Management Trends and Outcomes of Patients Undergoing Radical Cystectomy for Urothelial Carcinoma of the Bladder: Evolution of the University of Southern California Experience over 3,347 Cases

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Study Need and Importance: Perioperative and surgical management for invasive urothelial carcinoma of the bladder (UCB) has witnessed several advances over the past decades, including use of neoadjuvant chemotherapy (NAC), enhanced postoperative recovery pathways and robot-assisted radical cystectomy (RC). A 2001 study at our center highlighted the importance of surgical management for aggressive UCB. We herein update our experience and post-RC outcomes against the evolutionary background of developments in management patterns at a high-volume academic center.

What We Found: The contemporary cohort in this study doubles the size of what was previously reported from our center (see figure). Patients undergoing RC are now significantly older and with more comorbidities than in the past. Recent years have witnessed decreased postoperative hospital stay due to enhanced recovery pathways and increased use of robot-assisted RC. Despite maintaining a consistent lymphadenectomy template, lymph node submission as anatomical packets resulted in increased nodal yield. The last decade demonstrated a higher adoption of NAC with concomitant decrease in adjuvant chemotherapy use, resulting in improved complete response (ypT0N0M0) rates. While median overall survival following RC improved with time, recurrence-free outcomes have remained stable. However, patients with extravesical and/or nodal disease following NAC had worse outcomes compared with those who directly underwent RC.

Limitations: These include the study’s retrospective nature, and absence of patient performance metrics and comorbidities. This analysis also predates the increasing use of checkpoint inhibitors for UCB management.

Interpretation for Patient Care: Adherence to oncologically sound surgical principles can result in consistent outcomes for aggressive UCB. Refinements in perioperative management and operative techniques have resulted in improved postoperative experience. While a certain subset of UCB patients can derive great benefit from NAC, it is important to reliably identify those who may not respond to conventional NAC and therefore require other therapeutic interventions or early RC.
Management Trends and Outcomes of Patients Undergoing Radical Cystectomy for Urothelial Carcinoma of the Bladder: Evolution of the University of Southern California Experience over 3,347 Cases

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Purpose: There are conflicting reports on outcome trends following radical cystectomy (RC) for bladder cancer.

Materials and Methods: Evolution of modern bladder cancer management and its impact on outcomes was analyzed using a longitudinal cohort of 3,347 patients who underwent RC at an academic center between 1971 and 2018. Outcomes included recurrence-free survival (RFS) and overall survival (OS). Associations were assessed using univariable and multivariable models.

Results: In all, 70.9% of cases underwent open RC in the last decade, although trend for robot-assisted RC rose since 2009. While lymphadenectomy template remained consistent, nodal submission changed to anatomical packets in 2002 with increase in yield (p < 0.001). Neoadjuvant chemotherapy (NAC) use increased with time with concomitant decrease in adjuvant chemotherapy; this was notable in the last decade (p < 0.001) and coincided with improved pT0N0M0 rate (p = 0.013). Median 5-year RFS and OS probabilities were 65% and 55%, respectively. Advanced stage, NAC, delay to RC, lymphovascular invasion and positive margins were associated with worse RFS (all, multivariable p < 0.001). RFS remained stable over time (p = 0.73) but OS improved (5-year probability, 1990–1999 51%, 2010–2018 62%; p = 0.019). Among patients with extravesical and/or node-positive disease, those who received NAC had worse outcomes than those who directly underwent RC (p ≤ 0.001).

Conclusions: Despite perioperative and surgical advances, and improved pT0N0M0 rates, there has been no overall change in RFS trend following RC, although OS rates have improved. While patients who are downstaged with NAC derive great benefit, our real-world experience highlights the importance of preemptively identifying NAC nonresponders who may have worse post-RC outcomes.
Radical cystectomy (RC) is considered the gold-standard treatment for high-risk invasive urothelial carcinoma of the bladder (UCB). Population-based studies report conflicting UCB outcome trends over time, although these investigations do not differentiate between patients undergoing RC versus bladder-preserving therapies for less aggressive disease. Most studies concur that RC performed at high-volume and academic centers are associated with better outcomes.

