

Carcinomas of the Nasal Cavity and Paranasal Sinuses **Histopathology Reporting Guide**



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 NON-CORE. indicates multi-select values indicates single select values		
CLINICAL INFORMATION Information not provided Information provided (select all that apply) Previous therapy	Neck (lymph node) dissection, a specify Other, specify	
Surgery Chemotherapy Radiotherapy Targeted therapy, specify if available	TUMOUR SITE (select all that apply)	
•		
Immunotherapy, specify if available	○ Not specified□ Nasal cavity□ Septum□ Lateral wall	
Other clinical information, specify	☐ Floor ☐ Vestibule ☐ Paranasal sinus(es), maxillary ☐ Paranasal sinus(es), ethmoid	
OPERATIVE PROCEDURE (select all that apply) Not submitted Biopsy (excision, incisional, core needle), specify Resection Open En bloc Endoscopic Piecemeal	Cribriform plate Paranasal sinus(es), frontal Paranasal sinus(es), sphenoid Orbit Cranial cavity Other, specify	
Combined Neck (lymph node) dissection, a specify	TUMOUR LATERALITY (select all that apply)	
	Not specified	
Other, specify	Left Right Midline	
^a If a neck (lymph node) dissection is submitted, then a separate datase is used to record the information.	TUMOUR DIMENSIONS	
SPECIMEN(S) SUBMITTED (select all that apply)	Maximum tumour dimension (largest tumour) (pathology and/or imaging determination)	
Not specified Nasal cavity, specify	Additional dimensions (largest tumour)	
Paranasal sinus(es), specify	mm x mm	
Orbit, specify	BLOCK IDENTIFICATION KEY (List overleaf or separately with an indication of the nature and origin of all tissue blocks)	

HISTOLOGICAL TUMOUR TYPE (select all that apply)	LYMPHOVASCULAR INVASION
(Value list based on the World Health Organization	Not identified
Classification of Head and Neck Tumours (2024))	Present
Keratinising squamous cell carcinoma	Indeterminate, specify reason
Other squamous cell carcinoma subtype, <i>specify type</i>	V
Non-keratinising squamous cell carcinoma	
□ NUT carcinoma	PERINEURAL INVASION
☐ SWI/SNF complex-deficient sinonasal carcinoma	Not identified
☐ Sinonasal lymphoepithelial carcinoma	Present
Sinonasal undifferentiated carcinoma	Indeterminate, specify reason
☐ Teratocarcinosarcoma	V
HPV-related multiphenotypic sinonasal carcinoma	
Adenocarcinoma	
Intestinal-type adenocarcinoma	MARGIN STATUS
 ○ Non-intestinal-type adenocarcinoma □ Salivary gland-type carcinoma, b specify type 	Not involved by invasive carcinoma
Salivary gland-type carcinoma, specify type	Specify closest margin(s), if possible
Neuroendocrine neoplasm	
Small cell neuroendocrine carcinoma	Involved by invasive carcinoma
Large cell neuroendocrine carcinoma	Specify margin(s), if possible
Carcinoma mixed with neuroendocrine carcinoma	
Other, specify	
·	Cannot be assessed, specify
b	
For histological type of salivary gland-type carcinomas, refer to the Carcinomas of the major salivary glands dataset.	
HISTOLOGICAL TUMOUR GRADE	PRECURSOR LESIONS
(Not applicable to all tumours)	O Not applicable
Not applicable	○ Not present
Grade 1, well differentiated, low grade	Present (e.g., sinonasal papilloma (type), surface
Grade 2, moderately differentiated, intermediate grade	▼ dysplasia), <i>specify</i>
Grade 3, poorly differentiated, high grade	
Undifferentiated	
High grade transformation	
Grading system	
used, specify	ANCILLARY STUDIES
Cannot be assessed, specify	Not performed
	Performed
Conding of neuroandeering turneurs is non-core. Her only Crade 1	If performed, specify (select) all that apply
2 and 3 for neuroendocrine tumours; neuroendocrine carcinomas are	Non-keratinising squamous cell carcinoma
considered high grade by definition and are therefore not graded.	Opositive
EXTENT OF INVASION	Pancytokeratin p63
○ Not identified	☐ p40 ☐ CK5/6
Present (select all that apply)	Negative
Clinical observation Histologic	▼ □ CD99
and/or imaging	□ NKX2.2
■ Bone/cartilage invasion	NUT
Cortical bone erosion	○ INI1
Medullary bone involvement	Retained Deficient
Soft tissue infiltration	BRG1
Skull base involvement	Retained Deficient
☐ Invasion of skin	NUT carcinoma
☐ Invasion of orbital tissues	Ositive
Other,	▼ □ NUT immunohistochemistry
▼ specify	NUTM1 gene rearrangement, specify technique
Cannot be assessed, specify	

