

Adrenal Gland Tumours Histopathology Proforma



Mandatory questions (i.e. protocol standards) are in bold (e.g. **S1.03**).

Family name

Given name(s)

Date of birth

Patient identifiers

e.g. MRN, IHI or NHI (please indicate which)

Date of request

S1.03 Accession number

Sex

- Male
 Female
 Intersex/indeterminate

Ethnicity

- Unknown
 Aboriginal/Torres Strait Islander
 Other ethnicity:

Requesting doctor - name and contact details

Operative procedure

Functionality of the adrenal gland

Non-functional Functional

Type of presentation:

Any previous adrenal surgery

Site of lesion(s)

Laterality of the lesion

Left Right Both

Any accompanying specimens

Lymph nodes Kidney Other

Relevant clinical information

Pre-operative biopsy

Not performed Performed

Clinical stage

Involvement of adjacent organs

Not involved Involved

Clinical diagnosis or differential diagnosis

New primary cancer or recurrence

New primary Regional (local) recurrence

Distant metastases

S1.04 Principal clinician caring for the patient

S1.05 Surgeon's name & contact details

G1.01 Other relevant details

Macroscopic findings

S2.02 Specimen labelled as

G2.01 Specimen dimensions
length mm x width mm x thickness mm
OR Cannot be recorded

S2.03 Specimen weight
 g

S2.04 Tumour capsule
Intact
Not intact

S2.05 No. of macroscopically visible tumour(s)

S2.06 MACROSCOPIC APPEARANCE OF LESION(S)

LESION 1

Location
Cortex
Medulla
Indeterminate
Other

Borders
Encapsulated
Infiltrative

Description

Size in greatest dimension
 mm

Distance to nearest excision margin
 mm

LESION 2

Location
Cortex
Medulla
Indeterminate
Other

Borders
Encapsulated
Infiltrative

Description

Size in greatest dimension
 mm

Distance to nearest excision margin
 mm

LESION 3

Location
Cortex
Medulla
Indeterminate
Other

Borders
Encapsulated
Infiltrative

Description

Size in greatest dimension
 mm

Distance to nearest excision margin
 mm

S2.07 APPEARANCE OF UNINVOLVED ADRENAL GLAND

Cortex
Unremarkable
Atrophic (thin)
Hypertrophic (thickened)
Not identified

Medulla

- Unremarkable
Atrophic (thin)
Hypertrophic (thickened)
Not identified

Cortical nodules

- Not assessable
Absent
Present

Size of largest nodule mm

OR Cannot be recorded

Medullary hyperplasia/nodule

- Not assessable
Absent
Present

G2.02 ANY ACCOMPANYING SPECIMENS

Lymph nodes

- Absent
Present

Other adjacent structures

G2.03 Other macroscopic comment

Microscopic findings

S3.01 Tumour type (refer to adjacent list)

G3.01 Tumour type variants (refer to adjacent list)

S3.02 Diameter of largest tumour

 mm

S3.01 Tumour type /G3.01 Variant

Adrenal cortical tumours

Adrenal cortical carcinoma
Adrenal cortical adenoma

Adrenal medullary tumours

Malignant pheochromocytoma
Benign pheochromocytoma
Composite pheochromocytoma/
paraganglioma

Extra-adrenal paraganglioma

Carotid body
Jugulotympanic
Vagal
Laryngeal
Aortico-pulmonary
Gangliocytic
Cauda equina
Orbital Nasopharyngeal
Extra-adrenal sympathetic paraganglioma
Superior and inferior para-aortic paraganglioma
Urinary bladder
Intrathoracic and cervical paravertebral

Other adrenal tumours

Adenomatoid tumour
Sex-cord stromal tumour
Soft tissue and germ cell tumours
Myelolipoma
Teratoma
Schwannoma
Ganglioneuroma
Angiosarcoma

FOR PHEOCHROMOCYTOMA

S3.13 Cellular pattern

(For pheochromocytoma, diffuse architecture is defined as large nests or diffuse growth >10%.)

