

Protocol for the Examination of Biopsy Specimens From Patients With Carcinoma of the Urethra and Periurethral Glands

Version: 4.1.0.0

Protocol Posting Date: June 2021

The use of this protocol is recommended for clinical care purposes but is not required for accreditation purposes.

This protocol may be used for the following procedures AND tumor types:

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Procedure	Description	
Biopsy	Includes specimens designated biopsy or transurethral resection	
Tumor Type	Description	
	Includes invasive carcinomas of the urinary tract, including urothelial carcinoma and its morphological variants (squamous cell carcinoma, adenocarcinoma,	
	Müllerian carcinoma, neuroendocrine carcinoma, and sarcomatoid carcinoma)	

The following should NOT be reported using this protocol:

Procedure	
Resection (consider the Urethra Resection protocol)	
Transurethral resection	
Cytologic specimens	

The following tumor types should NOT be reported using this protocol:

Tumor Type
Lymphoma (consider the Hodgkin or non-Hodgkin Lymphoma protocols)
Sarcoma (consider the Soft Tissue protocol)

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With guidance from the CAP Cancer and CAP Pathology Electronic Reporting Committees.

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Accreditation Requirements

The use of this case summary is recommended for clinical care purposes but is not required for accreditation purposes. The core and conditional data elements are routinely reported. Non-core data elements are indicated with a plus sign (+) to allow for reporting information that may be of clinical value.

Summary of Changes

v 4.1.0.0

- General Reformatting
- Added LVI section
- Elements that are recommended for clinical care purposes are designated as Core and Conditional (indicated by bolded text), while Non-core elements are now indicated with a plus (+) sign

Reporting Template

Protocol Posting Date: June 2021 Select a single response unless otherwise indicated.			
CASE SUMMARY: (URETHRA: Biopsy) Standard(s): AJCC-UICC 8 This case summary is recommended for reporting biopsy specimens, but is not required for accreditation purposes.			
SPECIMEN (Note A)			
Specimen			
Urethra			
Other (specify):			
Not specified			
TUMOR			
Tumor Site (select all that apply)			
Male Penile urethra			
Bulbomembranous urethra			
Prostatic urethra			
Female			
Anterior urethra			
Posterior urethra			
Other			
Urethra, not otherwise specified:			
Histologic Type (Note B) (select all that apply)			
Urothelial Desille menunathadial accessor and a resolution assistant.			
Papillary urothelial carcinoma, noninvasive			
Papillary urothelial carcinoma, invasive Urothelial carcinoma in situ			
Urothelial carcinoma, invasive			
Urothelial carcinoma, nested (including large nested) variant			
Urothelial carcinoma, microcystic variant			
Urothelial carcinoma, micropapillary variant			
Urothelial carcinoma, lymphoepithelioma-like variant			
Urothelial carcinoma, plasmacytoid / signet ring cell / diffuse variant			
Urothelial carcinoma, sarcomatoid variant			
Urothelial carcinoma, giant cell variant			
Urothelial carcinoma, poorly differentiated variant			
Urothelial carcinoma, lipid-rich variant			
Urothelial carcinoma, clear cell variant			
Urothelial carcinoma with squamous differentiation			
+Percentage of Squamous Differentiation			
Specify percentage: %			
Other (specify):			
Cannot be determined			

Urothelial carcinoma with glandular differenti	ation
+Percentage of Glandular Differentiation	
Specify percentage:	%
Other (specify):	
Cannot be determined	
Urothelial carcinoma with trophoblastic differ	entiation
+Percentage of Trophoblastic Differentiation	
Specify percentage:	
Other (specify):	-
Cannot be determined	
Urothelial carcinoma with Müllerian differenti	ation
+Percentage of Müllerian Differentiation	
Specify percentage:	%
Other (specify):	. / •
Cannot be determined	
Squamous	
Squamous cell carcinoma	
Verrucous carcinoma	
Squamous cell carcinoma in situ (no invasive	e carcinoma identified)
Glandular	,
Adenocarcinoma	
Adenocarcinoma, enteric	
Adenocarcinoma, mucinous	
Adenocarcinoma, mixed	
Adenocarcinoma in situ (no invasive carcino	ma identified)
Tumors of Müllerian type	,
Clear cell carcinoma	
Endometrioid carcinoma	
Neuroendocrine Tumors	
Small cell neuroendocrine carcinoma	
+Percentage of Small Cell Neuroendocrine	Component
Specify percentage:	_ %
Other (specify):	
Cannot be determined	
Large cell neuroendocrine carcinoma	
+Percentage of Large Cell Neuroendocrine	Component
Specify percentage:	_ %
Other (specify):	
Cannot be determined	
Well-differentiated neuroendocrine tumor	
+Percentage of Well-differentiated Neuroen	docrine Component
Specify percentage:	%
Other (specify):	-
Cannot be determined	
Other	
Other histologic type not listed (specify):	
Carcinoma, type cannot be determined:	
+Histologic Type Comment:	

