Lung Cancer Histopathology Reporting Guide

International Collaboration on Cancer Reporting (ICCR)

Family/Last name	Date of birth DD – MM – YYYY
Given name(s)	
Patient identifiers	Date of request Accession/Laboratory number
	DD – MM – YYYY
Elements in black text are CORE. Elements in grey text are N	ON-CORE. SCOPE OF THIS DATASET
OPERATIVE PROCEDURE Wedge resection Lobectomy Segmentectomy Bilobectomy Other, specify Pneumonectomy	MAXIMUM TUMOUR DIMENSION mm TUMOUR INVOLVES MAIN BRONCHUS Not applicable Not identified Not assessable Present
SPECIMEN LATERALITY	Not applicable Not identified Not assessable Present
ATTACHED ANATOMICAL STRUCTURES	HISTOLOGICAL TUMOUR TYPE (Value list from the World Health Organisation Classification of Tumours. Pathology and Genetics of Tumours of the Lung, Pleura, Thymus and Heart. (2015)) (select all that apply)
ACCOMPANYING SPECIMENS None submitted Lymph nodes Other, specify	Squamous cell carcinoma Carcinoid Keratinizing Typical Non-keratinizing Atypical Basaloid Large cell neuroendocrine carcinoma Large cell carcinoma Small cell carcinoma
TUMOUR SITE Upper lobe Middle lobe Lower lobe Bronchus, specify site	Classification of Adenocarcinoma Adenocarcinoma Adenocarcinoma in situ (AIS) Non-mucinous Mucinous
SEPARATE TUMOUR NODULES Absent Cannot be assessed Synchronous primaries (CORE elements should be	 Minimally invasive adenocarcinoma (MIA) Non-mucinous Mucinous Invasive adenocarcinoma PREDOMINANT PATTERN Lepidic
Present Site Same lobe Different ipsilateral lobe Contralateral lung	Acinar Papillary Micropapillary Solid Invasive mucinous Colloid Fetal Enteric
MACROSCOPIC APPEARANCE OF PLEURA	OTHER PATTERNS (<i>if present</i>) $TYPE OF PATTERN \implies \%$
	TYPE OF PATTERN ➡ %
	TYPE OF PATTERN ➡ %
ATELECTASIS/OBSTRUCTIVE PNEUMONITIS EXTENDING	Other, <i>specify</i>
Present Absent Not assessable	

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DISTANCE OF TUMOUR TO CLOSEST	mm	SURGICAL MARGIN STAT	US
RESECTION MARGIN		Bronchial margin	
HISTOLOGICAL GRADE Well differentiated U Moderately differentiated Poorly differentiated	Indifferentiated lot applicable	 Involved by invasiv Involved by carcino Only peribronchial s Vascular margin	e carcinoma ONot involved oma in situ only Not applicable soft tissue involved
RESPONSE TO NEOADJUVANT THERA	PY	Involved Not i	involved ONot applicable Oft tissue involved
Not applicable Less than 10% residual viable tu Greater than 10% residual viable Treatment history not known	imour e tumour	Other margin 1 (specify e.	g. parenchymal, chest wall)
DIRECT INVASION OF ADJACENT ST		O Involved ONot i	involved ONot applicable
(select all that apply)		Other margin 2 (specify e.	g. parenchymal, chest wall)
 Trachea Chest wall Diaphragm Oesophagus Heart 	 Not identified Not applicable 	Involved ONOT	involved O Not applicable
Great vessels			
Phrenic nerve		LYMPH NODES STATUS	
		Station(s) examined,	specify
Mediastinal pleura			
Recurrent laryngeal nerve			
LYMPHOVASCULAR INVASION		Not involved Involved by micro	metastacis only
O Present O Not identified			
		▼ Involved	
VISCERAL PLEURAL INVASION		station 1	
Present Not identified			Number of involved
ţ	Cannot be assessed		Total number of lymph nodes from this site
Extent of pleural involveme	nt 📖		Number cannot be determined
O PL1		Involved	
PL2		station 2	
UPL3			Number of involved lymph nodes
PERINEURAL INVASION			Total number of lymph nodes from this site
	\bigcirc		O Number cannot be determined
OTHER NEOPLASTIC PROCESSES (e.g. tumourlets, NEH, AAH, dys	plasia)	Involved station 3	
			Number of involved lymph nodes
			Total number of lymph nodes from this site
NON-NEODI ASTIC LUNG DISEASE			Number cannot be determined