Advances in perioperative and surgical UCB management including enhanced recovery after surgery (ERAS) pathways and robot-assisted surgery have improved patient experience while maintaining oncological efficacy. Neoadjuvant chemotherapy (NAC) has been associated with modest survival benefit. However, pathological stage remains the most important predictor of outcomes following RC. This study examined clinicopathological and outcome trends of patients undergoing RC for UCB at the University of Southern California following a consistent surgical philosophy in the face of changing management paradigms over time. The aim was to juxtapose historical and contemporary patient groups to evaluate how temporal evolution of high-risk UCB management has impacted outcomes following RC at a high-volume academic center.

**MATERIALS AND METHODS**

**Patient Population and Management**

A total of 3,957 subjects who underwent RC between 1971 and 2018 were identified through a prospectively maintained database (IRB No. HS018014). Study criteria, preoperative evaluation and management are detailed in the supplementary methods (https://www.jurology.com). RC and urinary diversion were performed by open or robot-assisted approach per patient and surgeon preference. Per institutional standard, all eligible patients underwent extended pelvic lymphadenectomy. Since 2012, patients were started on ERAS pathway. Pathological stage was based on tumor-node-metastasis system at RC (supplementary table 1 and supplementary methods, https://www.jurology.com). Postoperative followup was at 4-month intervals in year 1, 6-month intervals in year 2, and annually thereafter with laboratory and imaging studies as indicated.

**Data Analysis**

Associations between clinicopathological characteristics and outcomes were assessed using univariable and multivariable models (supplementary methods, https://www.jurology.com). Trends were determined annually, or between decades or eras. Directed acyclic graphs were used to identify confounding variables. Oncological outcomes included recurrence-free survival (RFS) and overall survival (OS).

**RESULTS**

**Patient Characteristics and Management Trends**

A total of 3,347 patients underwent RC for UCB and met study criteria for final analysis. Population included 2,685 (80.2%) males, with median age of 68 (IQR, 61–75) years (supplementary table 2, https://www.jurology.com). Median age and proportion of non-Caucasians treated increased over time (p < 0.001; supplementary table 2, https://www.jurology.com; fig. 1). In all, 43.4% of all males had histological evidence of concomitant prostatic adenocarcinoma in their cystoprostatectomy specimen; 58 (1.7%) patients experienced perioperative mortality, ie death, within 30 days of surgery or prior to discharge, whichever was later.

Intravesical therapy and NAC administration increased over time, with concomitant decrease in adjuvant chemotherapy (AC); this was especially notable in last decade (all, p < 0.001; fig. 2 and supplementary table 2, https://www.jurology.com). Cisplatin-based combinations have been the mainstay for NAC (88% patients) and AC (84.4% patients) regimens. Associations between patient and management characteristics are summarized in supplementary figure 1 (https://www.jurology.com). Higher proportions of patients receiving NAC and AC were younger than those who did not receive these treatments (p < 0.005). Median time from diagnosis to RC was 6.2 (IQR 4.6–13.6) months for patients receiving NAC, and 2.7 (IQR 1.3–14) months for those directly undergoing surgery (p < 0.001). NAC administration was not associated with overall surgical margin status (p=0.46); 50.1% and 20.2% of patients receiving NAC achieved tumor downstaging and complete pathological response (pT0N0M0), respectively. The corresponding proportions for patients directly undergoing RC were 27% and 8.7%, respectively (both p < 0.001).

There has been an increasing trend for robot-assisted RC since 2009, although open approach was still used in 70.9% of patients in the last decade (p < 0.001; fig. 2 and supplementary table 2, https://www.jurology.com); 65.8% of patients underwent intracorporeal ileal conduit urinary diversion during robot-assisted RC and 59.6% of patients underwent orthotopic neobladder urinary diversion during open RC (p < 0.001). While there was a trend towards decreasing proportion of ileal conduit diversions and increasing proportion of orthotopic neobladders until 2009, the last decade witnessed an increase in ileal conduits associated with higher percentages of older patients with more comorbidities (p < 0.001). While our extended pelvic lymphadenectomy template has remained consistent
since the 1980s, specimen submission changed from en bloc to anatomically defined nodal packets in 2002 with consequent increase in nodal yield (p < 0.001). Proportion of patients with pT0N0M0 disease increased in the last decade (p = 0.013), with no significant trend differences in clinical stage, variant histology or tumor upstaging rates (supplementary table 2, https://www.jurology.com). Median hospital stay decreased to 5 (IQR 4–7) days since implementation of ERAS in 2012 from 9 (IQR 8–12) days (p < 0.001).