Histologic Grade (Note C)
For urothelial carcinoma, other variants, or divergent differentiation
Low-grade
High-grade
For squamous cell carcinoma or adenocarcinoma
G1, well differentiated
G2, moderately differentiated
G3, poorly differentiated
GX, cannot be assessed:
Other
Other (specify):
Cannot be determined:
Not applicable:
Tumor Extent (Note D)
Male
Carcinoma of penile and bulbomembranous urethra
Noninvasive urothelial papillary carcinoma
Carcinoma in situ
Invades subepithelial connective tissue
Invades adjacent structure(s)
Corpus spongiosum
Periurethral muscle
Corpus cavernosum
Bladder wall
Rectum
Necture Other (specify):
Carcinoma of the prostatic urethra
Carcinoma in situ, involving prostatic urethra
Carcinoma in situ, involving prostatic ducts
Invades urethral subepithelial connective tissue immediately underlying the urothelium
Invades prostatic stroma surrounding ducts either by direct extension from the urothelial surface o
by invasion from prostatic ducts
Invades periprostatic fat
Invades adjacent structure(s)
Extraprostatic invasion of the bladder wall
Rectum
Other (specify):
Female Nepipyagiya urathalial papillary carainama
Noninvasive urothelial papillary carcinoma
Carcinoma in situ
Invades subepithelial connective tissue
Invades adjacent structure(s)
Periurethral muscle (fibromuscular and adipose tissue)
Bladder wall
Rectum
Other (specify):
Other Connet he determined:
Cannot be determined:
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+Lymphovascular Invasion
Not identified
Present
Cannot be determined:
+Tumor Configuration (select all that apply)
Papillary
Solid / nodule
Flat
Ulcerated
Other (specify):
Cannot be determined:
+Tumor Comment:
ADDITIONAL FINDINGS
+Associated Epithelial Lesions (select all that apply)
None identified
Condyloma
Squamous dysplasia (low, intermediate, high grade)
Urothelial papilloma
Urothelial papilloma, inverted type
Papillary urothelial neoplasm, low malignant potential (PUNLMP)
Urothelial proliferation of uncertain malignant potential
Urothelial dysplasia
Other (specify):
Cannot be determined:
+Additional Findings (select all that apply)
Keratinizing squamous metaplasia
Inflammation / regenerative changes
Therapy-related changes (specify):
Cautery artifact
Urethritis cystica et glandularis
Intestinal metaplasia
Other (specify):
COMMENTS
Comment(s):

Explanatory Notes

A. History

A relevant history is important for interpretation of urethral biopsies. A history of renal stones, recent urinary tract procedures, infections, obstruction, or prior therapy (intravesical or systemic chemotherapy, local radiation) can lead to reactive epithelial changes potentially mimicking malignancy. Any neoplasms previously diagnosed should be specified, including the histologic type, primary site, and histologic grade.

B. Histologic Type

Carcinomas of the urethra vary in histologic type, depending on type of epithelium lining the urethra in a given anatomic location. 1.2.3.4 In women, squamous cell carcinoma is the most common histologic subtype (approximately 75%) and is most common in the anterior urethra (distal third). Urothelial carcinoma is next in frequency, followed by adenocarcinoma (approximately 10% to 15% each). Clear cell adenocarcinomas comprise a significant proportion of adenocarcinomas in women but are quite rare in men.⁵ In the male, most tumors involve the bulbomembranous urethra, followed by penile urethra and prostatic urethra. Most carcinomas of the male urethra (80%) are squamous cell carcinoma, followed by urothelial origin. As in women, urothelial carcinomas are typically more proximal. Primary urethral adenocarcinomas are rare in men. Adenocarcinomas may rarely arise from the periurethral Skene's (female) or Littre's (male) glands.4 The distinction between a urothelial carcinoma with divergent squamous, glandular, or Müllerian differentiation and a pure squamous cell carcinoma, adenocarcinoma or Müllerian is rather arbitrary. Most authorities, including the 2016 World Health Organization (WHO) classification, require a pure histology of squamous cell carcinoma, adenocarcinoma, or Müllerian to designate a tumor as such, all others with recognizable papillary, invasive, or flat carcinoma in situ (CIS) urothelial component being considered as urothelial carcinoma with divergent differentiation. A malignant neoplasm with small cell neuroendocrine carcinoma component arising in the urinary tract is designated as small cell carcinoma.6

2016 WHO Classification of Tumors of the Urothelial Tract

Urothelial tumors

Infiltrating urothelial carcinoma

Nested, including large nested

Microcystic

Micropapillary

Lymphoepithelioma-like

Plasmacytoid/signet ring cell/diffuse

Sarcomatoid

Giant cell

Poorly differentiated

Noninvasive urothelial lesions

Urothelial carcinoma in situ

Noninvasive papillary urothelial carcinoma, low grade

Noninvasive papillary urothelial carcinoma, high grade

Papillary urothelial neoplasm of low malignant potential

Urothelial papilloma

Inverted urothelial papilloma

Urothelial proliferation of uncertain malignant potential

Urothelial dysplasia

Squamous cell neoplasms

Squamous cell carcinoma Verrucous carcinoma Squamous cell papilloma

Glandular neoplasms

Adenocarcinoma, NOS

Enteric

Mucinous

Mixed

Villous adenoma

Urachal carcinoma

Tumors of Mullerian type

Clear cell carcinoma Endometrioid carcinoma

Neuroendocrine tumors

Small cell neuroendocrine carcinoma Large cell neuroendocrine carcinoma Well differentiated neuroendocrine tumor Paraganglioma

