Protocol for the Examination of Specimens From Patients With Carcinoma of the Esophagus

Protocol applies to all carcinomas of the esophagus, including esophagogastric junction carcinomas. Well-differentiated neuroendocrine tumors (carcinoid tumors) are not included.

Based on AJCC/UICC TNM, 7th edition
Protocol web posting date: January 2016

Procedures
- Endoscopic Resection
- Esophagectomy
- Esophagogastrectomy

Authors
Kay Washington, MD, PhD*
Department of Pathology, Microbiology and Immunology, Vanderbilt University Medical Center, Nashville, TN
Jordan Berlin, MD
Department of Medicine, Vanderbilt University Medical Center, Nashville, TN
Philip Branton, MD
Department of Pathology, Inova Fairfax Hospital, Falls Church, VA
Lawrence J. Burgart, MD
Allina Laboratories, Abbott Northwestern Hospital, Minneapolis, MN
David K. Carter, MD
Department of Pathology, St. Mary’s/Duluth Clinic Health System, Duluth, MN
Patrick Fitzgibbons, MD
Department of Pathology, St. Jude Medical Center, Fullerton, CA
Wendy L. Frankel, MD
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
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
Department of Pathology, Mayo Clinic, Jacksonville, FL
Chanjuan Shi, MD, PhD
Department of Pathology, Microbiology and Immunology, Vanderbilt University Medical Center, Nashville, TN
Laura H. Tang, MD, PhD
Department of Pathology, Memorial Sloan-Kettering Cancer Center, New York, NY
Carolyn C. Compton, MD, PhD†
Critical Path Institute, Tucson, AZ
For the Members of the Cancer Committee, College of American Pathologists

* Denotes primary author. † Denotes senior author. All other contributing authors are listed alphabetically.

Previous contributors: Randall G. Lee, MD; Leslie H. Sobin, MD; Donald Antonioli, MD; Harvey Goldman, MD; Rodger C. Haggitt, MD; Robert V. P. Hutter, MD; Klaus Lewin, MD; Pablo Ross, MD; Heidrun Rotterdam, MD; Stuart Spechler, MD; Christopher Willett, MD; Donald E. Henson, MD
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CAP Esophagus Protocol Revision History

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

Version: Esophagus 3.2.0.0

Summary of Changes
The following changes have been made since the October 2013 release.

The following data elements have been modified:
- Histologic Type
- Margins: Distal Margin
- Treatment Effect
- Lymph-Vascular Invasion
- Perineural Invasion
- Distant Metastasis (changed to required only if confirmed pathologically)
- Ancillary Studies (added note)

The following data element was added:
- Margins: Mucosal Margin
Surgical Pathology Cancer Case Summary

Protocol web posting date: January 2016

ESOPHAGUS: Endoscopic Resection, Esophagectomy, or Esophagogastrectomy (Note A)

Select a single response unless otherwise indicated.

Specimen (select all that apply)
___ Esophagus
___ Proximal stomach
___ Other (specify): _______________________
___ Not specified

Procedure
___ Endoscopic resection
___ Esophagectomy
___ Esophagogastrectomy
___ Other (specify): _______________________
___ Not specified

Tumor Site (select all that apply) (Note B)
___ Cervical (proximal) esophagus
___ Midesophagus
   + ___ Upper thoracic esophagus
   + ___ Mid thoracic esophagus
___ Distal esophagus (lower thoracic esophagus)
___ Esophagogastrectomy (EGJ)
___ Proximal stomach and esophagogastrectomy
___ Other (specify): _______________________
___ Not specified

Relationship of Tumor to Esophagogastroduodenal Junction (Note B)
___ Tumor is entirely located within the tubular esophagus and does not involve the esophagogastroduodenal junction
___ Tumor midpoint lies in the distal esophagus and tumor involves the esophagogastroduodenal junction
___ Tumor midpoint is located at the esophagogastroduodenal junction
___ Tumor midpoint lies in the proximal stomach or cardia and tumor involves the esophagogastroduodenal junction
___ Not specified
___ Cannot be assessed