Associations and Trends with Outcomes
Median followup was 10.1 years (range, 3 months to 40.4 years), during which 1,038 (31%) patients recurred and 1,900 (56.8%) patients died. Median±SE 5-year RFS and OS probabilities were 65±1% and 55±1%, respectively (fig. 3).

Univariable associations of patient characteristics with outcomes are detailed in supplementary table 3 (https://www.jurology.com). Advanced pathological stage was associated with worse RFS and OS (both p < 0.001; fig. 4). This overall trend persisted even after stratification by decade of surgery (table 1). NAC and AC were univariably associated with worse RFS (both p < 0.001), consistent with them being administered for more advanced disease (supplementary fig. 1, https://www.jurology.com). Longer delay to RC, variant histology, lymphovascular invasion, tumor upstaging, and positive surgical margins were also associated with worse outcomes (all p < 0.001).
Relevant univariably prognostic variables were included in multivariable models to estimate their independent relationships with outcomes (table 2). This confirmed that advanced pathological stage was associated with worse outcomes (RFS and OS, p < 0.001). Other independent RFS predictors included NAC and AC administration (both p < 0.001). Longer delay to RC, lymphovascular invasion, tumor upstaging, and positive margins remained independently associated with worse outcomes (all p ≤ 0.030). Given the interplay of various demographic and clinicopathological characteristics in determining outcomes, an exploratory variable analysis using directed acyclic graph was performed to outline causal relationships and eliminate confounders with the hypothesis that pathological stage is a primary outcome predictor following RC (supplementary fig. 2, https://www.jurology.com). This identified biasing paths that could be controlled by adjusting for age, comorbidity measured by American Society of Anesthesiologists® score, NAC administration and lymphovascular invasion (supplementary table 4, https://www.jurology.com), which would be sufficient to estimate the total effect of pathological stage on outcomes. The resulting multivariable model confirmed the independent associations of pathological stage (p < 0.001) and the other variables with outcomes (supplementary table 5, https://www.jurology.com). Despite causal relationship-based adjustment, NAC administration remained associated with worse RFS and OS (both, p < 0.001).

There was no change in RFS trend across decades (p=0.73; supplementary table 2, https://www.jurology.com) or annually (p=0.94; fig. 5, and...
supplementary fig. 3, https://www.jurology.com). For example, median±SE 5-year RFS probability was 64%±2% during 1990–1999 and 66%±2% during 2010–2018. However, OS trend improved across decades (p<0.019) and annually (p=0.009). For instance, median±SE 5-year OS probability improved from 51%±2% during 1990–1999 to 62%±2% during 2010–2018. When stratified by pathological stage, the trend in improved OS was significantly notable in patients with pT2-4N0M0 disease at RC (table 1).

Use of NAC
Trends in proportion of patients receiving NAC and associated pathological stages were analyzed (fig. 6, A). NAC administration increased across eras with correspondingly increased pT0N0M0 rate (supplementary table 2, https://www.jurology.com). Among patients who did not receive NAC, advanced pathological stage was associated with worse RFS and OS (overall p<0.001; supplementary table 6, https://www.jurology.com; stratified p<0.001; fig. 6, B,C). While this association with pathological stage was also noted among NAC recipients (overall p<0.001, supplementary table 6, https://www.jurology.com), stratified analysis between pT0N0M0 and organ-confined disease showed no difference in RFS (p=0.42) or OS (p=0.87; fig. 6, D and E). However, RFS and OS for these disease stages were better than for patients with extravesical and/or nodal metastatic disease who received NAC (stratified p<0.001). Among patients with extravesical and/or nodal metastatic disease at RC, those who previously received NAC had worse RFS (log-rank p<0.001) and OS (log-rank p=0.001) than those who directly underwent RC (supplementary table 6, https://www.jurology.com; data not shown).

Among patients with ≥cT2 disease, those who were upstaged following NAC had significantly worse RFS and OS (both, p<0.001) compared with those who directly underwent RC (table 3). However, those who were downstaged following NAC had a slightly worse RFS (p=0.046) and no difference in OS (p=0.99) compared with those who directly underwent RC.

