

# Intrahepatic Cholangiocarcinoma, Perihilar Cholangiocarcinoma and Hepatocellular Carcinoma Histopathology Reporting Guide



Family/Last name  Date of birth

Given name(s)

Patient identifiers  Date of request  Accession/Laboratory number

Elements in **black text** are **CORE**. Elements in **grey text** are **NON-CORE**.

☐ indicates multi-select values ☐ indicates single select values

SCOPE OF THIS DATASET

## SPECIMEN(S) SUBMITTED (select all that apply)

- ☐ Not specified  
☐ Indeterminate  
☐ Liver  
☐ Total hepatectomy  
☐ Segmental resection, *specify segment(s) or type of segmentectomy*

  


☐ Wedge resection, *specify site/segment*

- ☐ Extrahepatic bile duct  
☐ Gallbladder  
☐ Diaphragm  
☐ Lymph nodes, *specify site(s), distinguishing between portal and extra-portal nodes*

  


☐ Other, *specify*

## SPECIMEN DIMENSIONS

(Indicate greatest measurement for each parameter in an irregularly shaped specimen)

mm x  mm x  mm

Length of extrahepatic bile duct  
(Applicable to perihilar cholangiocarcinoma only)

 mm

## SPECIMEN WEIGHT

 g

## SATELLITOSIS

(Applicable to hepatocellular carcinoma only)

- ☐ Cannot be assessed ☐ Not identified ☐ Present

## MACROSCOPIC TUMOUR RUPTURE

(Applicable to hepatocellular carcinoma and perihilar cholangiocarcinoma only)

- ☐ Fragmented specimen ☐ Ruptured ☐ Intact

## TUMOUR SITE AND NUMBER

- ☐ No macroscopic residual tumour

Tumour ID	Specify	No./site, if possible
<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>

## MAXIMUM TUMOUR DIMENSION

- ☐ Cannot be assessed

Tumour ID	Maximum dimension
<input type="text"/>	<input type="text"/> mm
<input type="text"/>	<input type="text"/> mm
<input type="text"/>	<input type="text"/> mm
<input type="text"/>	<input type="text"/> mm
<input type="text"/>	<input type="text"/> mm

For a large number of tumours include a range  mm to  mm

Linear extent of tumour along the bile duct  
(Applicable to perihilar cholangiocarcinoma only, where possible)

 mm

## HISTOLOGICAL TUMOUR TYPE

(Value list from the World Health Organization Classification of Tumours of the Digestive System (2019))

- ☐ Hepatocellular carcinoma  
☐ Intrahepatic cholangiocarcinoma  
☐ Large duct ☐ Small duct ☐ Other  
☐ Perihilar cholangiocarcinoma  
☐ Combined hepatocellular – cholangiocarcinoma  
☐ Intraductal papillary neoplasm with an associated invasive carcinoma  
☐ Mucinous cystic neoplasm with an associated invasive carcinoma  
☐ Undifferentiated carcinoma  
☐ Carcinoma, type cannot be determined



**HEPATOCELLULAR CARCINOMA SUBTYPE**

- |   |                                       |
|---|---------------------------------------|
| <input type="radio"/> Steatohepatic           | <input type="radio"/> Chromophobe     |
| <input type="radio"/> Clear cell              | <input type="radio"/> Fibrolamellar   |
| <input type="radio"/> Macrotrabecular massive | <input type="radio"/> Neutrophil-rich |
| <input type="radio"/> Scirrhus                | <input type="radio"/> Lymphocyte-rich |
| <input type="radio"/> No special type         |                                       |

**TUMOUR GROWTH PATTERN**
**Hepatocellular carcinoma**

- ☐ Cannot be determined  
☐ Early hepatocellular carcinoma  
☐ Single distinct nodule  
☐ Large dominant nodule with multiple small satellite nodules  
☐ Cirrhotomimetic  
☐ Multiple distinct nodules

**Intrahepatic and perihilar cholangiocarcinoma**

- ☐ Cannot be determined  
☐ Mass-forming  
☐ Intraductal-growth  
☐ Periductal infiltrating  
☐ Mixed mass-forming and periductal infiltrating