Distance of tumor center from esophagogastroduodenal junction (specify, if applicable): ___ cm

Tumor Size
Greatest dimension: ___ cm
+ Additional dimensions: ___ x ___ cm
___ Cannot be determined (explain): _______________________

+ 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 (select all that apply) (Note C)
- [ ] Adenocarcinoma
- [ ] Squamous cell carcinoma
- [ ] Adenosquamous carcinoma
- [ ] High-grade neuroendocrine carcinoma
  - [ ] Large cell neuroendocrine carcinoma
  - [ ] Small cell neuroendocrine carcinoma
- [ ] Undifferentiated carcinoma
- [ ] Other (specify): __________________________
- [ ] Carcinoma, type cannot be determined

### Histologic Grade (Note D)
- [ ] Not applicable
- [ ] GX: Cannot be assessed
- [ ] G1: Well differentiated
- [ ] G2: Moderately differentiated
- [ ] G3: Poorly differentiated
- [ ] G4: Undifferentiated

### Microscopic Tumor Extension (Note E)
- [ ] Cannot be assessed
- [ ] No evidence of primary tumor
- [ ] High-grade dysplasia (carcinoma in situ)
- [ ] Tumor invades lamina propria
- [ ] Tumor invades muscularis mucosae
- [ ] Tumor invades submucosa
- [ ] Tumor invades muscularis propria
- [ ] Tumor invades through the muscularis propria into the periesophageal soft tissue (adventitia)
- [ ] Tumor directly invades adjacent structures (specify): __________________________

### Margins (select all that apply) (Note F)
If all margins uninvolved by invasive carcinoma:
Distance of invasive carcinoma from closest margin: ___ mm or ___ cm
  Specify margin: __________________________

#### Proximal Margin
- [ ] Cannot be assessed
- [ ] Uninvolved by invasive carcinoma
- [ ] Uninvolved by dysplasia
- [ ] Involved by dysplasia
  - [ ] Squamous dysplasia
    - [ ] Low grade
    - [ ] High grade
  - [ ] Intestinal metaplasia (Barrett’s esophagus) with dysplasia
    - [ ] Low grade
    - [ ] High grade
- [ ] Involved by intestinal metaplasia (Barrett’s esophagus) without dysplasia

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+ 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.
**Distal Margin**
- Cannot be assessed
- Uninvolved by invasive carcinoma
- Involved by invasive carcinoma
- Uninvolved by dysplasia
- Involved by dysplasia
  - Squamous dysplasia
    - Low grade
    - High grade
  - Intestinal metaplasia (Barrett's esophagus) with dysplasia
    - Low grade
    - High grade
- Involved by intestinal metaplasia (Barrett's esophagus) without dysplasia

**Circumferential (Adventitial) Margin (esophagectomy or esophagogastrectomy specimens) or Deep Margin (endoscopic resection specimens)**
- Cannot be assessed
- Uninvolved by invasive carcinoma
- Involved by invasive carcinoma

**Mucosal Margin (endoscopic resection specimens)**
- Cannot be assessed
- Uninvolved by invasive carcinoma
- Involved by invasive carcinoma
  + Distance of invasive carcinoma from closest mucosal margin: ___ mm or ___ cm
  - Involved by invasive carcinoma
  - Uninvolved by dysplasia
  - Involved by dysplasia
    - Squamous dysplasia
      - Low grade
      - High grade
    - Intestinal metaplasia (Barrett's esophagus) with dysplasia
      - Low grade
      - High grade
  - Involved by intestinal metaplasia (Barrett's esophagus) without dysplasia

**Other Margin(s) (required only if applicable)**
Specify margin(s): ______________________
- Cannot be assessed
- Uninvolved by invasive carcinoma
- Involved by invasive carcinoma

