Specific Enolase (NSE)</td>
<td>50% +</td>
<td>95%-100% +</td>
<td>94%-100% +</td>
</tr>
<tr>
<td>Synaptophysin</td>
<td>33% + ¹¹,¹²</td>
<td>86% + ¹¹,¹²</td>
<td>45%-83% + ³ ⁵,¹²</td>
</tr>
<tr>
<td>Serotonin</td>
<td>Rare, + for pancreatic</td>
<td>Prostatic acid</td>
<td>Prostatic acid</td>
</tr>
<tr>
<td></td>
<td>polypeptide, histamine,</td>
<td>phosphatase + in</td>
<td>phosphatase + in</td>
</tr>
<tr>
<td></td>
<td>gastrin, somatostatin,</td>
<td>20%-40% ¹¹,¹²</td>
<td>20%-82% ³ ⁵,¹²</td>
</tr>
<tr>
<td></td>
<td>vasoactive intestinal</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>peptide (VIP), or</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>adrenocorticotropic</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>hormone (ACTH)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>Rare</td>
<td>5%-39% ⁶,⁷</td>
<td>Rare</td>
</tr>
<tr>
<td>Immunohistochemical</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Markers</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Carcinoid Syndrome</td>
<td>Rare</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

B. Site-Specific Features
Gastric neuroendocrine tumors are divided into 4 types.³ Type 1 tumors arising in the setting of atrophic gastritis with associated hypergastrinemia are the most common. These lesions are composed of enterochromaffin-like (ECL) cells and are usually found as multiple small nodules in the body of the stomach and limited to mucosa and submucosa. Type 1 lesions are generally benign and may regress following antrectomy; lymph node metastases are very rare and occur only when the tumors are large (greater than 2 cm) and infiltrate the muscularis propria.

Type 2 gastric neuroendocrine tumors are rare. These multifocal small tumors, which are associated with multiple endocrine neoplasia (MEN) type 1 with Zollinger-Ellison syndrome, develop in the body of the stomach, are usually smaller than 1.5 cm, and are confined to the mucosa or submucosa. However, in contrast to type 1 tumors, 30% metastasize. Tumors greater than 2 cm and invading the muscularis propria and exhibiting vascular invasion are more likely to metastasize.

Type 3 gastric neuroendocrine tumors, the second most common neuroendocrine tumor in the stomach, are sporadic solitary tumors that are unassociated with atrophic gastritis or endocrine cell hyperplasia. These tumors may occur anywhere in the stomach. Metastasis is associated with larger mean size, angioinvasion, and invasion of muscularis propria. Surgical resection is usually advised for
solitary gastric carcinoid tumors, particularly those larger than 2.0 cm, but tumors smaller than 1.0 cm have been rarely reported to metastasize.\(^4\)

Type 4 gastric neuroendocrine tumors are rare high-grade neuroendocrine carcinomas that are usually bulky tumors with metastases at diagnosis (the CAP cancer protocol for gastric carcinoma applies\(^1\)).

C. Tumor Size
For neuroendocrine tumors in any part of the gastrointestinal tract, size greater than 2.0 cm is associated with a higher risk of lymph node metastasis. In the stomach, types 3 and 4 neuroendocrine tumors are significantly larger than type 1 tumors, which usually measure 1 cm or less (Table 2). Tumor size correlates with depth of invasion for gastric neuroendocrine tumors, with larger tumors more likely to be deeply infiltrative and thus at higher risk for metastases. Nodules measuring 0.5 mm or larger are defined as neuroendocrine tumors; lesions measuring less than 0.5 mm are regarded as representing in situ tumor, neuroendocrine cell dysplasia, or hyperplasia.

### Table 2. Types of Gastric Neuroendocrine Tumors

<table>
<thead>
<tr>
<th>Type 1</th>
<th>Type 2</th>
<th>Type 3</th>
<th>Type 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frequency</td>
<td>70%-80% of cases</td>
<td>Rare</td>
<td>10%-15% of cases</td>
</tr>
<tr>
<td>Multiplicity</td>
<td>Multifocal</td>
<td>Multifocal</td>
<td>Solitary</td>
</tr>
<tr>
<td>Size</td>
<td>0.5-1.0 cm</td>
<td>~1.5 cm or less</td>
<td>Variable; one-third are larger than 2 cm</td>
</tr>
<tr>
<td>Location</td>
<td>Corpus</td>
<td>Corpus</td>
<td>Anywhere in stomach</td>
</tr>
<tr>
<td>Associations</td>
<td>Hypergastrinemic states; chronic atrophic gastritis, enterochromaffin-like (ECL) cell hyperplasia, pernicious anemia</td>
<td>Multiple endocrine neoplasia (MEN) type 1, with hypergastrinemia or Zollinger-Ellison syndrome</td>
<td>Sporadic</td>
</tr>
<tr>
<td>Clinical Behavior</td>
<td>Usually benign</td>
<td>30% metastasize</td>
<td>71% of tumors &gt;2 cm with muscularis propria and vascular invasion have lymph node metastases</td>
</tr>
<tr>
<td>Demographic Profile</td>
<td>70%-80% are females in their 50s and 60s</td>
<td>Equally in males and females, mean age 50 y</td>
<td>More common in males, mean age 55 y</td>
</tr>
</tbody>
</table>

D. Histologic Type
The World Health Organization (WHO) classifies neuroendocrine neoplasms as well-differentiated neuroendocrine tumors (either the primary tumor or metastasis) and poorly differentiated neuroendocrine carcinomas.\(^5-8\) Historically, well-differentiated neuroendocrine tumors have been referred to as carcinoid tumors, a term which may cause confusion because clinically a carcinoid tumor is a serotonin-producing tumor associated with functional manifestations of carcinoid syndrome.

