

# A guide to Colorectal Cancer Histopathology Reporting

Includes the  International Collaboration on Cancer reporting dataset denoted by \*



Clinical details		Microscopic findings	
<b>S1.02</b>	<b>Clinical info. on request form</b> (complete as narrative or use the structured format below)	Text	
	*Known polyposis syndrome	See p2	
	*Lynch syndrome	Text	
	*Chronic inflammatory bowel disease	Ulcerative colitis Crohn disease	
	*Previous polyp(s)	Text	
	*Previous colorectal cancer	Text	
	*Other	Text	
	*Neoadjuvant therapy	Info not provided Not administered Administered, <i>describe</i>	
G1.01	Copy to doctor	Text	
<b>S1.03</b>	<b>Pathology accession number</b>	Text	
<b>S1.04</b>	<b>Principal clinician</b>	Text	
G1.02	Other clinical information	Text	
Macroscopic findings			
<b>S2.01</b>	<b>Specimen labelled as</b>	Text	
<b>S2.02</b>	<b>Clinical information</b>		
<b>S2.03</b>	*Operative procedure	See p2	
<b>S2.04</b>	<b>Specimen length</b>	__ mm	
<b>S2.05</b>	*Tumour site	See p2	
<b>S2.06</b>	*Tumour dimensions <small>*Repeat tumour identification and maximum dimension for each tumour identified</small>	Can't be assess'd, <i>specify</i> Tumour identification __mm max dimension (largest tumour)	
	*Additional dimensions (largest tumour)	__mm x __mm	
<b>S2.07</b>	<b>Distance of tumour to the nearer proximal or distal 'cut end'</b>	Can't be assess'd __ mm	
<b>S2.08</b>	<b>Distance of tumour to the nonperitonealised circumferential margin</b>	Can't be assess'd __mm	
<b>S2.09</b>	*Tumour perforation (defined as a macroscopically visible full thickness defect in the wall)	See p2	
<b>S2.10</b>	*Relation of tumour to anterior peritoneal reflection (rectal cancer specimens only)	See p2	
<b>S2.11</b>	*Plane of mesorectal excision (rectal cancer specimens only)	See p2	
G2.01	*Plane of sphincter excision (abdominoperineal excision specimens only)	See p2	
G2.02	*Plane of mesocolic excision (colon cancer specimens only)	See p2	
G2.03	Peritoneum	See p2	
G2.04	Other macroscopic comment	Text	
<b>S3.01</b>	*Histological tumour type	See p2	
<b>S3.02</b>	*Histologic tumour grade	See p2	
<b>S3.03</b>	*Extent of invasion	See p2	
G3.01	*Measurement of invasion beyond muscularis propria (for pT3 tumours) Dist. beyond muscularis propria	Can't be assess'd, OR __ mm	
G3.02	Inflammatory cell infiltrate	Text	
<b>S3.04</b>	*Lymphatic and venous invasion	See p3	
<b>S3.05</b>	*Perineural invasion	Not identified Present	
<b>S3.06</b>	*Lymph node (LN) status  No. of LNs examined  No. of positive LNs	No nodes submitted or found OR __ Not involved Involved __	
G3.03	Apical node involvement	Not applicable Absent Present	
G3.04	Ratio of involved/total LNs	Involved LNs _ / total LNs _	
<b>S3.07</b>	*Tumour deposits  <i>Number of tumour deposits</i>	See p3 __	
G3.05	*Tumour budding	See p3	
<b>S3.08</b>	*Response to neoadjuvant therapy	See p3	
<b>S3.09</b>	*Margin status  *Longitudinal margin status <i>Distance to closer margin</i> <i>Proximal or distal margin</i>	See p3 __mm __mm	
	*Circumferential margin status <i>Not involved - Dist. to nearest 1 mm or ≥10 mm</i> <i>Involved (≤1 mm) - specify 0 mm or dist. to nearest 0.1 mm</i>	See p3 __mm OR ≥10 mm __mm OR By primary tumour By other, <i>specify</i>	
<b>S3.10</b>	*Distant metastases	Not identified Present, <i>specify sites(s)</i>	
G3.06	*Coexistent pathology	None identified OR Polyp(s), <i>specify</i> Synchronous carcinoma(s), <i>specify</i> Other, <i>specify</i>	
<b>S3.11</b>	Microscopic residual tumour status (completeness of resection)	Text	
G3.07	Additional microscopic comment	Text	

