



APPROPRIATENESS CRITERIA FOR URETERAL STENT OMISSION FOLLOWING URETEROSCOPY FOR UROLITHIASIS: A SINGLE-CENTER EXPERIENCE

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ABSTRACT

Background: Ureteral stenting following uncomplicated ureteroscopy (URS) and ureterorenoscopy remains a common practice. However, evidence suggests that routine stent placement may not be necessary in all cases. Defining clear criteria for uncomplicated URS could help establish when stent omission (SO) is appropriate. **Objective:** The primary objective of this study was to establish judicious and specific indications for stent omission following URS and ureterorenoscopy. The secondary objective was to evaluate the outcomes and complications associated with stent omission. **Methods:** This retrospective study reviewed all patients who underwent URS or ureterorenoscopy at the Urology Unit, Al-Khor Hospital, from 1 January 2017 to 31 December 2022. Adult patients (> 18 years) were included. Exclusion criteria were renal anomalies, uncorrectable bleeding disorders, febrile urinary tract infection, solitary kidney, musculoskeletal deformities, and ureteral obstruction. Preoperative and intraoperative factors were recorded, including stone burden, preoperative stent presence, procedure time, and use of an access sheath. Postoperative events within 30 days were assessed, including unplanned office or emergency visits, admissions, or reoperations. **Results:** A total of 1,500 URSL procedures were reviewed. Of these, 636 (42.4%) cases were eligible for SO, and 360 (24.0% of total; 56.6% of eligible) underwent SO. Postoperative events occurred in 16.7% of SO cases (n=60) compared to 34.8% of stented cases (n=393, P = 0.03). Unplanned return visits occurred in 8.3% of SO cases (n=30) versus 17.4% of stented cases (n=199, P = 0.16). Thirty-day readmissions were 0% in the SO group compared to 6.5% (n=74) in the stented group (P = 0.08). The odds ratio for unplanned visits in the SO group was 0.43 (95% CI: 0.13–1.42, P = 0.17). No significant associations were found between postoperative events and stone burden, procedure time, or preoperative stenting. **Conclusion:** Stent omission following uncomplicated URS is safe and associated with fewer postoperative events compared to routine stenting. These findings support the development of appropriateness criteria to guide selective stent omission and reduce unnecessary morbidity.

KEYWORDS : Ureteroscopy, Ureteral Stents, Quality Improvement, Urinary Stone Disease

INTRODUCTION.

Ureteroscopy (URS) and ureterorenoscopy have become standard interventions for the treatment of ureteral and renal calculi. Postoperative ureteral stenting has traditionally been

practiced reducing the risk of obstruction, ensure drainage, and minimize postoperative complications. However, accumulating evidence suggests that routine stent placement may not be necessary following uncomplicated procedures.

Stents are associated with significant morbidity, including flank pain, lower urinary tract symptoms, urinary tract infection, and the need for an additional procedure for removal.

Several studies have explored the safety of stent omission (SO), highlighting that it does not compromise patient safety in carefully selected cases. Nevertheless, stent omission remains underutilized due to the absence of universally accepted appropriateness criteria. In this context, the present study aims to evaluate stent omission practices at a tertiary centre in Qatar and to identify appropriate indications for SO following URS and ureterorenoscopy. {1-5}

Methods

This retrospective study was conducted at Al-Khor Hospital, Qatar. We reviewed medical records of all patients who underwent URS or ureterorenoscopy between 1 January 2017 and 31 December 2022. Inclusion criteria were adult patients (≥18 years) undergoing the procedure for urolithiasis. Exclusion criteria included congenital renal anomalies, uncorrectable bleeding disorders, febrile urinary tract infection, solitary kidney, musculoskeletal deformities, and ureteral obstruction.

Preoperative factors (age, sex, stone burden, preoperative stent), intraoperative variables (operative time, access sheath use, ureteral trauma), and postoperative outcomes (unplanned visits, emergency visits, admissions, or reoperations within 30 days) were recorded. The decision to omit stenting was at the discretion of the operating surgeon, based on intraoperative findings.