**HISTOLOGICAL TUMOUR GRADE**

- ☐ Not applicable  
☐ Cannot be assessed  
☐ Grade 1: Well differentiated  
☐ Grade 2: Moderately differentiated  
☐ Grade 3: Poorly differentiated

**EXTENT OF INVASION**

- ☐ Cannot be assessed  
☐ No evidence of primary tumour  
☐ Macroscopic invasion
 

☐ Tumour confined to liver  
☐ Tumour confined to the extrahepatic bile ducts (carcinoma in situ/high grade dysplasia) (*Applicable to perihilar cholangiocarcinoma only*)  
☐ Tumour involves visceral peritoneum  
☐ Tumour directly invades gallbladder  
☐ Invasion of periductal tissue - either adipose or hepatic tissue (*Applicable to perihilar cholangiocarcinoma only*)  
☐ Tumour directly invades other adjacent organs, *specify*
- ☐ Microscopic invasion
 

☐ Tumour confined to liver  
☐ Tumour confined to the bile duct mucosa histologically (carcinoma in situ/high grade dysplasia) (*Applicable to perihilar cholangiocarcinoma only*)  
☐ Tumour involves visceral peritoneum  
☐ Tumour directly invades gallbladder  
☐ Invasion of periductal tissue - either adipose or hepatic tissue (*Applicable to perihilar cholangiocarcinoma only*)  
☐ Tumour directly invades other adjacent organs, *specify*

**PERINEURAL INVASION**

(Applicable to intrahepatic and perihilar cholangiocarcinoma)

- ☐ Not identified   ☐ Indeterminate   ☐ Present

**VASCULAR INVASION**

- ☐ Not identified  
☐ Indeterminate  
☐ Present macroscopically (large portal or hepatic veins)  
☐ Present microscopically (small portal or hepatic veins or microvessels)

**COEXISTENT PATHOLOGY**
**Other histopathological features** (select all that apply)

- ☐ None identified  
☐ Steatosis  
☐ Steatohepatitis  
☐ Iron overload  
☐ Biliary disease, *specify if known*

- ☐ Chronic hepatitis, *specify type if known*

- ☐ Other, *specify*

**Fibrosis**

- ☐ Not identified   ☐ Indeterminate   ☐ Present

ISHAK stage	/6
OR	
KLEINER stage	/4
OR	
METAVIR stage	/4
OR	
BATTS-LUDWIG stage	/4
OR	
SAF system	/4

**Dysplastic/pre-malignant lesions**

- ☐ None identified

**BILIARY INTRA-EPITHELIAL NEOPLASIA (BiIN)**

- ☐ Absent   ☐ Present
 

☐ High grade BiIN  
☐ Low grade BiIN

**INTRADUCTAL PAPILLARY NEOPLASM OF THE BILE DUCTS (IPNB)**

- ☐ Absent   ☐ Present
 

☐ High grade IPNB  
☐ Low grade IPNB

**LOW GRADE HEPATOCELLULAR DYSPLASTIC NODULE**

- ☐ Absent   ☐ Present

**HIGH GRADE HEPATOCELLULAR DYSPLASTIC NODULE**

- ☐ Absent   ☐ Present



## RESPONSE TO NEOADJUVANT THERAPY

- ☐ No neoadjuvant treatment
- ☐ Complete response – no viable cancer cells
- ☐ Partial response – residual cancer with some tumour regression
- Percentage necrosis  %
- ☐ No response – extensive residual cancer with no evident tumour regression
- ☐ Cannot be assessed, *specify*

## MARGIN STATUS

- ☐ Cannot be assessed
- ☐ Not involved by invasive carcinoma
- Distance of tumour to closest margin  mm
- OR
- ☐ Clearance is  $\geq 10$  mm
- ☐ Involved by invasive carcinoma
- Specify margin(s), if possible
- ☐ Involved by BiIN
- (Applicable to cholangiocarcinoma only)
- Specify margin(s), if possible

