INTRODUCTION

Prostate cancer (PCa) remains the most common non-cutaneous malignancy and second most common cause of cancer-related mortality among American men in the 21st century.[1] As PCa management pathways continue to evolve, the process of treatment selection can become a challenging endeavor. Often, notwithstanding a shared decision-making approach, treatment is biased toward the consultant’s area of expertise.[2] Alternatively, PCa multidisciplinary cancer clinic (MDC) comprehensively addresses all applicable treatment options and can occur in the setting of separate risk/benefit discussions with PCa experts in the fields of urologic surgery,
medical and radiation oncology following a review of imaging and histopathology by participating radiologists and genitourinary pathologists.\textsuperscript{[3-6]} For some institutions, the use of MDC for PCa management has been a critical part of urologic practice for decades and a mandated modality for the evaluation of newly diagnosed cancer patients.\textsuperscript{[7,8]}

Despite the advantages of MDC for PCa management, patient attendance and the decision to subsequently undergo treatment at the MDC center is variable.\textsuperscript{[9]} However, a distinct benefit to the patient is the multidisciplinary expert review of preexisting pathologic and radiologic data that can result in reclassification of disease, alteration in the treatment algorithm,\textsuperscript{[4]} as well as incur the possibility of a survival benefit.\textsuperscript{[7]} Thus, as a consequence of vetting PCa treatment options within the context of MDC, the subsequent utilization of diagnostic and therapeutic resources is executed in an efficient and guideline adherent manner.

Given the inherent complexity, demand for time and physician resources the institutionalization of MDC is not ubiquitous. Herein, we present a query and analysis of physician characteristics and opinion regarding MDC, and discuss potential predictors of MDC utilization for the evaluation and treatment of PCa.

MATERIAL AND METHODS

Survey

A 14-item questionnaire was created to collect baseline characteristics and practice pattern data among the Society of Urologic Oncology (SUO) and the Endourological Society (ES) members regarding MDC for the management of PCa. We did not include a definition of MDC in the survey, thus the specific MDC setting was unknown. Urologist demographic data included age, sex, practice type, fellowship training, years in practice, and number of PCa patients seen in 1 month’s time. Response options included the selection of a single response, select all that apply, or freehand responses depending on the query. A complete version of the survey is included as a Supplementary Appendix.

Study design

Between January and February 2018, the 14-item questionnaire was disseminated on two separate occasions through an email containing a hyperlink to the web-based survey platform (SurveyMonkey\textsuperscript{®}, Palo Alto, CA, USA) to members of both the SUO and ES. Approximately 2–3 weeks’ time elapsed between survey disseminations. No incentive was offered for completion of the survey. We estimate approximately 6000–7000 total members of the SUO and ES collectively received the request to participate in the study. Survey responses were then collected and analyzed anonymously. For the purposes of discussion of our analysis, we utilized PCa disease risk categorization defined by the National Comprehensive Cancer Network (NCCN).\textsuperscript{[9]} The study was determined to be exempt from review by Institutional Review Board.

Statistical analysis

Statistical analysis was performed using R (R Core team, 2017). Mann–Whitney U-test was used to compare distribution of continuous variables. Pearson Chi-square test and Fisher’s exact tests were used to compare proportions of categorical variables. Multivariate proportional odds models were fitted to identify predictors for urologists’ perception of benefit of MDC in treating PCa patients. The predictors studied were age, type of practice (academic vs. non-academic), years in practice, urologic oncology fellowship training, and number of new PCa patients seen per month. Statistical significance was defined as $P < 0.05$.

RESULTS

Respondents baseline characteristics are shown in Table 1. A total of 211 responses were obtained: 87 respondents from the SUO (41%) and 124 (59%) respondents from the ES. Thus, the estimated response rate was approximately 3%. Respondents were most commonly academic (61.4%), male urologists (97.6%) with median age of 48.5 years, fellowship training in urologic oncology (50%), and greater than 20 years of practice (40.2%). One hundred and seven respondents (51%) reported that they participated in MDC.

| Table 1: Baseline respondent characteristics ($n = 208$). |
|----------------------------------------------------------|
| Median age (IQR) | 48.5 (18.5) |
| No. of male (%) | 203 (96.7) |
| Practice context (%) | |
| Academic | 129 (61.1) |
| Non-academic | 82 (38.8) |
| Years in practice | |
| 0–5 | 31 (14.7) |
| 6–10 | 35 (16.6) |
| 11–20 | 60 (28.4) |
| >20 | 85 (40.2) |
| Urologic oncology fellowship training (%) | |
| Yes | 105 (50.0) |
| No | 105 (50.0) |
| No. of patients seen with newly diagnosed prostate cancer monthly (%) | |
| None | 4 (1.9) |
| 1–5 | 68 (32.3) |
| 6–10 | 68 (32.3) |
| 11–20 | 46 (21.8) |
| >20 | 25 (11.8) |
| IQR: Interquartile range |
of which 53% were fellowship trained. The majority of reported MDCs existed in an academic institution (65.3%).