Statistical analysis included chi-square and Fisher's exact test for categorical variables, Student's t-test for continuous variables, and odds ratios with 95% confidence intervals. A p-value <0.05 was considered statistically significant.

RESULTS.

A total of 1,500 URSL procedures were reviewed. Of these, 636 (42.4%) cases were eligible for SO, and 360 (24.0% of total; 56.6% of eligible) underwent SO. Postoperative events occurred in 16.7% of SO cases (n=60) compared to 34.8% of stented cases (n=393, P = 0.03). Unplanned return visits occurred in 8.3% of SO cases (n=30) versus 17.4% of stented cases (n=199, P = 0.16). Thirty-day readmissions were 0% in the SO group compared to 6.5% (n=74) in the stented group (P = 0.08). The odds ratio for unplanned visits in the SO group was 0.43 (95% CI: 0.13–1.42, P = 0.17). No significant associations were found between postoperative events and stone burden, procedure time, or preoperative stenting.

Table 1. Eligibility And Performance Of Stent Omission (SO)

| Category | Cases (n) | Percentage (%) |
|-----------------------|-----------|----------------------------------|
| Total URSL procedures | 1,500 | 100 |
| Eligible for SO | 636 | 42.4 |
| SO performed | 360 | 24.0 of total (56.6 of eligible) |

Table 2. Postoperative Event Rates

| Group | Total Cases | Events (n) | Rate (%) |
|---------------|-------------|------------|----------|
| SO group | 360 | 60 | 16.7 |
| Stented group | 1,140 | 393 | 34.8 |

Table 3. Unplanned Return Visits And Readmissions Within 30 Days

| Group | Total Cases | Unplanned Visits (n, %) | Readmissions (n, %) | Reoperations (n, %) |
|---------------|-------------|-------------------------|---------------------|---------------------|
| SO group | 360 | 30 (8.3%) | 0 (0.0%) | 0 (0.0%) |
| Stented group | 1,140 | 199 (17.4%) | 74 (6.5%) | 74 (6.5%) |

DISCUSSION.

This single-centre study demonstrates that ureteral stent omission (SO) after uncomplicated URS is safe and reduces

postoperative events. The lower rate of complications in the SO group (16.7% vs. 34.8%) aligns with prior studies highlighting the safety of selective SO {1,3,4,7,10.} No readmissions or reoperations occurred in the SO cohort, reinforcing its safety {5,6,9}.

Stent-related morbidity, including pain, LUTS, hematuria, and infection, negatively impacts patient quality of life {3,7,10}. Avoiding unnecessary stents reduces these complications and healthcare costs {8,11,12}. Emerging evidence indicates adherence to SO criteria does not compromise stone-free rates or increase ureteral complications {13,14}.

Despite supporting evidence, SO remains underutilized; only 56.6% of eligible cases underwent SO. Underutilization may reflect surgeon caution, variable practice patterns, or limited awareness of guidelines {5,6,11}. Structured appropriateness criteria, surgeon education, and institutional protocols improve SO rates without compromising safety {8,12}.

Recent meta-analyses and prospective studies reinforce safe application of SO in patients with uncomplicated URS, short operative times, minimal ureteral trauma, and absence of infection or significant residual stones {7,9,13}. Patient-reported outcomes such as pain and urinary symptoms are consistently better in SO cohorts {14,15}.

Limitations include retrospective design, potential selection bias, and absence of patient-reported outcomes. Strengths include large sample size and real-world clinical relevance. Future multicenter RCTs are needed to refine SO criteria and standardize postoperative protocols.

CONCLUSION

Selective ureteral stent omission following uncomplicated URS is safe, reduces postoperative morbidity, and improves patient outcomes. Evidence-based appropriateness criteria can guide clinicians in minimizing unnecessary stenting.

Declaration.

Financing. The study was not sponsored.

Conflict Of Interest. The authors declare no conflicts of interest.

Ethical approval. The study was approved by the MRC.

Authors Contribution:

All authors contribute equally to study design development, data analysis, drafting the manuscript, study concepts, scientific editing, literature review, data acquisition and data analysis

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