DISCUSSION
A prior report of our institutional RC experience highlighted the importance of surgical management for invasive UCB. However, it predated important changes in perioperative and surgical management including NAC use, lymph node packeting, ERAS, and robot-assisted RC. This study presents our long-term experience using a meticulously annotated population of 3,347 patients to compare historical and contemporary outcomes following RC in light of evolving changes to UCB management. Through the decades, our patients have trended to being older and with more comorbidities. NAC use has increased, peaking at 36.8% in 2017, with concomitantly decreased AC use that is consistent with national trends; 29.2% of patients overall (40.8% of patients with ≥cT2 disease) received NAC in the last decade compared with the 22.3% national average, which translated to an improved 23.7% overall pT0N0M0 rate in this subpopulation in the same timeframe. Conformity to ERAS resulted in shorter inpatient stay. Despite these advances, there was no change in RFS over time, although OS showed a trend towards improvement. While patients with organ-confined disease have excellent 5-year RFS and OS probabilities after RC (80% and 71%, respectively), corresponding rates in patients with nodal-metastatic disease remain dismal (30% and 25%, respectively). This reflects “real-world” trends and outcomes following RC at a high-volume academic center with a tradition of pioneering UCB management. It underscores the value of aggressive surgical treatment that incorporates advances in operative and perioperative management while emphasizing the need for innovations that improve disease-specific outcomes.

Prior studies have reported an association between improved surgical outcomes and higher number of dissected lymph nodes. However, our data indicate that, notwithstanding a consistent template, nodal counts can vary based on sample submission methodology. Our nodal yield increased with concomitant decrease in nodal density without any significant RFS change since specimen submission was changed in 2002 from en bloc to anatomically defined packets. Others have emphasized the importance of a thorough lymphadenectomy over...
nodal counts for achieving optimal oncologic outcomes.\textsuperscript{16} Our RFS and OS are comparable to the series from Ulm, Germany, where patients underwent super-extended lymphadenectomy since 2001 with a mean of 18 resected nodes.\textsuperscript{9} We believe that our adherence to a meticulous extended lymphadenectomy template using open and robot-assisted approaches has resulted in high nodal yields and putatively curative resections. The first phase 3 trial to address the extent of node dissection in RC was unable to provide conclusive evidence of oncologic benefit for extended lymphadenectomy over

![Comparison of clinical outcomes of patients with bladder cancer stratified by pathological stage.](image)

**Figure 4.** Comparison of clinical outcomes of patients with bladder cancer stratified by pathological stage. Kaplan-Meier curves show recurrence-free (A) and overall survival probabilities (B) of patients with pT0N0M0 (green), nonmuscle-invasive (magenta), pT2N0M0 (aqua), extravesical (orange), and nodal metastatic (purple) disease. Overall p value calculated by log rank test.
Table 1. Univariable associations of pathological stage with clinical outcomes in patients undergoing radical cystectomy for bladder cancer, stratified across decades