Survey responses regarding patient characteristics and the logistics of MDC are depicted in Table 2. MDCs most commonly existed in an academic center (65.3%) and were managed by a similar distribution of urologists, radiation, and medical oncologists. Most respondents agreed that MDC was “very beneficial” for PCa treatment and research [Figure 1]. Criticisms against use of MDC included lack of infrastructure (65.4%) and time consumption (32.7%) [Figure 2].

Table 3 shows both univariate and multivariate analyses to identify predictors for respondents to agree that MDC benefits management of PCa. On multivariate analysis years in practice tended, the likelihood of positively viewing MDC, that is, the longer a respondent was in clinical practice, the less beneficial MDCs were reported (11–20 years in practice OR 0.32, 95% CI 0.113–0.882, P < 0.028; >20 years in practice OR 0.14, 95% CI 0.031–0.616, P < 0.009). Specifically, respondents with 11–20 years and >20 years in practice were 68% and 86% less likely to find MDC beneficial for the management of PCa patients. Although respondent age approached statistical significance, no other variables on multivariate analysis were predictive of reporting benefit of MDC.

### DISCUSSION

MDC evaluation for men with newly diagnosed PCa represents a modality of management unlike that achievable within a single area of expertise. The premise of MDC lies within the comprehensive review of clinicopathologic and radiologic data such that patient management is deliberated and the most appropriate treatment options, albeit on the basis of national guideline recommendations, are presented to the patient. We sought contemporary opinion of MDC for PCa management from the urologist’s perspective. Our results suggest a positive view of MDC and notable barriers for implementation. We found that the modern PCa MDC was reported to occur within the context of an academic center and is staffed by experienced urologists with fellowship training. The majority of respondents supported MDC with respect to the benefits for patient care and research, whereas criticisms of MDC adoption included complex logistics and the resource demand that is required. Importantly, on multivariate analysis years in clinical practice tended to decrease the probability, a provider would find MDC beneficial for the treatment of PCa.

A tenant of MDC is adherence to current guideline-directed practice. To this end, several centers have explored their MDC experience, compared treatment pathways, and outcomes to a matched cohort of men from the Surveillance, Epidemiology, and End Results (SEER) database. The National Cancer Institute retrospectively compared treatment outcome of their MDC patients with SEER patients and reported overall survival approaching 100% at 5 and 10 years for localized PC, as well as significantly improved survival for locally advanced PCa.\(^{[7]}\) Tang et al.\(^{[8]}\) found PCa treatment selection differed than that occurring on a national basis; MDC patients with low-risk disease were more often treated with active surveillance and those with high-risk disease were treated with definitive treatment (e.g., prostatectomy or radiation therapy) compared to matched SEER controls over the period of study. Importantly, specific discrepancies were identified in the treatment of intermediate- and high-risk African-American men compared to Caucasian men, which are well-known phenomena.\(^{[10]}\) Since MDC provides unbiased

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**Table 2: Survey responses for participants in multidisciplinary clinic.**

| Query                                                                 | Respondents* (%) |
|----------------------------------------------------------------------|------------------|
| Where are most referrals to your PCa MDC directed from?             |                  |
| Urology                                                             | 84 (78.5)        |
| Radiation oncology                                                  | 14 (13.1)        |
| Medical oncology                                                    | 3 (2.8)          |
| Primary care provider                                               | 2 (2)            |
| Other provider                                                      | 4 (3.7)          |
| How often do you schedule PCa MDC?                                 |                  |
| Once per month                                                      | 23 (21.5)        |
| Twice per month                                                     | 29 (27.1)        |
| Four times per month                                                | 42 (39.3)        |
| Greater than 4 times per month                                      | 11 (10.3)        |
| Unknown                                                             | 2 (2)            |
| What patient population comprises your PCa MDC†                     |                  |
| Clinically localized low-risk disease (Gleason sum 6)               | 73 (68.2)        |
| Clinically localized intermediate-risk disease (Gleason sum 7)       | 86 (80.4)        |
| Clinically localized high-risk disease (Gleason sum 8–10)           | 89 (83.1)        |
| Active surveillance candidates                                       | 61 (57.0)        |
| Advanced/metastatic prostate cancer                                 | 78 (72.9)        |
| Recurrent prostate cancer                                           | 40 (37.4)        |
| What professionals participate in PCa MDC at your institution?‡      |                  |
| Urologist                                                           | 106 (99.1)       |
| Radiation oncologist                                               | 100 (93.5)       |
| Medical oncologist                                                  | 91 (85)          |
| Radiologist                                                         | 55 (51.4)        |
| Pathologist                                                         | 54 (50.0)        |
| Nurse practitioner                                                  | 50 (46.7)        |
| Nutritionist                                                        | 13 (12.1)        |
| Social worker                                                       | 23 (21.5)        |
| Other                                                               | 7 (6.5)          |