**Treatment Effect (applicable to carcinomas treated with neoadjuvant therapy) (select all that apply)**
(Note G)
- No prior treatment
- Present
  + No viable cancer cells (complete response, score 0)
  + Single cells or rare small groups of cancer cells (near complete response, score 1)
  + Residual cancer with evident tumor regression, but more than single cells or rare small groups of cancer cells (partial response, score 2)
- Extensive residual cancer with no evident tumor regression (poor or no response, score 3)
- Treatment history not known
Lymph-Vascular Invasion
___ Not identified
___ Present
___ Cannot be determined

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

Pathologic Staging (pTNM) (Note H)

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

Primary Tumor (pT)
___ pTX: Cannot be assessed
___ pT0: No evidence of primary tumor
___ pTis: High-grade dysplasia
___ pT1: Tumor invades lamina propria, muscularis mucosae, or submucosa
___ pT1a: Tumor invades lamina propria or muscularis mucosae
___ pT1b: Tumor invades submucosa
___ pT2: Tumor invades muscularis propria
___ pT3: Tumor invades adventitia
___ pT4: Tumor invades adjacent structures (specify): ________________________
___ pT4a: Resectable tumor invading pleura, pericardium, or diaphragm
___ pT4b: Unresectable tumor invading other adjacent structures, such as aorta, vertebral body, trachea, etc

Regional Lymph Nodes (pN) (Note I)
___ pNX: Cannot be assessed
___ pN0: No regional lymph node metastasis
___ pN1: Regional lymph node metastasis involving 1 to 2 nodes
___ pN2: 3 to 6 nodes involved
___ pN3: 7 or more nodes involved
___ No nodes submitted or found

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) (required only if confirmed pathologically in this case)
___ pM1: Distant metastasis
   Specify site(s), if known: ________________________

+ 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.
Additional Pathologic Findings (select all that apply) (Note J)
___ None identified
___ Intestinal metaplasia (Barrett’s esophagus)
___ Dysplasia
   ___ Low grade
   ___ High grade
+ ___ Esophagitis (type): ___________________________
+ ___ Gastritis (type): ________________________________
+ ___ Other (specify): ________________________________

+ Ancillary Studies
   Note: For HER2 reporting, the CAP Gastric HER2 template should be used. Pending biomarker studies should be listed in the Comments section of this report.
   + Specify: ___________________________________

+ Clinical History (select all that apply) (Note J)
+ ___ Barrett’s esophagus
+ ___ Other (specify): ________________________________
+ ___ Not known

+ Comment(s)
Explanatory Notes

A. Application
This protocol applies to all carcinomas arising in the esophagus and to carcinomas involving the esophagogastric junction (EGJ), including tumors that cross the EGJ but are predominantly located in the proximal stomach. Lymphomas, well-differentiated neuroendocrine tumors (carcinoid tumors), and sarcomas are not included (separate TNM staging systems’ and CAP protocols apply).

B. Location
The location of the tumor in the esophagus (cervical, upper thoracic, midthoracic, lower thoracic, abdominal) and with respect to the macroscopic EGJ (defined as where the tubular esophagus meets the stomach, as measured from the top of the gastric folds) should be noted whenever possible (Figure 1). The macroscopic EGJ often does not correspond to the junction of esophageal squamous mucosa and columnar mucosa because of the common finding in esophageal resection specimens of glandular mucosa involving the distal esophagus. Because anatomic divisions of the esophagus are defined by anatomic boundaries and relationships to other structures, it may not be possible for the pathologist to determine exact tumor location from the resection specimen.

