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Brief Correspondence

Adult Ureteroscopy (A-URS) Checklist: A New Tool To Standardise Reporting in Endourology

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Abstract

Ureteroscopy is increasingly being used for urolithiasis. Technological innovations have been accompanied by wide variations in practice patterns. At the same time, a common finding in many studies, especially systematic reviews, is that the heterogeneity of outcome measurements and lack of standardisation can limit both the reproducibility and generalisability of study findings. While many checklists are available to improve study reporting, there are no ureteroscopy-specific ones. The Adult-Ureteroscopy (A-URS) checklist is a practical aid for both researchers and reviewers for studies in this field. It contains five main sections (study details, preoperative, operative, postoperative, and long term data) and a total of 20 items. **Patient summary:** We developed a checklist to improve how studies on ureteroscopy (insertion of a telescope through the urethra to inspect the urinary tract) in adults are reported. This could help in advancing the field and improving patient outcomes, as all the key information is captured.

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Checklists are well established in the setting of study reporting. Examples include Strengthening the Reporting of Observational studies in Epidemiology (STROBE) guideline for observational studies and the Consolidated Standards of Reporting Trials (CONSORT) statement [1,2]. These are valuable tools for both authors and reviewers when making an assessment for publication. However, procedure-specific tools are lacking in surgery, which is also the case for endourology. We recently developed the Paediatric Ureteroscopy (P-URS) checklist to improve standardisation of parameters reported in this clinical area [3]. As for paediatric URS, practice patterns vary widely in adult URS [4]. Topics of continued debate and disagreement include the role of access sheaths, laser settings, and prestenring. A core reason for this variation is the heterogeneity of data collection and reporting. This is a common conclusion in literature reviews [5,6].

While there is an ever-increasing volume of URS publications, which is of course welcome, the path towards gaining clarity and resolution will be slow without more efforts to address these issues. Results can also be misleading when subtleties in study protocols are not clearly defined. This can cause confusion for readers, especially for more junior faculty without the experience to “read between the lines” regarding a study’s results. For example, it can be claimed that a particular technique for URS is superior as the stone-free rate (SFR) was higher than in another study. However, the latter study may have pooled SFR data for renal and ureteral stones but not made it clear that 90% of the burden was represented by distal ureteric stones. Even such seemingly small points can lead to quick dissemination of the conclusion that one modality is clearly superior. This is just one example of many.



| Item | Recommendation |
|--|--|
| <i>Study details</i> | |
| Study aim | <input type="checkbox"/> Description of primary and secondary aims of the study |
| Study setting | <input type="checkbox"/> Hospital setting and volume of cases performed each year |
| Study design | <input type="checkbox"/> Retrospective (clinical audit), prospective (comparative) study (non-randomised), randomised controlled trial |
| Selection criteria | <input type="checkbox"/> Description of how patients were enrolled e.g., consecutively <input type="checkbox"/> Inclusion criteria <input type="checkbox"/> Exclusion criteria <input type="checkbox"/> Indication for surgery |
| <i>Pre-operative</i> | |
| Operating team | <input type="checkbox"/> Experience of surgeon e.g., case volume <input type="checkbox"/> Number of surgeons involved <input type="checkbox"/> State whether resident involvement |
| Patient information | <input type="checkbox"/> Previous treatments undergone by patient during that stone episode <input type="checkbox"/> Comorbidities <input type="checkbox"/> Pre-operative urine culture <input type="checkbox"/> ASA grade |
| Medical therapy | <input type="checkbox"/> Specify if any patients had MET (either pre or post operatively) <input type="checkbox"/> Use of any medications (alpha blocker or anticholinergics) for stent symptoms |
| Imaging | <input type="checkbox"/> Breakdown of imaging modalities used <input type="checkbox"/> Time interval of imaging pre and post treatment |
| Preoperative stone status | <input type="checkbox"/> Stone size, largest diameter (for biggest stone if multiple) <input type="checkbox"/> Stone volume (include formula for calculation) <input type="checkbox"/> Stone density (Hounsfield units) <input type="checkbox"/> Stone location (both in kidney -UC, MC, LC and central RP - and ureter) <input type="checkbox"/> Preoperative stone obstruction (hydronephrosis/proximal dilatation) <input type="checkbox"/> Stone multiplicity <input type="checkbox"/> Stone access (e.g., diverticulum or urinary diversion) |
| Prior drainage | <input type="checkbox"/> Proportion of patients with indwelling stent <input type="checkbox"/> Proportion of patients with indwelling nephrostomy |
| <i>Operative</i> | |
| Timing | <input type="checkbox"/> Breakdown of elective and emergency cases <input type="checkbox"/> Operative and laser time <input type="checkbox"/> Type of anaesthesia used |
| Equipment and description of URS procedure | <input type="checkbox"/> Type and dimensions of ureteroscope(s) <input type="checkbox"/> State use of safety guidewire (mandatory, optional, not in use) <input type="checkbox"/> Energy source for lithotripsy <input type="checkbox"/> Laser type and power output <input type="checkbox"/> Start-up settings <input type="checkbox"/> Extras: Laser activation time, total laser energy <input type="checkbox"/> Fragmentation strategy: fragmentation or dusting or both <input type="checkbox"/> Stone retrieval: Y/N (if yes, state device) <input type="checkbox"/> Use of access sheath (including size) <input type="checkbox"/> Irrigation used (preferably information on temperature – pre-warmed, room-tempered or chilled – and irrigation pumps/gravitational) <input type="checkbox"/> Total radiation dose used during the procedure |