| Outcome by Pathological Stage | 1971-1979 | 1980-1989 | 1990-1999 | 2000-2009 | 2010-2018 |
|-------------------------------|-----------|-----------|-----------|-----------|-----------|
| % Probability±SE              | 3-yr      | 5-yr      | 3-yr      | 5-yr      | 3-yr      | 5-yr      |
| RFS                           |           |           |           |           |           |           |
| pT0NO0M0                      | 93±6      | 93±6      | 99±6      | 95±3      | 93±4      | 95±2      |
| Nonmuscle-invasive pT2N0M0     | 81±6      | 79±6      | 84±3      | 81±3      | 87±3      | 86±2      |
| Extravesical pT0N0M0           | 84±9      | 78±10     | 82±4      | 80±4      | 81±4      | 76±4      |
| Nodal metastatic              | 32±13     | 32±13     | 55±5      | 50±4      | 54±5      | 58±4      |
| OS                            | 29±8      | 23±7      | 32±5      | 30±5      | 34±4      | 35±4      |
| Overall log rank p <0.001 for each clinical outcome in each decade. |
| Risk of Recurring              |           |           |           |           |           |           |
| HR (95% CI)                    |           |           |           |           |           |           |
| Age (yrs):                     |           |           |           |           |           |           |
| <70                           | 1.00 (reference) |          | 1.00 (reference) |          |<0.001     |
| >70                           | 0.93 (0.82—1.05) |        | 1.66 (1.50—1.83) |        |           |
| NAC                           |           |           |           |           |           |           |
| Not administered              | 1.00 (reference) |        | 1.00 (reference) |        |0.072      |
| Administered                  | 1.56 (1.31—1.86) |       | 1.15 (0.99—1.34) |       |           |
| Diagnosis to cystectomy (days): |           |           |           |           |           |           |
| <120                          | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| >120                          | 1.30 (1.14—1.48) |      | 1.20 (1.09—1.32) |      |           |
| Surgical approach:            |           |           |           |           |           |           |
| Open                          | 1.00 (reference) |        | 1.00 (reference) |        |0.21       |
| Robot-assisted                | 1.12 (0.90—1.38) |     | 0.87 (0.70—1.08) |     |           |
| Pathological stage:           |           |           |           |           |           |           |
| pT0N0M0                       | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| Nonmuscle-invasive pT2N0M0     | 1.69 (1.16—2.47) |      | 1.12 (0.91—1.37) |      |           |
| Extravesical                  | 2.11 (1.42—3.13) |      | 1.32 (1.06—1.65) |      |           |
| Nodal metastatic              | 4.99 (3.38—7.36) |      | 2.27 (1.80—2.86) |      |           |
| Tumor upstaging               | 9.14 (6.19—13.48) |      | 3.57 (2.39—4.51) |      |           |
| Risk of Overall Mortality     |           |           |           |           |           |           |
| HR (95% CI)                    |           |           |           |           |           |           |
| Age (yrs):                     |           |           |           |           |           |           |
| <70                           | 1.00 (reference) |        | 1.00 (reference) |        |0.002      |
| >70                           | 1.18 (1.02—1.37) |      | 1.20 (1.07—1.35) |      |           |
| NAC                           |           |           |           |           |           |           |
| Not administered              | 1.00 (reference) |        | 1.00 (reference) |        |           |
| Administered                  | 1.11 (0.96—1.28) |      | 0.97 (0.87—1.09) |      |           |
| Variant histology:            |           |           |           |           |           |           |
| Absent                        | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| Present                       | 1.11 (0.96—1.28) |      | 0.97 (0.87—1.09) |      |           |
| Lymphovascular invasion:      |           |           |           |           |           |           |
| Absent                        | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| Present                       | 1.52 (1.31—1.75) |      | 1.42 (1.27—1.59) |      |           |
| All surgical margin status:   |           |           |           |           |           |           |
| Neg                           | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| Pos                           | 1.79 (1.49—2.18) |      | 1.47 (1.26—1.72) |      |           |
| AC                            |           |           |           |           |           |           |
| Administered                  | 1.00 (reference) |        | 1.00 (reference) |        |<0.001     |
| Not administered              | 1.53 (1.31—1.80) |      | 1.86 (1.63—2.13) |      |           |

* p value based on Cox proportional hazards model.

Table 2. Multivariable associations of patient, disease and management characteristics with clinical outcomes in patients undergoing radical cystectomy for bladder cancer

standard dissection, although it was not a non-inferiority design and likely underpowered to discern small but clinically relevant differences. Results of the SWOG S1011 trial that randomized patients with muscle-invasive disease to standard versus extended lymphadenectomy are pending.

In contrast to national trends, majority of patients at our institution underwent orthotopic ileal...
neobladder as the modality of urinary diversion.\textsuperscript{18} This is also oncologically safe and feasible in patients with prostatic stromal invasion and prior pelvic irradiation.\textsuperscript{19,20} Individual choice of diversion is ultimately a shared decision based on preoperative counseling, patient and disease characteristics, lifestyle, priorities, and other intangible patient preferences.

