

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:

Procedure	Description
Biopsy	Includes specimens designated biopsy or transurethral resection
Tumor Type	Description
Carcinomas	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)

Authors

Gladell P. Paner, MD*; Jesse K. McKenney, MD*; Ming Zhou, MD, PhD*; Lara R. Harik, MD; Robert Allan, MD; Mahul B. Amin, MD; Jonathan I. Epstein, MD; David J. Grignon, MD; Peter A. Humphrey, MD, PhD; Esther Oliva, MD; Jason Pettus, MD; Victor E. Reuter, MD; John R. Srigley, MD.

With guidance from the CAP Cancer and CAP Pathology Electronic Reporting Committees. * Denotes primary author.

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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

- 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 differentiation
+Percentage of Glandular Differentiation
Specify percentage: %
Other (specify):
Cannot be determined
Urothelial carcinoma with trophoblastic differentiation
+Percentage of Trophoblastic Differentiation
Specify percentage: %
Other (specify):
Cannot be determined
Urothelial carcinoma with Müllerian differentiation
+Percentage of Müllerian Differentiation
Specify percentage: %
Other (specify):
Cannot be determined
Squamous
Squamous cell carcinoma
Verrucous carcinoma
Squamous cell carcinoma in situ (no invasive carcinoma identified)
Glandular
Adenocarcinoma
Adenocarcinoma, enteric
Adenocarcinoma, mucinous
Adenocarcinoma, mixed
Adenocarcinoma in situ (no invasive carcinoma identified)
Tumors of Müllerian type
Clear cell carcinoma
Endometrioid carcinoma
Neuroendocrine Tumors
Small cell neuroendocrine carcinoma
+Percentage of Small Cell Neuroendocrine Component Specify percentage: %
F J
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 Neuroendocrine Component
Specify percentage. %
Specify percentage:%
Other (specify):
Other (specify): Cannot be determined
Other (specify): Cannot be determined <i>Other</i>
Other (specify): Cannot be determined <i>Other</i> Other histologic type not listed (specify):
Other (specify): Cannot be determined <i>Other</i>

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
 - 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 or by invasion from prostatic ducts
 - Invades periprostatic fat
 - Invades adjacent structure(s)
 - Extraprostatic invasion of the bladder wall
 - Rectum
 - ____ Other (specify): _____

Female

- ___ 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

- ____ Cannot be determined: _____
- ____ No evidence of primary tumor

	+Lym	phovas	cular	Invas	ion
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- Not identified
- Present
- Cannot be determined: ____

+Tumor Configuration (select all that apply)

- ____ Papillary
- Solid / nodule
- ___ Flat
- Ulcerated
- ____ Other (specify): ___
- ____ Cannot be determined: _____

+T	umor	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 men⁵. 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

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- 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.
- 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.
- 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.
- 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.¹ 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 pT2.² 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