EAU GUIDELINES ON PRIMARY URETHRAL CARCINOMA

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Epidemiology

Primary Urethral Carcinoma is a rare cancer, accounting for < 1% of all genitourinary malignancies. The age-standardised ratio is 4.3/million in men and 1.5/million in women, with a male to female ratio of 2.9:1.

Aetiology

Predisposing factors in males include urethral strictures, chronic irritation after intermittent catheterisation/ urethroplasty, external beam irradiation therapy, radioactive seed implantation, chronic urethral inflammation following sexually transmitted diseases (especially human papilloma virus) and lichen sclerosus. In females, urethral diverticula and recurrent urinary tract infections have been associated with the development of primary urethral carcinoma.

Staging and Grading systems

The 2017 TNM classification (8th edition) is used for the staging of urethral carcinoma. Of note, a separate staging system exists for urothelial carcinoma (UC) of the prostatic urethra.

T - Primary Tumour			
TX	Primary tumour cannot be assessed		
T0	No evidence of primary tumour		
Urethr	Urethra (male and female)		
Та	Non-invasive papillary, polypoid, or verrucous		
	carcinoma		
Tis	Carcinoma in situ		
T1	Tumour invades subepithelial connective tissue		
T2	Tumour invades any of the following: corpus		
	spongiosum, prostate, periurethral muscle		
T3	Tumour invades any of the following: corpus		
	cavernosum, beyond prostatic capsule, anterior		
	vagina, bladder neck (extraprostatic extension)		
T4	Tumour invades other adjacent organs (invasion of		
	the bladder)		
	lial (transitional cell) carcinoma of the prostate		
Tis pu	Carcinoma in situ, involvement of prostatic urethra		
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Tis pd	Carcinoma in situ, involvement of prostatic ducts		
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M - Distant Metastasis	
M0	No distant metastasis
M1	Distant metastasis

Histopathology

Urothelial carcinoma of the urethra is the predominant histological type in men with primary urethral carcinoma followed by squamous cell carcinoma (SCC) and adenocarcinoma (AC).

In women, recent studies report higher rates of adenocarcinoma, followed by SCC rather than UC. Specimen handling should follow the general rules as published by the International Collaboration on Cancer Reporting.

Recommendation for staging and grading	Strength rating
Use the 2017 TNM classification and	Strong
2004/2016 WHO grading systems for	
pathological staging and grading of primary	
urethral carcinoma.	

Diagnosis

Diagnosis of primary urethral carcinoma is based on clinical examination, urine cytology, urethroscopy with biopsy and cross-sectional imaging for the assessment of the primary tumour, lymph nodes (LNs) and distant organs. Patients with clinically enlarged inguinal or pelvic LNs often exhibit pathological LN metastasis.

Recommendations	Strength rating
Use urethrocystoscopy with biopsy and urinary cytology to diagnose urethral carcinoma.	Strong
Assess the presence of distant metastases by computed tomography of the thorax and abdomen/pelvis.	Strong
Use pelvic magnetic resonance imaging to assess the local extent of urethral tumour and regional lymph node enlargement.	Strong

Prognosis

The majority of patients are diagnosed late, with local symptoms due to advanced disease and the prognosis is poor.

Risk factors for survival include age, race, tumour stage, grade, nodal stage, presence of distant metastasis, histological type, tumour size, tumour location, concomitant bladder cancer and the type and modality of treatment, as well as the type of treating facility. In locally advanced UC and SCC of the urethra, treatment in academic centres improves overall survival (OS).

Disease management

Primary disease in males

Distal urethral tumours exhibit significantly improved survival rates compared with proximal tumours. Therefore, optimising treatment of distal urethral carcinoma has become the focus of clinicians to improve functional outcome and quality of life, while preserving oncological safety. Penile-preserving surgery for tumours confined to the corpus spongiosum (stage ≤ T2) using various reconstructive techniques has been investigated. In distal urethral tumours performing a partial urethrectomy with a minimal safety margin does not increase the risk of local recurrence when complete circumferential assessment of the margins shows no evidence of disease.

Recommendations	Strength rating
Offer distal urethrectomy as an alternative	Weak
to penile amputation in localised distal	
urethral tumours, if negative surgical	
margins can be achieved intra-operatively.	
Ensure complete circumferential	Strong
assessment of the proximal urethral margin	
if penile-preserving surgery is intended.	