ANCILLAF	RY STUDIES	EML4-ALK result	
Immunoh	istochemical markers	 Rearrangement absent Result indeterminate 	
Po	isitive Abs		
Ne	egative Abs		
Eq	juivocal Abs		
Cor	nclusions:		
		Other, specify	
		Test Result	
Molecular	data 💷		
EG	FR result		
0) Mutation absent O Result indeterminate		
$ $ \bigcirc	Mutation present		
	Describe		
PATHOL	OGICAL STAGING (TNM 8th edition)##	N - Regional lymph nodes	
🗌 m - r	nultiple primary tumours at a single site	N0 No regional lymph node metastasis	
y - c	classification is performed during or following	N1 Metastasis in ipsilateral peribronchial and/ or ipsilateral hilar lymph nodes and intrapulmonary nodes, including	
r	multimodality treatment	involvement by direct extension	
T - P	rimary tumour	N2 Metastasis in ipsilateral mediastinal and/ or subcarinal lymph node(s)	
⊖тх	Primary tumour cannot be assessed, or tumour	N3 Metastasis in contralateral mediastinal, contralateral	
	or bronchial washings but not visualized by imaging	supraclavicular lymph node(s)	
○ ТО	or bronchoscopy. No evidence of primary tumour		
Tis	Carcinoma in situ ^a	M - Distant metastasis	
○ T1	Tumour 3 cm or less in greatest dimension, surrounded by lung or visceral pleura, without	 Not applicable M0 No distant metastasis 	
	bronchoscopic evidence of invasion more proximal	M1 Distant metastasis	
	bronchus) ^b	M1a Separate tumour nodule(s) in a contralateral lobe; tumour with pleural or pericardial nodules or malignant	
	Minimally invasive adenocarcinoma ^c Tumour 1 cm or less in greatest dimension ^b	pleural or pericardial effusion ^e	
T1b	Tumour more than 1 cm but not more than 2 cm in	M1c Multiple extrathoracic metastasis in a single organ	
∩T1c	greatest dimension [®] Tumour more than 2 cm but not more than 3 cm in	organs	
	greatest dimension ^b		
	tumour more than 3 cm but not more than 5 cm; or tumour with any of the following features ^d	 a. Tis includes adenocarcinoma in situ and squamous carcinoma in situ 	
	 Involves main bronchus regardless of distance to the carina, but without involvement of the carina 	b. The uncommon superficial spreading tumour of any size with	
	 Invades visceral pleura 	its invasive component limited to the bronchial wall, which may extend proximal to the main bronchus, is also classified as T1a.	
	 Associated with atelectasis or obstructive pneumonitis that extends to the hilar region either involving part of or the entire lung 	c. Solitary adenocarcinoma (not more than 3 cm in greatest dimension), with a predominantly lepidic pattern and not more	
◯ T2a	Tumour more than 3 cm but not more than 4 cm in	 d. T2 tumours with these features are classified T2a if 4 cm or less, 	
⊖ T2b	greatest dimension. Tumour more than 4 cm but not more than 5 cm in	or if size cannot be determined and T2b if greater than 4 cm but not larger than 5 cm.	
<u>О та</u>	greatest dimension.	e. Most pleural (pericardial) effusions with lung cancer are due	
015	greatest dimension or one that directly invades any	examinations of pleural (pericardial) fluid are negative for	
	of the following: parietal pleura, chest wall (including superior sulcus tumours) phrenic nerve, parietal	tumour, and the fluid is non-bloody and is not an exudate. Where these elements and clinical judgment dictate that the effusion is	
	pericardium; or separate tumour nodule(s) in the same lobe as the primary	not related to the tumour, the effusion should be excluded as a staging descriptor.	
◯ T4	Tumour more than 7 cm or of any size that invades	f. This includes involvement of a single non-regional node.	
	any of the following: diaphragm, mediastinum, heart, great vessels, trachea, recurrent larvngeal nerve.	## Reproduced with permission, Source:Brierley ID, Gospodarowicz MK	
	oesophagus, vertebral body, carina; separate tumour nodule(s) in a different ipsilateral lobe to that of the primary.	and Wittekind C (eds) (2016). UICC TNM Classification of Malignant Tumours, 8th Edition, Wiley-Blackwell.	