## LYMPH NODE STATUS

- ☐ Cannot be assessed
- ☐ No nodes submitted or found
- Number of lymph nodes examined
- ☐ Not involved
- ☐ Involved
- Number of involved lymph nodes
- ☐ Number cannot be determined

## ANCILLARY STUDIES

- ☐ Not performed
- ☐ Performed, *specify*

## PATHOLOGICAL STAGING (UICC TNM 8<sup>th</sup> edition)<sup>a</sup>

### Primary tumour (pT)

#### INTRAHEPATIC CHOLANGIOCARCINOMA<sup>b</sup> (Intrahepatic bile ducts)

- ☐ TX Primary tumour cannot be assessed
- ☐ T0 No evidence of primary tumour
- ☐ Tis Carcinoma in situ (intraductal tumour)
- ☐ T1a Solitary tumour 5 cm or less in greatest dimension without vascular invasion
- ☐ T1b Solitary tumour more than 5 cm in greatest dimension without vascular invasion
- ☐ T2 Solitary tumour with intrahepatic vascular invasion or multiple tumours, with or without vascular invasion
- ☐ T3 Tumour perforating the visceral peritoneum
- ☐ T4 Tumour involving local extrahepatic structures by direct hepatic invasion

## HEPATOCELLULAR CARCINOMA

(Liver excluding intrahepatic and perihilar bile ducts)

- ☐ TX Primary tumour cannot be assessed
- ☐ T0 No evidence of primary tumour
- ☐ T1a Solitary tumour 2 cm or less in greatest dimension with or without vascular invasion
- ☐ T1b Solitary tumour more than 2 cm in greatest dimension without vascular invasion
- ☐ T2 Solitary tumour more than 2 cm dimension with vascular invasion or multiple tumours none more than 5 cm in greatest dimension
- ☐ T3 Multiple tumours any more than 5 cm in greatest dimension
- ☐ T4 Tumour(s) involving a major branch of the portal or hepatic vein or with direct invasion of adjacent organs (including the diaphragm), other than the gallbladder or with perforation of visceral peritoneum

## PERIHILAR CHOLANGIOCARCINOMA

(Perihilar bile ducts)

- ☐ TX Primary tumour cannot be assessed
- ☐ T0 No evidence of primary tumour
- ☐ Tis Carcinoma in situ
- ☐ T1 Tumour confined to the bile duct, with extension up to the muscle layer or fibrous tissue
- ☐ T2a Tumour invades beyond the wall of the bile duct to surrounding adipose tissue
- ☐ T2b Tumour invades adjacent hepatic parenchyma
- ☐ T3 Tumour invades unilateral branches of the portal vein or hepatic artery
- ☐ T4 Tumour invades main portal vein or its branches bilaterally; or the common hepatic artery; or unilateral second-order biliary radicals with contralateral portal vein or hepatic artery involvement

### Regional lymph nodes (pN)

- ☐ No nodes submitted or found

## HEPATOCELLULAR CARCINOMA & INTRAHEPATIC CHOLANGIOCARCINOMA

(Liver including intrahepatic bile ducts and excluding perihilar bile ducts)

- ☐ NX Regional lymph nodes cannot be assessed
- ☐ N0 No regional lymph node metastasis
- ☐ N1 Regional lymph node metastasis

## PERIHILAR CHOLANGIOCARCINOMA

(Perihilar bile ducts)

- ☐ NX Regional lymph nodes cannot be assessed
- ☐ N0 No regional lymph node metastasis
- ☐ N1 Metastases to 1-3 regional lymph nodes
- ☐ N2 Metastases to 4 or more regional lymph nodes

### Distant metastasis (pM)

- ☐ Not applicable
- ☐ M1 Distant metastasis

### TNM Descriptors (only if applicable) (select all that apply)

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

<sup>a</sup> Reproduced with permission. Source: UICC TNM Classification of Malignant Tumours, 8<sup>th</sup> Edition, eds by James D. Brierley, Mary K. Gospodarowicz, Christian Wittekind. 2016, Publisher Wiley-Blackwell.

<sup>b</sup> Combined Hepatocellular-Cholangiocarcinomas are staged as per Intrahepatic Cholangiocarcinoma.