Fig. 1 – Adult Ureteroscopy (A-URS) checklist. Items of high priority are highlighted in bold font.

| | |
|---------------------------------------|---|
| | <input type="checkbox"/> Rate of negative ureteroscopy (i.e., no stone found) |
| Access success | <input type="checkbox"/> Success at accessing upper urinary tract at the initial surgery <input type="checkbox"/> If active dilatation (e.g., balloon) - provide details of equipment and settings |
| Complications | <input type="checkbox"/> Report any intra operative complications and status of ureter on exit <input type="checkbox"/> Use a validated grading tool wherever possible <input type="checkbox"/> State if procedure was interrupted or not due to the occurrence of intraoperative complication/adverse event |
| Exit strategy | <input type="checkbox"/> Breakdown of patients receiving stent, ureteral catheter or nephrostomy <input type="checkbox"/> Specify if stent modification used e.g., Stent on string <input type="checkbox"/> Duration of indwelling stent (or other) <input type="checkbox"/> Number of patients receiving post op urinary catheter |
| <i>Post operative</i> | |
| Follow up | <input type="checkbox"/> Length of hospital stay <input type="checkbox"/> Time interval when follow up performed <input type="checkbox"/> Stone composition (if available) <input type="checkbox"/> Use of patient reported outcome measure (PROM) |
| SFR | <input type="checkbox"/> Definition and imaging used to calculate SFR <input type="checkbox"/> Include zero fragment definition <input type="checkbox"/> Breakdown of SFR according to ureteral and renal stones rather than pooled result only <input type="checkbox"/> Give initial SFR after first procedure as well as final SFR after any additional URS treatments required. <input type="checkbox"/> Provide total and average no. of URS procedures to become stone free. |
| Auxiliary treatment | <input type="checkbox"/> Give details on any further intervention e.g., PCNL or SWL required to become stone free and provide a further SFR result including this accordingly. |
| Complications | <input type="checkbox"/> Use a validated grading tool (e.g., Clavien-Dindo) <input type="checkbox"/> Specify if complications were per patient, procedure or renal unit <input type="checkbox"/> Include complications occurring during all stages of stone treatment i.e., pre-stenting, formal stone surgery, post operative and stent removal <input type="checkbox"/> Indicate if complication rate is for URS procedure only or whether it includes additional procedures such as stent removal <input type="checkbox"/> Mention how the complications were resolved (e. antibiotics, hospital readmission, stent insertion etc.) |
| <i>Long term</i> | |
| Additional information (if available) | <input type="checkbox"/> Stricture rate <input type="checkbox"/> Any stone recurrence noted in study period <input type="checkbox"/> Any metabolic screening and changes noted <input type="checkbox"/> Use of any preventative medications (e.g., potassium citrate etc.) <input type="checkbox"/> Cost of the procedure (including ancillary equipment used) <input type="checkbox"/> Long-term complications <input type="checkbox"/> Quality of life assessment and changes |