*In 107 respondents. †Respondents asked to select all that apply. PCa: Prostate cancer, MDC: Multidisciplinary clinic.
PCa counseling, it can help mitigate racial disparities seen in treatment pathways.\textsuperscript{11} Similar guideline adherence in appropriately risk stratified PCa treatment selection in the setting of MDC was also identified by Aizer \textit{et al.}, who demonstrated that evaluation at MDC was a significant independent predictor for choosing active surveillance in the setting of low-risk disease.\textsuperscript{12}

Our survey responses found most patients presenting to MDC had at least intermediate-risk or advanced PCa. As such, the impact of MDC evaluation for patients with complex disease is seemingly relevant. Reichard \textit{et al.}'s\textsuperscript{13} comparison of their MDC patients with NCCN high-risk and very high-risk PCa to a matched SEER cohort who underwent primary surgical or radiation-based treatment

### Table 3: Logistic regression analysis; referent how beneficial is multidisciplinary clinic for treating patients with prostate cancer.

| Variable                          | Benefit of MDC for treatment |
|-----------------------------------|------------------------------|
|                                   | Univariate OR (CI) | P-value | Multivariate OR (CI) | P-value |
| Age                               | 0.99 (0.97–1.01) | 0.574 | 1.04 (0.99–1.09) | 0.063 |
| Practice type                     |                             |       |                     |        |
| 0=Non-academic                    |                             |       |                     |        |
| 1=Academic                        | 0.87 (0.50–1.52) | 0.630 | 0.85 (0.45–1.6) | 0.54   |
| Fellowship training\*             |                             |       |                     |        |
| 0=No                              |                             |       |                     |        |
| 1=Yes                             | 0.89 (0.54–1.44) | 0.632 | 0.77 (0.43–1.36) | 0.37   |
| Years in practice                 |                             |       |                     |        |
| 1=0–5                             |                             |       |                     |        |
| 2=6–10                            | 0.98 (0.41–2.35) | 0.969 | 0.74 (0.26–2.04) | 0.561 |
| 3=11–20                           | 0.71 (0.32–1.55) | 0.391 | 0.31 (0.11–0.88) | 0.028 |
| 4≥20                              | 0.59 (0.28–1.24) | 0.168 | 0.13 (0.03–0.61) | 0.001 |
| No. of newly diagnosed PCa seen   |                             |       |                     |        |
| 0=None                            |                             |       |                     |        |
| 1=1–5                             | 0.68 (0.11–4.25) | 0.687 | 0.07 (0.10–4.79) | 0.721 |
| 2=6–10                            | 0.55 (0.09–3.39) | 0.519 | 0.70 (0.10–4.75) | 0.716 |
| 3=11–20                           | 1.07 (0.17–6.79) | 0.941 | 1.24 (0.18–8.65) | 0.826 |
| 4≥20                              | 1.56 (0.23–10.54) | 0.648 | 2.03 (0.27–14.92) | 0.486 |

*Fellowship training in urologic oncology. MDC: Multidisciplinary clinic, OR: Odds ratio, CI: Confidence interval

Figure 1: Distribution of opinion regarding the benefits of multidisciplinary clinic for prostate cancer treatment or research. *For 211 responses.
found a mean 17-month survival advantage for all patient's treated at their MDC. The authors reported that in addition to their radiation treatment protocols, use of neoadjuvant and multimodal treatment algorithms was felt to be the result of appropriately managing the “volume of information processed in a clinical visit” and mitigating “cognitive biases that can affect decision-making.”