Classification of neuroendocrine tumors (NETs) is based upon size, functionality, site, and invasion. Functioning tumors are those associated with clinical manifestations of hormone production or secretion of measurable amounts of active hormone; immunohistochemical demonstration of hormone production is not equivalent to clinically apparent functionality.
Histologic Patterns
Although specific histologic patterns in well-differentiated neuroendocrine tumors, such as trabecular, insular, and glandular, roughly correlate with tumor location, 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.

E. Histologic Grade
Cytologic atypia in low-grade neuroendocrine tumors has no impact on clinical behavior of these tumors. However, grading systems based on mitotic activity have been shown to have utility for foregut tumors. The following grading system is recommended by both the European Neuroendocrine Tumor Society (ENETS) and the WHO:

<table>
<thead>
<tr>
<th>Grade</th>
<th>Mitotic Rate (per 10 HPF)*</th>
<th>Ki-67 Index (%)##</th>
</tr>
</thead>
<tbody>
<tr>
<td>G1</td>
<td>&lt;2</td>
<td>≤2</td>
</tr>
<tr>
<td>G2</td>
<td>2 to 20</td>
<td>3 to 20</td>
</tr>
<tr>
<td>G3</td>
<td>&gt;20</td>
<td>&gt;20</td>
</tr>
</tbody>
</table>

*Mitotic rate should be based upon counting 50 high-power (40x objective) fields in the area of highest mitotic activity and reported as number of mitoses per 10 HPF.

##Ki-67 index is reported as percentage of positive tumor cells in area of highest nuclear labeling. It has been recommended that 2000 tumor cells be counted to determine the Ki-67 index; however, this practice may not be practical for routine clinical purposes, and it is acceptable to estimate the labeling index.

G1 and G2 are well-differentiated tumors with diffuse intense chromogranin/synaptophysin positivity. Punctate necrosis is more typical of G2 tumors. G3 tumors are high-grade neuroendocrine carcinomas (the CAP carcinoma protocol for carcinomas of the stomach applies).

F. Circumferential (Radial) Margin
For surgical resection specimens, margins include the proximal, distal, and radial margins. The radial margins represent the nonperitonealized soft tissue margins closest to the deepest penetration of tumor. In the stomach, the lesser omental (hepatoduodenal and hepatogastric ligaments) and greater omental resection margins are the only radial margins. For endoscopic resection specimens, margins include mucosal margins and the deep margin of resection. It may be helpful to mark the margin(s) closest to the tumor with ink. Margins marked by ink should be designated in the macroscopic description.

G. TNM and Anatomic Stage/Prognostic Groupings
The TNM staging system for gastric neuroendocrine tumors of the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC) is recommended.

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

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

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

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

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

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

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

N Category Considerations
The specific nodal areas of the stomach are listed below.

**Greater curvature of stomach:** Greater curvature, greater omental, gastroduodenal, gastroepiploic, pyloric, and pancreaticoduodenal

**Pancreatic and splenic area:** Pancreaticolienal, peripancreatic, splenic

**Lesser curvature of stomach:** Lesser curvature, lesser omental, left gastric, cardioesophageal, common hepatic, celiac, and hepatoduodenal

Involvement of other intra-abdominal lymph nodes, such as hepatoduodenal, retropancreatic, mesenteric, and para-aortic, is classified as distant metastasis.

**TNM Anatomic Stage/Prognostic Groupings**

<table>
<thead>
<tr>
<th>Stage</th>
<th>T category</th>
<th>N category</th>
<th>M category</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>Tis</td>
<td>N0</td>
<td>M0*</td>
</tr>
<tr>
<td>I</td>
<td>T1</td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>IIa</td>
<td>T2</td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>IIb</td>
<td>T3</td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>IIIa</td>
<td>T4</td>
<td>N0</td>
<td>M0</td>
</tr>
<tr>
<td>IIIb</td>
<td>Any T</td>
<td>N1</td>
<td>M0</td>
</tr>
<tr>
<td>IV</td>
<td>Any T</td>
<td>Any N</td>
<td>M1</td>
</tr>
</tbody>
</table>

* M0 is defined as no distant metastasis.

H. Ancillary Studies
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, neuron-specific enolase, synaptophysin, and CD56. Because of their relative sensitivity and specificity, chromogranin A and synaptophysin are recommended.
Immunohistochemistry for Ki-67 may be useful in establishing tumor grade (Note E) and prognosis but is not currently considered standard of care.

Immunohistochemistry for specific hormone products, such as glucagon, gastrin, and somatostatin, may be of interest in some cases. However, immunohistochemical demonstration of hormone production does not equate with clinical functionality of the tumor.

I. Additional Pathologic Findings

Most gastric neuroendocrine tumors arise in the setting of chronic atrophic gastritis (see Note B). Atrophic gastritis may be associated with glandular dysplasia, and in rare cases, gastric adenocarcinoma. Coagulative tumor necrosis, usually punctate, may indicate more aggressive behavior and should be reported.

References