URS = Ureteroscopy

PCNL = Percutaneous nephrolithotomy

MET = Medical expulsive therapy

SFR = Stone free rate

Fig. 1 (continued)

With these issues in mind, we created the Adult Ureteroscopy (A-URS) checklist to complement the P-URS [3]. The development process mirrored the methods used for the

P-URS, including a literature review (P.J.-J. and B.K.S.) and the development of a list of core items. The draft version was critiqued by all of the authors and then revised accord-

ingly. This process was repeated a total of four times until consensus was finally achieved. Before its development, the team decided that the general layout for the A-URS would match that of the P-URS. While many of the points included in the final version may seem obvious at first glance, a review of the literature on adult URS revealed that such information is often missing.

The final version is shown in [Figure 1](#). Key items deemed by the authors to be of high priority are highlighted in bold font. There are five main sections, which cover information on the following areas: study details, and preoperative, operative, postoperative, and long-term data.

Section 1 (Study details) covers generic information relevant to the study such as the aim, design, setting (eg, academic or community and annual case volume), as well as the study selection criteria and indication for the surgery.

Section 2 (Preoperative data) includes details for the operating team (eg, experience) and the number of surgeons involved in the study. This has relevance as it helps in differentiating when a series has been performed by single expert surgeon or whether a group that includes residents was involved. Information on patient demographics is also included here (eg, preoperative urine culture). Another important area covered is preoperative stone status, including stone dimensions, multiplicity, and location, as well as imaging modality. Maximal stone diameter is still the most common measure of size, but stone volume is increasingly being used.

Section 3 contains operative information. Key areas include information relating to the energy source (laser type, power output, and start-up settings). Intraoperative complications can occur, and tools exist that can aid classification at the time of reporting, such as ClassIntra [7]. In cases in which ureteral mucosal trauma has occurred, this can be graded, for example, using the Traxer and Thomas system [8]. The exit strategy should be made clear, including use of any modifications such as a stent on a string.

Section 4 covers postoperative information and follow-up. Consensus is lacking on the ideal approach for reporting stone-free status. From a practical perspective, the inclusion of at least a zero-fragment definition allows for a useful baseline when comparing study results. Use of noncontrast computed tomography allows for more accurate assessment of the residual stone burden and is preferable if available. As well as use of a classification tool for postoperative complications, we recommend recording of further details, rather than only reporting the percentage breakdown for major and minor adverse events, as is often observed. This should also include the time period over which complications were recorded (eg, 30 d).

Section 5 covers long-term information as well as more general items that are valuable for analysis, if available. With this in mind, it is fully appreciated that when undertaking retrospective data collection, not all items can be gathered at a later date; for example, access to prescription records or patients treated out of the area with loss to longer-term follow-up may not be possible.

While there is clear overlap with the P-URS, there are a number of subtle but important differences. In the paediatric setting, for example, further breakdown of the sample

by age group is suggested (eg, infants, children, prepuberty, adolescents) and/or weight. Another difference is the highlighting any additional intraoperative radiation protection measures in paediatric URS and transparency regarding repeat URS at the time of stent removal, given that this is typically performed under general anaesthesia in the paediatric age group. The A-URS also includes an additional fifth section that covers long-term information such as the stricture rate and quality of life that is not present in the P-URS checklist.

The A-URS checklist is not aimed at being an exhaustive list and authors should also not feel obligated to include every item. The main goal is to provide a practical supplement that authors and reviewers can use to provide more robust studies and reports in the setting of URS and stone lithotripsy.

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Study concept and design: Juliebø-Jones, Somani, Ulvik, Beisland.

Acquisition of data: Juliebø-Jones, Somani, Ulvik, Beisland.

Analysis and interpretation of data: Juliebø-Jones, Somani, Ulvik, Beisland.

Drafting of the manuscript: Juliebø-Jones, Somani, Ulvik, Beisland.

Critical revision of the manuscript for important intellectual content: Juliebø-Jones, Somani, Ulvik, Beisland.

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