Ancillary test findings		
<a href="#">G4.01</a>	<b>*Ancillary studies</b>	See p3
<b>MMR IHC</b>		
<a href="#">G4.02</a>	<b>*BRAF (V600E) mutation testing</b>	See p3
	<b>*MLH1 promoter methylation testing</b>	
<a href="#">G4.03</a>	<b>*MMR status by microsatellite instability testing</b>	See p3
<a href="#">G4.04</a>	<b>RAS gene mutation testing</b> <i>(KRAS exons 2, 3 or 4, NRS exons 2, 3 or 4 or RAS mutation)</i>	See p3
	Comments	Text
	<i>Laboratory performing test and report number</i>	Text
<a href="#">G4.05</a>	<b>*Neuroendocrine neoplasm markers</b>	Text
	Ki-67 (labelling index)	___ %
	Other	Text
Synthesis and overview		
<a href="#">S5.01</a>	<b>*PATHOLOGICAL STAGING</b>	See p3
<a href="#">S5.02</a>	<b>Residual tumour status</b>	See p4
<a href="#">S5.03</a>	<b>Year and edition of staging system</b>	Text
<a href="#">G5.01</a>	<b>Diagnostic summary</b> Include: Specimen label; Tumour site; Histological tumour type; Margin status; Tumour stage.	Text
<a href="#">S5.04</a>	<b>Overarching comment</b>	Text
<a href="#">G5.02</a>	<b>Edition/version of RCPA protocol</b>	Text

## S1.02 Clinical information

Text  
OR  
Information not provided

### Known polyposis syndrome

*(Select all that apply)*

- Familial adenomatous polyposis (FAP)
- *MUTYH*-associated polyposis (MAP)
- Serrated polyposis
- Other, *specify*

## S2.03 Operative procedure

Not specified  
OR

Select all that apply:

- Total colectomy
- Proctocolectomy
- Right hemicolectomy
- Extended right hemicolectomy
- Transverse colectomy
- Left hemicolectomy
- Anterior resection (*specify if possible*)
  - o High
  - o Low/ultralow
- Abdominoperineal resection
- Other, *specify*

## S2.06 Tumour site\*

If multiple tumours are present, separate protocols should be used to record this and all following elements for each tumour.

- Not specified
- Caecum
- Ascending colon
- Hepatic flexure
- Transverse colon
- Splenic flexure
- Descending colon
- Sigmoid colon
- Rectosigmoid\*
- Rectum
- Other, *specify*

*\*Note: Reserved for cases in which an accurate determination between rectum and sigmoid cannot be made by pathological assessment and clinical information regarding site is not available.*

## S2.09 Tumour perforation

(defined as a macroscopically visible full thickness defect in the wall)

- Not identified
- Present
  - o Through tumour (tumour perforation)
  - o Not involving tumour

## S2.10 Relation of tumour to anterior peritoneal reflection

(rectal cancer specimens only)

- Not applicable
- Entirely above
- Entirely below
- Astride

## S2.11 Plane of mesorectal excision

(rectal cancer specimens only)

- Not applicable
- Mesorectal fascia (complete)
- Intramesorectal (near complete)
- Muscularis propria (incomplete)

## G2.01 Plane of sphincter excision

(abdominoperineal excision specimens only)

- Extralevator plane
- Sphincter plane
- Intrasphincter plane
- 

## G2.02 Plane of mesocolic excision

(colon cancer specimens only)

- Mesocolic plane
- Intramesocolic plane
- Muscularis propria plane

## G2.03 Peritoneum

- Tumour invades to the peritoneal surface
- Tumour has formed nodule(s) discrete from the tumour mass along the serosal surface

## S3.01 Histological tumour type

- Adenocarcinoma, not otherwise specified (NOS)
- Mucinous adenocarcinoma
- Signet-ring cell adenocarcinoma
- Medullary carcinomar
- Serrated adenocarcinoma
- Micropapillary adenocarcinoma
- Adenoma-like adenocarcinoma
- Other, *specify*

## S3.02 Histological tumour grade

- Not applicable
- Low grade  $\geq 50\%$  (formerly well to moderately differentiated)
- High grade  $< 50\%$  (formerly poorly differentiated and undifferentiated)

## S3.03 Extent of invasion

Cannot be assessed

No evidence of primary tumour

OR

Select all that apply:

- High grade dysplasia/non-invasive neoplasia
- Invasion into submucosa
- Invasion into muscularis propria
- Invasion into subserosa or into periocolic or perirectal tissues
- Invasion onto the surface of the visceral peritoneum
- Invasion directly into other structures/organs, *specify*

### S3.04 Lymphatic and venous invasion

- Not identified
- Present
  - o Small vessel (lymphatic, capillary or venular)
  - o Large vessel (venous)
    - Intramural
    - Extramural