- Non-diffuse
Zellballen
Large and irregular cell nests
Pseudorosette

S3.14 Cellularity

- Low
Moderate
High

S3.10 Central or confluent necrosis

- Absent
Present

S3.03 Adrenal vein or vena cava invasion

- Absent
Present

S3.04 Capsular invasion

- Absent
Present

S3.06 Extension into adipose tissue

Absent
Present

S3.07 Adjacent organs

Not involved
Involved

Involved organs:

S3.14 Cellular monotony

Absent
Present

Tumour cell spindling

Absent
Present

Profound nuclear pleomorphism

Absent
Present

Nuclear hyperchromasia

Absent
Present

S3.08 Mitotic rate

S3.09 Atypical mitotic figures

Absent
Present

FOR ADRENOCORTICAL TUMOURS

S3.11 Nuclear grade (Fuhrman)

grade 1
grade 2
grade 3
grade 4

S3.12 Tumour comprising clear or vacuolated cells

S3.08 Mitotic rate

S3.09 Atypical mitotic figures

Absent
Present

S3.13 Cellular pattern

(For conventional adrenocortical neoplasms diffuse growth is defined as more than 1/3 of the tumour forms patternless sheets of cells; trabecular, cord, columnar, alveolar or nesting is not considered to be diffuse)

Non-diffuse
Diffuse architecture

S3.10 Central or confluent necrosis

Absent
Present

S3.03 Adrenal vein or vena cava invasion

Absent
Present

S3.04 Capsular invasion

Absent
Present

S3.07 Adjacent organs

Not involved
Involved

Involved organs:

S3.05 Sinusoidal invasion (only if conventional or oncocytic adrenocortical neoplasm)

Absent
Present

FOR ALL TUMOURS

G3.02 MALIGNANT POTENTIAL

Scoring system (eg PASS, Weiss etc)

Score for malignant potential

S3.15 Non-tumour adrenal gland

Unremarkable

OR

Not identified/not assessable

OR select all that apply:

Adrenal cortical atrophy

Hyperplasia

Cortical nodules

Medullary hyperplasia/nodule

S3.16 Margin status

Clear Distance to closest margin

mm

Involved Involved margin(s)

S3.17 Lymph node status

Clear

Involved Number of positive nodes:

/

(No. of positive nodes out of total no. of nodes)

G3.03 Coexistent pathological abnormalities

Absent

Present Description:

G3.04 Other relevant microscopic comments

Ancillary test findings

S4.01 Ancillary tests

Not performed

Performed Complete the following:

Test result type 1 eg FISH, IHC

Result

Conclusion

Laboratory performing the test

Method

Substrate (eg cytology smears, fluid in special media, paraffin block, fresh

Person responsible for reporting

Test result type 2 eg FISH, IHC

Result

Conclusion

Laboratory performing the test

Method

Substrate (eg cytology smears, fluid in special media, paraffin block, fresh

Person responsible for reporting

Test result type 3 eg FISH, IHC

Result

Conclusion

Laboratory performing the test

Method

Substrate (eg cytology smears, fluid in special media, paraffin block, fresh

Person responsible for reporting

Synthesis and overview

S5.01 AJCC Tumour stage (see opposite)

T

N

M

Stage Grouping

S5.02 Year of publication and edition of cancer staging system

G5.01 Diagnostic summary

Include: specimen type, tumour type, diameter of largest tumour, tumour stage, completeness of excision.

S5.03 Overarching comment

S5.01 Tumour stage (AJCC)^{##}

Primary Tumour (T)

TX Primary tumour cannot be assessed

T0 No evidence of primary tumour

T1 Tumour 5 cm or less in greatest dimension, no extra-adrenal invasion

T2 Tumour greater than 5 cm, no extra-adrenal invasion

T3 Tumour of any size with local invasion, but not invading adjacent organs*

T4 Tumour of any size with invasion of adjacent organs*

*Adjacent organs include kidney, diaphragm, great vessels, pancreas, spleen, and liver.

Regional Lymph Nodes (N)

NX Regional lymph nodes cannot be assessed

N0 No regional lymph node metastasis

N1 Metastasis in regional lymph node(s)

Distant Metastasis (M)

M0 No distant metastasis

M1 Distant metastasis

Stage Grouping

Stage	T	N	M
I	T1	N0	M0
II	T2	N0	M0
III	T1	N1	M0
	T2	N1	M0
III	T3	N0	M0
IV	T3	N1	M0
	T4	N0	M0
	T4	N1	M0
	Any T	Any N	M1

^{##} Used with the permission of the American Joint Committee on Cancer (AJCC), Chicago, Illinois. The original source for this material is the AJCC Cancer Staging Manual, Seventh Edition (2010) published by Springer Science and Business Media LLC, www.springerlink.com

Worksheet prepared by:

On:

DD – MM – YYYY