NCILLARY STUDIES continued	Other ancillary studies, record test(s), methodology
SWI/SNF complex-deficient sinonasal carcinoma	and results
◯ INI1	
Retained Deficient	
BRG1	
Retained Deficient	
Sinonasal undifferentiated carcinoma	Representative blocks for ancillary studies, specify those blocks best representing tumour and/or normal tissue
Positive Pancytokeratin CK7	for further study
Negative	
p40/p63 NKX2.2	
☐ CK5/6 ☐ NUT	
☐ CD99	4
◯ INI1	PATHOLOGICAL STAGING (UICC TNM 8th edition) D
Retained Deficient	TNM Descriptors (only if applicable) (select all that apply)
○ BRG1	m - multiple primary tumours
Retained Deficient	r - recurrent
HPV-related multiphenotypic sinonasal carcinoma	
Positive	Primary tumour (pT) ^e
p16 immunohistochemistry (screening)	TX ^f Primary tumour cannot be assessed
HPV-specific testing, specify technique	Tis Carcinoma in situ
	MAXILLARY SINUS
Neuroendocrine carcinoma	T1 Tumour limited to the mucosa with no erosion or
O Positive	destruction of bone T2 Tumour causing bone erosion or destruction,
CAM5.2/CK-pan Synaptophysin	including extension into the hard palate and/or
Chromogranin INSM1	middle nasal meatus, except extension to posterior
Ki-67 proliferation index %	wall of maxillary sinus and pterygoid plates T3 Tumour invades any of the following: bone of
☐ Ki-67 proliferation index %	posterior wall of maxillary sinus, subcutaneous
Rb	tissues, floor or medial wall of orbit, pterygoid fossa or ethmoid sinuses
Retained Deficient	T4a Tumour invades any of the following: anterior
Keratinising squamous cell carcinoma	orbital contents, skin of cheek, pterygoid plates,
Positive	infratemporal fossa, cribriform plate, sphenoid or frontal sinuses
Pancytokeratin p63 p40 CK5/6	T4b Tumour invades any of the following: orbital apex,
	dura, brain, middle cranial fossa, cranial nerves
Sinonasal lymphoepithelial carcinoma Positive	other than maxillary division of trigeminal nerve (V2), nasopharynx, or clivus
Pancytokeratin	
p16	T1 Tumour restricted to one subsite of nasal cavity or
Teratocarcinosarcoma	ethmoid sinus, with or without bony invasion
Positive	T2 Tumour involves two subsites in a single site or
Nuclear β-catenin	extends to involve an adjacent site within the nasoethmoidal complex, with or without bony
O BRG4 (SMARCA4)	invasion
Retained Deficient	T3 Tumour extends to invade the medial wall or floor of
Intestinal-type sinonasal adenocarcinoma	the orbit, maxillary sinus, palate, or cribriform plate
Positive	 T4a Tumour invades any of the following: anterior orbita contents, skin of nose or cheek, minimal extension
CDX2 SATB2	to anterior cranial fossa, pterygoid plates, sphenoid
☐ CK20 ☐ Villin	or frontal sinuses
CK7	T4b Tumour invades any of the following: orbital apex, dura, brain, middle cranial fossa, cranial nerves other
Non-intestinal-type sinonasal adenocarcinoma	than V2, nasopharynx, or clivus
Positive	d Reproduced with permission. Source: UICC TNM Classification of
CK7 S100 protein	Malignant Tumours, 8th Edition, eds by James D. Brierley, Mary K. Gospodarowicz, Christian Wittekind. 2016, Publisher Wiley
☐ SOX10 ☐ Nuclear β-catenin	(incorporating any errata published up until 12 th July 2024).
☐ DOG1	e Note that the results of neck (lymph node) dissection are derived from a
Negative	separate dataset.
▼ CK20 CDX2	TX should be used only if absolutely necessary.