For tumors involving the esophagogastric junction, specific observations should be recorded in an attempt to establish the exact site of origin of the tumor. The EGJ is defined as the junction of the tubular esophagus and the stomach, irrespective of the type of epithelial lining of the esophagus. The pathologist should record the maximum longitudinal dimension of the tumor mass, the distance of the tumor midpoint from the EGJ, and the relative proportions of the tumor mass located in the esophagus and in the stomach.
Tumors involving the EGJ are classified for purposes of staging as esophageal carcinomas. Although the nature of these tumors (gastric versus esophageal) has been controversial (reviewed by Carneiro and Chaves), recent data support their classification as esophageal carcinomas. The World Health Organization (WHO) defines esophageal tumors as those located entirely above the EGJ and proximal gastric tumors as those located entirely below the EGJ. Tumors crossing the EGJ are classified as EGJ tumors. An alternative system proposed by Siewart and colleagues divides adenocarcinomas involving the EGJ into 3 categories, based upon location of the midpoint of the tumor:

Type I: Adenocarcinoma of the distal esophagus, with or without infiltration of the EGJ from above
Type II: True carcinoma of the gastric cardia, arising from the cardiac epithelium or short segments with intestinal metaplasia at the EGJ
Type III: Subcardial gastric carcinoma, which infiltrates the EGJ and distal esophagus from below

Application of the Siewart system is complicated by lack of consensus as to the definition and nature of the gastric cardia, with some investigators regarding it as a normal anatomic finding, and others as a metaplastic response to injury from esophagogastric reflux.

WHO Classification of Carcinoma of the Esophagus
- Squamous cell carcinoma
- Verrucous (squamous) carcinoma
- Spindle cell (squamous) carcinoma
- Adenocarcinoma
- Adenosquamous carcinoma
- Mucoepidermoid carcinoma
- Adenoid cystic carcinoma
- High-grade neuroendocrine carcinoma
  - Large cell neuroendocrine carcinoma
  - Small cell neuroendocrine carcinoma
- Undifferentiated carcinoma
- Others

These types are not generally graded.

The term carcinoma, NOS (not otherwise specified) is not part of the WHO classification.

D. Histologic Grade
The histologic grades for esophageal squamous cell carcinomas are as follows:

Grade X  Grade cannot be assessed
Grade 1   Well differentiated
Grade 2   Moderately differentiated
Grade 3  Poorly differentiated

If there are variations in the differentiation within the tumor, the highest (least favorable) grade is recorded. In general, mucoepidermoid carcinoma and adenoid cystic carcinoma of the esophagus are not amenable to grading.

For adenocarcinomas, a suggested grading system based on the proportion of the tumor that is composed of glands is as follows:

- Grade X  Grade cannot be assessed
- Grade 1  Well differentiated (greater than 95% of tumor composed of glands)
- Grade 2  Moderately differentiated (50% to 95% of tumor composed of glands)
- Grade 3  Poorly differentiated (49% or less of tumor composed of glands)

Undifferentiated tumors cannot be categorized as squamous cell carcinoma or adenocarcinoma (or other) type. They are classified as "undifferentiated carcinomas" in the WHO classification of tumor types (see above) and may be assigned grade 4. Small cell carcinomas are not typically graded but are high-grade tumors and would correspond to grade 4.

The revised TNM staging system for esophageal carcinomas incorporates tumor grade and histologic type in the stage groupings (see Note H). For purposes of staging, grade 4 carcinomas (undifferentiated carcinomas) are staged as grade 3 squamous cell carcinomas. Grade X tumors are grouped as grade 1 carcinomas.

E. Tumor Extension
For purposes of data reporting, Barrett's esophagus with high-grade dysplasia in an esophageal resection specimen is reported as "carcinoma in situ." The term carcinoma in situ is not widely applied to glandular neoplastic lesions in the gastrointestinal tract but is retained for tumor registry reporting purposes as specified by law in many states. Invasion of the lamina propria may be difficult to assess for glandular neoplasms in the esophagus. The muscularis mucosae (Figure 2) is commonly duplicated and thickened in Barrett's esophagus; invasion of this layer should not be misinterpreted as invasion of the muscularis propria. It should be noted that the muscularis mucosae varies in organization from relatively sparse bundles of smooth muscle in the cervical esophagus to a thickened reticulated network in the distal esophagus.