References

- 1. Amin MB, Young RH. Primary carcinomas of the urethra. *Semin Diag Pathol.* 1997;14(2):147-160.
- 2. Reuter V.E. Urethra. In: Bostwick DG, Eble JN, eds. *Urologic Surgical Pathology.* St. Louis, MO: Mosby Year Book, Inc; 1997:223-230.
- 3. Reuter VE. The urothelial tract: renal pelvis, ureter, urinary bladder and urethra. In: Mills SE, Carter D, Greenson JK, Oberman HA, Reuter VE, Stoler MH, eds. *Sternberg's Diagnostic Surgical Pathology.* 4th ed. Philadelphia, PA: Lippincott Williams and Wilkins; 2004:2035-2081.
- 4. Murphy WM, Grignon DJ, Perlman EJ. Tumors of the kidney, bladder, and related urinary structures. In: *Atlas of Tumor Pathology*. 4th series. Fascicle 1. Washington, DC: American Registry of Pathology; 2004.
- 5. Oliva E, Young RH. Clear cell adenocarcinoma of the urethra: a clinicopathologic analysis of 19 cases. *Mod Pathol.* 1996;9:513-520.
- 6. Lopez-Beltran A, Sauter G, Gasser T, et al. Infiltrating urothelial carcinoma. In: Eble JN, Sauter G, Epstein JI, Sesterhenn IA, eds. *World Health Organization Classification of Tumours: Pathology and Genetics of Tumours of the Urinary System and Male Genital Organs.* Lyon, France: IARC Press; 2004:97.

C. Histologic Grade

Squamous cell carcinoma and adenocarcinoma are graded on a 3-tiered system as well differentiated (grade 1), moderately differentiated (grade 2), or poorly differentiated (grade 3).

For urothelial neoplasia, flat intraepithelial lesions and papillary and invasive lesions are graded separately. Due to variable classification systems and the need for a universally acceptable system, the World Health Organization/International Society of Urological Pathology (WHO/ISUP) consensus classification was proposed and has been adopted in the 2016 WHO classification 1.2 and has been validated by many studies to be prognostically significant. Other systems (that were being used

previously) may still be used according to institutional preferences Tumor grade according to both the WHO/ISUP (1998) system and the older WHO (1973) system may be concurrently used.^{3.4}

Flat and papillary urothelial hyperplasia has been renamed as "urothelial proliferation of uncertain malignant potential" in the 2016 WHO classification.

References

- 1. Moch H, Humphrey PA, Ulbright TM, Reuter VE. *WHO Classification of Tumours of the Urinary System and Male Genital Organs.* Geneva, Switzerland: WHO Press; 2016.
- 2. Sauter G, Algaba F, Amin MB, et al. Non-invasive urothelial tumours. In: Eble JN, Sauter G, Epstein JI, Sesterhenn IA, eds. *World Health Organization Classification of Tumours: Pathology and Genetics of Tumours of the Urinary System and Male Genital Organs.* Lyon, France: IARC Press; 2004:110.
- 3. Epstein JI, Amin MB, Reuter VR, Mostofi FK, the Bladder Consensus Conference Committee. The World Health Organization/ International Society of Urological Pathology Consensus classification of urothelial (transitional cell) neoplasms of the urinary bladder. *Am J Surg Pathol.* 1998;22(12):1435-1448.
- 4. Mostofi FK. Histological typing of urinary bladder tumours. In: *WHO Histological Classification of Tumours*. No. 10. Geneva, Switzerland: World Health Organization; 1973.

D. Extent of Invasion

A critical role of the surgical pathologist is to diagnose the depth/extent of invasion into the tissues surrounding the urethra. The surrounding anatomic structures vary by gender and location within the urethra but include the subepithelial connective tissue, corpus spongiosum, corpus cavernosum, prostate, periurethral muscle, extraprostatic soft tissue, anterior vagina, bladder neck, or other adjacent organs. In the prostatic urethra, invasion may arise from a tumor lining the urethral lumen or from carcinoma in situ colonizing prostatic ducts. The pT1 designation should only be applied to superficial invasion arising from the urethral lining; invasion arising from the prostatic ducts is designated as at least pT2. In papillary urothelial tumors, invasion occurs most often at the base of the tumor and less frequently in the stalk.

References

- 1. Mostofi FK. Histological typing of urinary bladder tumours. In: *WHO Histological Classification of Tumours*. No. 10. Geneva, Switzerland: World Health Organization; 1973.
- 2. Amin MB, Edge SB, Greene FL, et al., eds. *AJCC Cancer Staging Manual*. 8th ed. New York, NY: Springer; 2017