

Protocol for the Examination of Tumor Resection Specimens for Which a Site-Specific Protocol is Not Available

Version: 1.1.0.0

Protocol Posting Date: June 2021

The use of this protocol is recommended for clinical care purposes but is not required for accreditation purposes.

This protocol may be used for the following procedures AND tumor types:

Procedure	Description
Resection	
Tumor Type	Description
Solid tumor	May be used for any malignancy only when an appropriate organ-specific resection protocol is not available

The following should NOT be reported using this protocol:

Procedure	
Biopsy (consider Generic Biopsy protocol)	
Tumor	
Any tumor for which an appropriate organ-specific protocol is available	

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With guidance from the CAP Cancer and CAP Pathology Electronic Reporting Committees. * Denotes primary author.

Accreditation Requirements

The use of this case summary is recommended for clinical care purposes but is not required for accreditation purposes. The core and conditional data elements are routinely reported. Non-core data elements are indicated with a plus sign (+) to allow for reporting information that may be of clinical value.

Summary of Changes

v 1.1.0.0

- General Reformatting
- Revised Margins Section
- Revised Lymph Nodes Section

Reporting Template

Protocol Posting Date: June 2021 Select a single response unless otherwise indicated.

CASE SUMMARY: (GENERAL TUMOR RESECTION: Reporting Template)

May be used for any malignancy for which an appropriate organ-specific resection protocol is not available.

SPECIMEN
+Procedure:
+Specimen Laterality Right Left Not specified Not applicable
+Lymph Node Sampling Performed +Lymph Nodes Sampled: Not performed Not known
TUMOR
+Tumor Site(s):
+Histologic Type:
+Histologic Grade:
+Tumor Size Greatest dimension in Centimeters (cm): cm +Additional Dimension in Centimeters (cm): x cm Cannot be determined (explain):
+Tumor Extent (specify structures or organs involved by tumor):
+Mitotic Rate:
+Tumor Necrosis Not identified Present +Percentage of Tumor Necrosis Specify percentage:% Other (specify):% Cannot be determined Cannot be determined:

thumphoweeouler Invesion	
+Lymphovascular Invasion Not identified	
Present	
Cannot be determined:	
+Treatment Effect	
No known preoperative therapy	
Not identified	
Present	
+Preoperative Therapy Given (spec	cifv):
Cannot be determined:	
• • • • • • • • • • • • • • • • •	
+Tumor Comment:	_
MARGINS	
+Margin Status	
All margins negative for tumor	
+Distance from Tumor to Closest N	largin
Specify in Millimeters (mm)	
Exact distance:	
Greater than:	
At least: mi	
Less than:	mm
Less than 1 mm	
Other (specify):	
Cannot be determined (explain):	
+Closest Margin(s) to Tumor	
Specify closest margin(s):	
Cannot be determined (explain):	
Tumor present at margin	
+Margin(s) Involved by Tumor	
Specify involved margin(s):	
Cannot be determined (explain):	
Other (specify):	
Cannot be determined (explain):	
Not applicable	
+Margin Comment:	_
REGIONAL LYMPH NODES	
+Regional Lymph Node Status	
Not applicable (no regional lymph needed)	odes submitted or found)
Regional lymph nodes present	
All regional lymph nodes negative	
Tumor present in regional lymph	
+Number of Lymph Nodes with 1	
Exact number (specify):	
At least (specify):	

____Other (specify): ______ Cannot be determined (explain): ______ +Extranodal Extension _____Not identified _____Present _____Cannot be determined: ______ ___Other (specify): ______ Cannot be determined (explain): ______ Cannot be determined (explain): ______ +Number of Lymph Nodes Examined _____Exact number (specify): ______ ___At least (specify): ______ ____Other (specify): _______ Cannot be determined (explain): _______ Hegional Lymph Node Comment: ______

+Stage or Classification System (specify): _____

ADDITIONAL FINDINGS

+Additional Findings (specify): _____

SPECIAL STUDIES

List pending biomarker studies in the Comments section of this report. Biomarkers Tested (repeat as needed)

+Biomarker Tested: _____ +Results: _____

+Testing Method: _____

COMMENTS

Comment(s): _____