![Microscopic anatomy of the esophagus. From Edge et al. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Illinois. The original source for this material is the AJCC Cancer Staging Manual, 7th edition (2009) published by Springer Science and Business Media LLC, www.springerlink.com.](image)

**Figure 2.** Microscopic anatomy of the esophagus. From Edge et al. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Illinois. The original source for this material is the AJCC Cancer Staging Manual, 7th edition (2009) published by Springer Science and Business Media LLC, www.springerlink.com.

F. Margins
Margins include the proximal, distal, and radial margins. The radial margin represents the adventitial soft tissue margin closest to the deepest penetration of tumor. Sections to evaluate the proximal and distal resections margins can be obtained in 2 orientations: (1) en face sections parallel to the margin or (2) longitudinal sections.
perpendicular to the margin. Depending on the closeness of the tumor to the margin, select the orientation(s) that will most clearly demonstrate the status of the margin. The distance from the tumor edge to the closest resection margin(s) should be measured. Proximal and distal resection margins should be evaluated for Barrett’s esophagus and for squamous and glandular dysplasia. It may be helpful to mark the margin(s) closest to the tumor with ink. Margins marked by ink should be so designated in the macroscopic description.

G. Treatment Effect
Response of tumor to previous chemotherapy or radiation therapy should be reported. Several systems for tumor response have been advocated, and a modified Ryan scheme is suggested, which has been shown to provide good interobserver reproducibility provide prognostic significance in rectal cancer. 

<table>
<thead>
<tr>
<th>Modified Ryan Scheme for Tumor Regression Score</th>
<th>Tumor Regression Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>No viable cancer cells (complete response)</td>
<td>0</td>
</tr>
<tr>
<td>Single cells or rare small groups of cancer cells (near complete response)</td>
<td>1</td>
</tr>
<tr>
<td>Residual cancer with evident tumor regression, but more than single cells or rare small groups of cancer cells (partial response)</td>
<td>2</td>
</tr>
<tr>
<td>Extensive residual cancer with no evident tumor regression (poor or no response)</td>
<td>3</td>
</tr>
</tbody>
</table>

Sizable pools of acellular mucin may be present after chemoradiation but should not be interpreted as representing residual tumor.

This protocol does not preclude the use of other systems for assessment of tumor response. 

H. TNM and Anatomic Stage/Prognostic Groupings
The TNM staging system for esophageal carcinoma of the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC) is recommended (Figure 3).

According to 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 infeasible) 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 after 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 before 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**
A mediastinal lymphadenectomy specimen will ordinarily include 7 or more regional lymph nodes.

### Stage Groupings: Squamous Cell Carcinoma

<table>
<thead>
<tr>
<th>Stage</th>
<th>T</th>
<th>N</th>
<th>M</th>
<th>G</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stage 0</td>
<td>Tis</td>
<td>N0</td>
<td>M0*</td>
<td>1</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IA</td>
<td>T1</td>
<td>N0</td>
<td>M0</td>
<td>1</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IB</td>
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<td>N0</td>
<td>M0</td>
<td>2 or 3</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>T2 or T3</td>
<td>N0</td>
<td>M0</td>
<td>1</td>
<td>Lower</td>
</tr>
<tr>
<td>Stage IIA</td>
<td>T2 or T3</td>
<td>N0</td>
<td>M0</td>
<td>1</td>
<td>Upper, middle</td>
</tr>
<tr>
<td></td>
<td>T2 or T3</td>
<td>N0</td>
<td>M0</td>
<td>2 or 3</td>
<td>Lower</td>
</tr>
<tr>
<td>Stage IIB</td>
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<td>N0</td>
<td>M0</td>
<td>2 or 3</td>
<td>Upper, middle</td>
</tr>
<tr>
<td></td>
<td>T1 or T2</td>
<td>N1</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IIIA</td>
<td>T1 or T2</td>
<td>N2</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>T3</td>
<td>N1</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>T4a</td>
<td>N0</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IIIB</td>
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<td>N2</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IIIC</td>
<td>T4a</td>
<td>N1 or N2</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
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<td></td>
<td>T4b</td>
<td>Any</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>Any</td>
<td>N3</td>
<td>M0</td>
<td>Any</td>
<td>Any</td>
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<td>Stage IV</td>
<td>Any T</td>
<td>Any N</td>
<td>M1</td>
<td>Any</td>
<td>Any</td>
</tr>
</tbody>
</table>

* M0 is defined as no distant metastasis.