Implementation of MDC demands financial resources and a significant time commitment from respective personnel. Our survey respondents reported that most MDCs were at least staffed with a urologist and radiation oncologist and occurred at least more than once per month, which is consistent with the literature.\textsuperscript{[5,12,14,15]} Importantly, nearly 40% of respondents declared a lack of infrastructure as limiting initiation of MDC followed by “insufficient time” and “cost is prohibitive.” Cost and efficiency measures of MDC were studied by De Leso et al.\textsuperscript{[10]} at 52 MDCs across various types of malignancies in the United Kingdom over a 1-month period. Cost analysis accounted for the hourly cost of physicians and ancillary personnel at the MDC meeting, overhead facility cost, and the cost of time spent preparing radiologic and pathologic patient data for review. The authors reported cost estimates between £14,000 and £38,000 ($21,000 to $57,000 based on 2011/2012 years figures) per month depending on the MDC. Interestingly, 42% of patients required greater than 1 MDC meeting, which was most often attributed to lack of acquisition and/or preparation of radiologic and pathologic studies before MDC. Indeed, lack of preparation implicating increased cost of MDC was echoed through interviews regarding protocolization and content of a urologic MDC with member physicians.\textsuperscript{[15]} Criticisms most commonly reported from the study included the lack of dedicated time for MDC preparation, especially for participating radiologists and pathologists, haphazard preparation on the part of the urologist, the lack of a nominated consultant in charge to make executive decisions, a referral pattern rendering patients possibly not meeting MDC inclusion criteria, and the overall time constraint to conduct the MDC. As demonstrated in the abovementioned study, our survey respondents also considered the cost and time commitments as impediments to wider adoption of MDC for PCa.

Although the use of MDC exists outside of PCa management, it is not ubiquitous among all other oncologic specialties in the United States. In the United Kingdom, however, MDCs are a mandated practice for the management of newly diagnosed malignancy.\textsuperscript{[14]} In an effort to improve MDC, Lamb et al.\textsuperscript{[16]} solicited for consensus among the United Kingdom MDC participants across a number of visceral and hematologic malignancies by querying opinions for various aspects of MDC. Their study included >1000 participants and identified consensus for ~85% of queries (113/136) with respect to infrastructure, governance, meeting logistics, and the shared decision-making process of MDC. Therefore, it may be prudent to perform a similar consensus study in the United States across various MDCs to create solutions to our respondents’ concerns that limit MDC adoption.

The results of our multivariate analysis suggest that urologist respondents in practice for longer than 10 years are less likely to find MDC beneficial for PCa management. We found this to be an interesting finding, as fellowship training can impact years in practice and the practice setting but did not significantly affect our queries of MDC. Notwithstanding the limits of our survey to further dissect this finding, we speculate that perhaps urologists, who one can conjecture that after a decade have established themselves in practice, may have created a referral pattern of consultants that would supplement MDC. Moreover, it can be inferred that the decision to establish or participate in an MDC has been already occurred. As previously discussed, the implementation of MDC for cancer care in the United States is not as ubiquitous or mandatory as elsewhere, and therefore, the urologist who is unfamiliar with the logistics and commitments of MDC may in itself be a reason to not participate.

The role of the radiologist in MDC will continue to grow as diagnostic and therapeutic interventions for PCa evolve. The advent of multiparametric magnetic resonance imaging (mpMRI) and the prostate imaging reporting and data system\textsuperscript{[17]} has led to increasing adoption of and advocacy for mpMRI-targeted prostate biopsy. A recent consensus of the literature estimates a negative predictive value close to 90% with over a third of men who undergo pelvic mpMRI avoiding the cost, morbidity, and potential overtreatment associated with prostate biopsy.\textsuperscript{[18]} With respect to treatment, partial or focal gland ablation has become a formidable therapeutic modality highly reliant on the interpretation of pelvic mpMRI. In 2020, a European consensus survey of urologic oncologists (72%) and radiologists (28%) was performed to standardize partial gland ablation management of focal PCa, thus highlighting the importance of a continued multidisciplinary approach to PCa. Moreover, as evident in
recent survey of radiologists who participate in oncologic MDCs, and echoing our survey results from urologists, time constraints and lack of incentive limit the preparation for and participation in MDC. Ultimately, multidisciplinary efforts will be required to resolve these multidisciplinary issues.