Localised primary disease in females

In women with distal tumours, urethra-sparing surgery and local radiotherapy (RT) present alternatives to primary urethrectomy but are associated with increased risk of tumour recurrence and local toxicity.

Recommendations	Strength rating
Offer urethra-sparing surgery, as an	Weak
alternative to primary urethrectomy, to	
females with distal urethral tumours, if	
negative surgical margins can be achieved	
intra-operatively.	
Offer local radiotherapy, as an alternative to	Weak
urethral surgery, to females with localised	
urethral tumours, but discuss local toxicity.	

Multimodal therapy in advanced disease in both males and females

Multimodal therapy in primary urethral carcinoma consists of definitive surgery plus chemotherapy with the option of additional RT. Multimodal therapy is often underutilised in locally advanced disease. It confers an OS benefit in primary urethral carcinoma of urothelial origin.

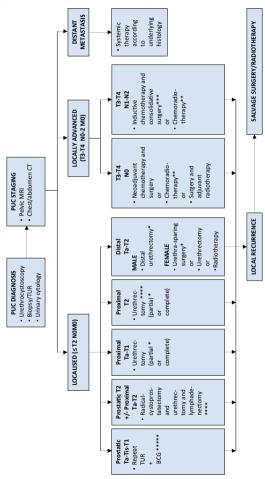
Recommendations	Strength rating
Refer patients with advanced urethral	Strong
carcinoma to academic centres.	
Discuss treatment of patients with locally	Strong
advanced urethral carcinoma within a	
multidisciplinary team of urologists,	
radiation-oncologists, and oncologists.	
In locally advanced urethral carcinoma,	Weak
use cisplatin-based chemotherapeutic	
regimens with curative intent prior to	
surgery.	
In locally advanced squamous cell	Weak
carcinoma (SCC) of the urethra, offer the	
combination of curative radiotherapy (RT)	
with radiosensitising chemotherapy for	
definitive treatment and genital	
preservation.	
Offer salvage surgery or RT to patients	Weak
with urethral recurrence after primary	
treatment.	
Offer inguinal lymph node (LN) dissection	Weak
to patients with limited LN-positive urethral	
SCC.	

Treatment of urothelial carcinoma of the prostate

Local conservative treatment with extensive transurethral resection (TUR) and subsequent bacillus Calmette-Guerin (BCG) instillation is effective in patients with Ta or Tis prostatic urethral carcinoma. Patients undergoing TUR of the prostate for prostatic urethral carcinoma prior to BCG treatment show superior complete response rates compared to those who do not.

Recommendations	Strength rating
Offer a urethra-sparing approach with	Strong
transurethral resection (TUR) and bacillus	
Calmette-Guérin (BCG) to patients with	
non-invasive urethral carcinoma or	
carcinoma in situ of the prostatic urethra	
and prostatic ducts.	
In patients not responding to BCG, or in	Weak
patients with extensive ductal or stromal	
involvement, perform a cystoprostatectomy	
with extended pelvic lymphadenectomy.	

Figure 1: Management of primary urethral carcinoma



- Ensure complete circumferential assessment if penilepreserving/urethra-sparing surgery or partial urethrectomy is intended.
- ** Squamous cell carcinoma.
- *** Regional lymphadenectomy should be considered in clinically enlarged lymph nodes.
- **** Consider neoadjuvant chemotherapy.
- ***** In extensive or BCG-unresponsive disease: consider (primary) cystoprostatectomy +/- urethrectomy + lymphadenectomy.

BCG = bacillus Calmette-Guérin; CT = computed tomography; MRI = magnetic resonance imaging; PUC = primary urethral carcinoma; TUR = transurethral resection.

Follow-up

Given the low incidence of primary urethral carcinoma, followup has not been systematically investigated. Therefore, it seems reasonable to tailor surveillance regimens according to patients' individual risk factors. In patients undergoing urethra-sparing surgery, it seems prudent to advocate a more extensive follow-up with urinary cytology, urethrocystoscopy and cross-sectional imaging despite the lack of specific data.

This short booklet text is based on the more comprehensive EAU Guidelines (ISBN 978-94-92671-16-5) available to all members of the European Association of Urology at their website, http://www.uroweb.org/guidelines/.