### Stage Grouping: Adenocarcinoma

<table>
<thead>
<tr>
<th>Stage</th>
<th>T</th>
<th>N</th>
<th>M</th>
<th>G</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stage 0</td>
<td>Tis (HGD)</td>
<td>N0</td>
<td>M0</td>
<td>1</td>
</tr>
<tr>
<td>Stage IA</td>
<td>T1</td>
<td>N0</td>
<td>M0</td>
<td>1 or 2</td>
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<tr>
<td>Stage IB</td>
<td>T1</td>
<td>N0</td>
<td>M0</td>
<td>3</td>
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<td></td>
<td>T2</td>
<td>N0</td>
<td>M0</td>
<td>1 to 2</td>
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<tr>
<td>Stage IIA</td>
<td>T2</td>
<td>N0</td>
<td>M0</td>
<td>3</td>
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<td>Stage IIB</td>
<td>T3</td>
<td>N0</td>
<td>M0</td>
<td>Any</td>
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<tr>
<td></td>
<td>T1 or T2</td>
<td>N1</td>
<td>M0</td>
<td>Any</td>
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<tr>
<td>Stage IIIA</td>
<td>T1 or T2</td>
<td>N2</td>
<td>M0</td>
<td>Any</td>
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<td>Stage IIIC</td>
<td>T4a</td>
<td>N1 or N2</td>
<td>M0</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>T4b</td>
<td>Any</td>
<td>M0</td>
<td>Any</td>
</tr>
<tr>
<td></td>
<td>Any</td>
<td>N3</td>
<td>M0</td>
<td>Any</td>
</tr>
<tr>
<td>Stage IV</td>
<td>Any T</td>
<td>Any N</td>
<td>M1</td>
<td>Any</td>
</tr>
</tbody>
</table>

* HGD, high-grade dysplasia.
Additional Descriptors

Lymph-Vascular Invasion
Lymph-vascular invasion (LVI) indicates whether microscopic lymph-vascular invasion is identified in the pathology report. LVI includes lymphatic invasion, vascular invasion, or lymph-vascular invasion. By AJCC/UICC convention, LVI does not affect the T category indicating local extent of tumor unless specifically included in the definition of a T category.

I. Regional Lymph Nodes
Regional lymph nodes (Figure 4) extend from periesophageal cervical nodes for the cervical esophagus to celiac lymph nodes for the distal esophagus. Number of involved lymph nodes has consistently emerged as a prognostic indicator on multivariate analysis. Extranodal extension may identify a subset of node-positive patients with a particularly poor prognosis.

Figure 4. Regional lymph nodes of the esophagus. From Edge et al. Used with permission of the American Joint Committee on Cancer (AJCC), Chicago, Illinois. The original source for this material is the AJCC Cancer Staging Manual, 7th edition (2009) published by Springer Science and Business Media LLC, www.springerlink.com.

J. Additional Findings
Most esophageal adenocarcinomas develop in the setting of Barrett’s esophagus, which is defined as alteration of the mucosal lining of the esophagus from the normal squamous epithelium to metaplastic columnar epithelium in response to esophagogastric reflux. Although in some cases the columnar epithelium may resemble gastric oxyntic or cardiac mucosa, only the specialized columnar epithelium with goblet cells is considered to carry significant risk of cancer and is designated as Barrett’s esophagus for diagnostic purposes.

References