### S3.07 Tumour deposits

- Not identified
- Present, *specify type*
  - o Vascular
  - o Other

### G3.05 Tumour budding

Cannot be assessed

OR

Number of tumour buds \_\_\_

\*Note: After scanning 10 fields on a 20x objective lens, the hotspot field normalised to represent a field of 0.785 mm<sup>2</sup>

Tumour budding score

- Bd1 - low budding (0-4 buds)
- Bd2 - intermediate budding (5-9 buds)
- Bd3 - high budding ( $\geq 10$  buds)

### S3.08 Response to neoadjuvant therapy

- No neoadjuvant treatment
- Complete response - No viable cancer cells (score 0)
- Near complete response - Single cells or rare small groups of cancer cells (score 1)
- Partial response - Residual cancer with evident tumour regression, but more than single cells or rare small groups of cancer cells (score 2)
- Poor or no response - Extensive residual cancer with no evident tumour regression (score 3)
- Cannot be assessed, *specify*

### S3.09 Margin status

Longitudinal margin status

- Cannot be assessed
- Not involved
- Involved

Note: Includes assessment of any separately submitted anastomotic ring(s).

Circumferential margin status

- Cannot be assessed
- Not involved
- Involved

### G4.01 Ancillary studies

Mismatch repair (MMR) immunohistochemistry

Not tested

OR

Not interpretable

OR

- MMR proficient
- MMR deficient
  - o *MLH1/PMS2* loss
  - o *MSH2/MSH6* loss
  - o *MSH6* loss
  - o *PMS2* loss
  - o Other, *specify*

### G4.02 BRAF (V600E) mutation testing

Not tested

OR

- Test failed
- Mutated
- Wild type

MLH1 promoter methylation testing

Not tested

OR

- Test failed
- Methylated
- Not methylated
- Inconclusive

### G4.03 MMR status by microsatellite instability (MSI) testing

Not tested

OR

- Test failed
- MSI-high
- MSI-low
- MSI-stable

### G4.04 RAS gene mutation testing (*KRAS* exons 2, 3 or 4, *NRAS* exons 2, 3 or 4 or *RAS* mutation)

- Mutated
- Wild type
- Not tested

### S5.01 Stage & stage group##

**Suffixes**

m - multiple primary tumours; y - post therapy; r - recurrent

**Primary Tumour (T)**

- TX Primary tumour cannot be assessed
- T0 No evidence of primary tumour
- Tis Carcinoma in situ, intramucosal carcinoma (involvement of lamina propria with no extension through muscularis mucosae)
- T1 Tumour invades the submucosa (through the muscularis mucosa but not into the muscularis propria)
- T2 Tumour invades the muscularis propria
- T3 Tumour invades through the muscularis propria into pericolorectal tissues
- T4 Tumour invades\* the visceral peritoneum or invades or adheres\*\* to adjacent organ or structure
- T4a Tumour invades\* through the visceral peritoneum (including gross perforation of the bowel through tumour and continuous invasion of tumour through areas of inflammation to the surface of the visceral peritoneum)
- T4b Tumour directly invades\* or adheres\*\* to adjacent organs or structures

\*\*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 tumour 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).

\*\*Tumour that is adherent to other organs or structures, grossly, is classified cT4b. However, if no tumour 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.

### **Regional lymph node (pN)**

- NX Regional lymph nodes cannot be assessed
- N0 No regional lymph node metastasis
- N1 One to three regional lymph nodes are positive (tumour in lymph nodes measuring  $\geq 0.2$  mm), or any number of tumour deposits are present and all identifiable lymph nodes are negative
  - N1a One regional lymph node is positive
  - N1b Two or three regional lymph nodes are positive
  - N1c No regional lymph nodes are positive, but there are tumour deposits in the subserosa mesentery or nonperitonealized pericolic, or perirectal/mesorectal tissues.
- N2 Four or more regional nodes are positive
  - N2a Four to six regional lymph nodes are positive
  - N2b Seven or more regional lymph nodes are positive

### **Distant metastasis (pM)**

- M0 No distant metastases by imaging, etc.; no evidence of tumour in distant sites or organs (This category is not assigned by pathologists.)
- M1 Metastasis to one or more distant sites or organs or peritoneal metastasis is identified
  - M1a Metastasis to one site or organ is identified without peritoneal metastasis
  - M1b Metastasis to two or more sites or organs is identified without peritoneal metastasis
  - M1c Metastasis to the peritoneal surface is identified alone or with other site or organ metastases

## **S5.02 Residual tumour status**

- RX Presence of residual tumour cannot be assessed
- R0 No residual tumour
- R1 Microscopic residual tumour
- R2 Macroscopic residual tumour at the primary cancer site or regional nodal sites. (This designation is not used to indicate metastatic disease identified but not resected at surgical exploration.)

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