Our findings have some important differences from the recently reported Memorial Sloan Kettering Cancer Center (MSKCC) experience.\textsuperscript{21} Detailed pathological staging for the MSKCC population is unavailable, but their overall nodal metastasis rate of 20\% suggests a comparable cohort composition as ours. While RFS in our cohort did not improve over time, MSKCC reported a significant improvement in 5-year RFS probabilities from 58\% in 1997 to 66\% in 2013. Our corresponding probabilities during those decades remained consistent at 64\% and 66\%, respectively (supplementary table 2, https://www.jurology.com). Indeed, the cumulative 5-year RFS probability across our cohort is 65\%, thereby explaining the perceived lack of change. However, it is unclear whether this represents a plateau in overall benefit with the current armamentarium of surgical and medical innovations. The MSKCC report also predated their institutional implementation of ERAS and robot-assisted RC. The authors attributed their RFS improvement to its temporal association with increasing NAC use but did not present any multivariable analyses. Our multivariable analyses employing classical and directed acyclic graphical approaches highlight the independent prognostic roles of age, comorbidities, pathological stage, NAC administration and lymphovascular invasion. Among patients who received NAC, our ypT0N0M0 rate of 21.7\% in the last 2 decades was comparable to MSKCC (22\%) and other

\section*{Figure 5. Annualized trends of clinical outcomes following radical cystectomy for bladder cancer. Five-year probability of recurrence-free (A) and overall survival (B) for patients (solid line) with corresponding standard error estimates (shaded area). Estimates determined for patients undergoing cystectomy until 2013 to allow for adequate followup. Univariable log rank test for trend p=0.94 and 0.009, respectively.}

\section*{Figure 6. NAC administration and association with clinical outcomes. A, flow diagram depicts the relative proportion of patients in each 6-year era who received (purple) or did not receive (orange) NAC, and corresponding proportions who were found to have pT0N0M0 (green), organ-confined (blue), and extravascular and/or nodal metastatic (red) disease on radical cystectomy. Thickness of each colored curve corresponds to relative proportion for respective originating node. Kaplan-Meier curves show probabilities of recurrence-free (B, D), and overall survival (C, E) among patients who did not (B, C) and did (D, E) receive NAC, when stratified by pathological stage. Log rank p values compared across strata shown.}
### Table 3. Univariable associations of NAC administration with clinical outcomes in patients with ≥cT2 bladder cancer undergoing radical cystectomy, stratified by pathological tumor staging status

| Pathological Tumor Staging Status | No. (%) | 3-yr RFS Probability±SE (%) | 5-yr RFS Probability±SE (%) | p Value* | 3-yr OS Probability±SE (%) | 5-yr OS Probability±SE (%) | p Value* |
|----------------------------------|---------|-----------------------------|-----------------------------|----------|-----------------------------|-----------------------------|----------|
| Downstaged to ≤pT1N0M0 at cystectomy: | | | | | | | |
| NAC not administered | 397 (68.9) | 88±2 | 85±2 | 0.046 | 80±2 | 75±2 | 0.99 |
| NAC administered | 179 (31.1) | 79±3 | 79±3 | <0.001 | 79±3 | 78±4 | <0.001 |
| Upstaged at cystectomy: | | | | | | | |
| NAC not administered | 713 (87.0) | 45±2 | 44±2 | <0.001 | 38±2 | 32±2 | <0.001 |
| NAC administered | 107 (13.0) | 28±5 | 28±5 | | 19±5 | 12±4 | |

*p Value based on log rank test.

international academic centers (22.7%), and higher than the overall national average (10.6%) during the corresponding time frame.21–23 Our findings indicate that there was discernible benefit for patients receiving NAC who achieved pT0N0M0 or organ-confined disease at RC with no significant outcome differences between these subgroups. In contradistinction, those with extravesical and/or nodal metastatic disease after NAC experienced delay to definitive surgery and worse outcomes than those who directly underwent RC. While these findings should be interpreted cautiously given the study’s retrospective nature, it is clear that current clinical tools for predicting pathological stage and NAC response are relatively imprecise.24 This highlights the need to validate and incorporate novel biomarkers that identify patients who may likely not respond to traditional NAC, and may therefore benefit more from novel agents or expedited surgery.25–27 The need to achieve remission following definitive surgery is crucial as post-cystectomy recurrences are associated with dismal prognosis despite salvage therapy.28

This study has some limitations that warrant consideration. Precise metrics of performance status and comorbidities were not accurately recorded on all patients given the study’s retrospective nature and were therefore excluded from analysis. Subcategories of variant histology were not examined in detail as they comprised a small minority of the population and this was beyond the analytic scope; outcomes of these subgroups have been characterized previously.29,30 This population also predates the increasing use of immune checkpoint inhibitors and other targeted therapies for advanced UCB management. The strength of this study rests on the well-curated clinical information on patients with UCB undergoing RC from a prospectively maintained database. Standards for clinical decision making, indications for RC, surgical intervention and technical fundamentals, and pathological evaluation have remained consistent throughout the study period. Operative principles were meticulously followed by a group of high-volume surgeons, and all relevant clinicopathological annotations were standardized. Granular analysis of the population on a decade-by-decade and annual basis allowed for combined comparison of historical and contemporary data to map the evolution of surgical management of UCB in the context of medical, technical and perioperative advances.