Limitations of this study include the inherent subjectivity in a questionnaire-based investigation and the low survey response rate. Importantly, the majority of respondents were academic urologists participating in an academic institution MDC, which introduces sampling bias to our results. Future sampling of contemporary opinion for MDC should target urologists in the private setting, as private urologists comprise the majority of practicing urologists and also include the perspective of participating and non-participating medical and radiation oncologists, radiologists, and pathologists to address this bias. Our decision to survey only urologists was in part to better understand their perspective as it pertains to initiating mechanisms of change in support of MDC; the contrary could prove valuable should future surveys include radiologists or pathologists exclusively. In addition, a majority of respondents identified men with high-risk PCa as the greatest population of MDC patients, which suggests selection bias. The greater proportion of high-risk PCa patients is likely because men in this group are less often candidates for unimodal treatment or active surveillance and more likely to require a multimodal multidisciplinary approach. Furthermore, the management of low- and intermediate-risk patients can also benefit from a multidisciplinary approach, especially as the role of prostate mpMRI in excluding men from prostate biopsy. In the past decade, the role of ablative focal and partial gland therapy is becoming part of the treatment armamentarium of PCa, which cannot be pursued without obtaining a pelvic mpMRI and suspicious lesions identified by a dedicated radiologist. Finally, depending on the urologists’ practice environment, MDC may be defined differently. For example, patients may encounter a specialist from each respective discipline (surgery, medical, and radiation oncology) in a single visit, require more than a single visit through referral to a consultant following an initial encounter, or “MDC” may refer to a tumor board-like meeting of providers who discuss treatment options before patient interaction, or any combination thereof. Thus, the interpretation of the objective findings and opinions in the present study is likely distributed among different MDC settings and cannot be generalized. Ultimately, providing a definition of MDC would benefit a more targeted approach to determining what works and what does not with respect to creating and maintaining MDC.

CONCLUSION

Most urologists consider MDC to be beneficial for PCa management with respect to the enhanced evaluation and collaboration among physicians, improved patient communication, and quality of care. Endorsement of and participation in MDC can be dependent on the setting in which the urologist initiates practice; interestingly, urologists in practice for longer than 10 years were less likely to find MDC beneficial for the management of PCa. The challenge in incentivizing providers, the logistic, and financial barriers to partake in MDC are shared among urologists, radiologists, and likely other MDC personnel. With a shift toward a targeted tissue diagnosis of PCa, the benefits of a centralized multidisciplinary evaluation and involvement of diagnosticians will become more necessary.

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There are no conflicts of interest.

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SUPPLEMENTARY APPENDIX

1. What is your age?

2. What is your gender (choose one of the following)?
   - Male
   - Female.

3. What is the context of your practice (choose one of the following)?
   - Academic
   - Non-academic.

4. How many years have you been practicing urology (choose one of the following)?
   - 0–5 years
   - 6–10 years
   - 11–20 years
   - Greater than 20 years.

5. Are you fellowship trained in urologic oncology (choose one of the following)?
   - Yes
   - No.

6. On average, how many patients do you see per month with newly diagnosed prostate cancer (choose one of the following)?
   - None
   - Between 1 and 5 patients
   - Between 6 and 10 patients
   - Between 11 and 20 patients
   - Greater than 20 patients.

7. How beneficial do you feel prostate cancer multidisciplinary clinics are in the management of prostate cancer (choose one of the following)?
   - No benefit at all
   - Slightly beneficial
   - Moderately beneficial
   - Very beneficial
   - Extremely beneficial.

8. How beneficial do you feel prostate cancer multidisciplinary clinics are in promoting prostate cancer research and patient enrollments in prostate cancer clinical trials (choose one of the following)?
   - No benefit at all
   - Slightly beneficial
   - Moderately beneficial
   - Very beneficial
   - Extremely beneficial.

9. Does your center/practice use prostate cancer multidisciplinary clinic to manage prostate cancer (choose one of the following)?
   - Yes
   - No.

10. If your practice does not utilize prostate cancer multidisciplinary clinic what is the reason (select all that apply)?
    - Lack of benefit
    - Insufficient time
    - Cost is prohibitive
    - Lack of infrastructure
    - Other ______.

11. If your practice does utilize prostate cancer multidisciplinary clinic at your institution what professional(s) participate in patient evaluation (select all that apply)?
    - Urologist
• Radiation oncologist
• Medical oncologist
• Radiologist
• Pathologist
• Advanced practice registered nurse
• Social worker
• Other ______.

12. On average, how often do you schedule prostate cancer multidisciplinary clinics (choose one of the following)?
• Once per month
• Twice per month
• Four times per month
• Greater than 4 times per month.

13. What patient population comprises your prostate cancer multidisciplinary clinic (select all that apply)?
• Clinically localized low-risk patients (Gleason Grade 3+3)
• Clinically localized intermediate-risk patients (Gleason Grade 3+4 or 4+3)
• Clinically localized high-risk patients (>Gleason 4+4)
• Active surveillance candidates
• Advanced/metastatic prostate cancer
• Recurrent prostate cancer.

14. Where are most referrals to your prostate cancer multidisciplinary clinic directed from (choose one of the following)?
• Urology clinic
• Primary care clinic
• Radiation oncology clinic
• Medical oncology clinic
• Other provider ________.