**CONCLUSIONS**

This study presents a homogeneous RC series with long followup where consistency of surgical philosophy and standardized reporting permitted focused comparisons. Taken together, these data suggest that adherence to oncologically sound surgical principles can result in consistent outcomes following RC for UCB. While there has been an improvement in OS, this trend has not been recapitulated with RFS over time. Our real-world evolutionary experience with NAC suggests that a certain subset of UCB patients can derive great benefit. The challenge is to build reliable tools that accurately identify this subgroup, while developing novel therapeutic strategies for those who may not respond to conventional NAC in order to augment the oncologic benefit of radical surgery.

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**EDITORIAL COMMENT**

Mitra et al provide an update to a large and mature cystectomy series 2 decades after Stein and colleagues first detailed what have become benchmarks for stage-stratified recurrence risk and survival after cystectomy (reference 8 in article). While the 2001 publication has become the standard-bearer for outcome probabilities and prognosis following cystectomy, results from this update are less instructive. The authors provide an abounding array of analyses that seem at once too broad and unfocused. In brief, they report decreased length of stay influenced by ERAS protocols, increases in lymph node counts following changes in specimen procurement, and modest uptake in use of NAC and robotic assisted approaches at their institution. Despite implementing these changes, cancer recurrence outcomes remain stubbornly stagnant in this decades-long series. A closer look at the data offers some clues for this disappointing lack of progress. Although the use of NAC at the University of Southern California has increased, its rate still hovers around only 30%. At many other centers, including our own, NAC usage exceeds 50% (reference 21 in article). Without complete knowledge of the clinical staging, the insinuations by the authors that patients with advanced disease suffered worse outcomes following NAC compared to those who directly underwent cystectomy should be interpreted judiciously. While this alone may not fully account for the lack of improvement in oncologic outcomes, it points broadly to room for improvement for all physicians and surgeons who manage bladder cancer. While emerging advances in molecular profiling and immunoncology will undoubtedly unlock the outcomes stalemate,
equally important is the adherence to best practice guidelines and efficient implementation of strategies to maximize treatment efficacy. We congratulate the investigators on assembling such an exhaustive surgical series and commend them on their transparent reporting of outcomes. However, it is time for us to move beyond the plateau in outcomes achieved with surgical treatment.

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REPLY BY AUTHORS

We appreciate the comments on the update of our RC experience for high-risk UCB. This study was designed to investigate 1) evolution of management patterns and their impact, if any, on outcomes, 2) long-term outcome trends, and 3) real-world experience with perioperative chemotherapy use.

We offer detailed and transparent reporting of one of the world’s largest RC series. We understand the concerns raised when evaluating *prima facie* evidence regarding NAC use and response. However, we caution readers from drawing premature conclusions without contextually examining the data. NAC usage should be assessed against the appropriate denominator. NAC rates increased in the last decade among patients with ≥cT2 disease (p <0.001, supplementary table 2 in article), exceeding 50% in the last few years in this subset. With increasing NAC use and rising nodal yield, another study reported 5-year RFS improvement from 58% (1997) to 66% (2013; reference 21 in article). Following a consistent surgical philosophy, our overall 5-year RFS of 65% has not changed significantly over decades, although OS has improved. We categorically are not suggesting that all patients with advanced disease have worse outcomes following NAC; indeed, our increasing NAC rates are evidence to the contrary. However, patients with ≥cT2 disease who do not respond to NAC perform worse than those who directly undergo RC. This underscores the pressing need for personalized approaches and accurate determinants of NAC response, and forms the basis for biomarker-based investigations that are currently underway. We remain optimistic that addition of novel targeted therapeutics to the armamentarium can move the needle in the future.