

7. Weiner AB, Patel SG, Etzioni R, Eggener SE. National trends in the management of low and intermediate risk prostate cancer in the United States. *J Urol*. 2015;193:95–102.
8. Cooperberg MR, Carroll PR. Trends in management for patients with localized prostate cancer, 1990–2013. *JAMA*. 2015;314:80–82.
9. Mottet N, Bellmunt J, Bolla M, et al. EAU-ESTRO-SIOG guidelines on prostate cancer. Part 1: screening, diagnosis, and local treatment with curative intent. *Eur Urol*. 2017;71:618–629.
10. Oberlin DT, Casalino DD, Miller FH, Meeks JJ. Dramatic increase in the utilization of multiparametric magnetic resonance imaging for detection and management of prostate cancer. *Abdominal Radiol*. 2017;42:1255–1258.
11. Ahmed HU, El-Shater Bosaily A, Brown LC, et al. Diagnostic accuracy of multi-parametric MRI and TRUS biopsy in prostate cancer (PROMIS): a paired validating confirmatory study. *Lancet*. 2017;389:815–822.
12. Coakley FV, Oto A, Alexander LF, et al. ACR appropriateness criteria(R) prostate cancer-pretreatment detection, surveillance, and staging. *J Am College Radiol*. 2017;14:S245–s257.
13. Warren JL, Klabunde CN, Schrag D, Bach PB, Riley GF. Overview of the SEER-Medicare data: content, research applications, and generalizability to the United States elderly population. *Med Care*. 2002;40. IV-3-18.
14. D'Amico AV, Whittington R, Malkowicz SB, et al. Biochemical outcome after radical prostatectomy, external beam radiation therapy, or interstitial radiation therapy for clinically localized prostate cancer. *JAMA*. 1998;280:969–974.
15. Tyson MD, Graves AJ, O'Neil B, et al. Urologist-level correlation in the use of observation for low- and high-risk prostate cancer. *JAMA Surg*. 2017;152:27–34.
16. Hoffman KE, Niu J, Shen Y, et al. Physician variation in management of low-risk prostate cancer: a population-based cohort study. *JAMA Internal Med*. 2014;174:1450–1459.
17. Elixhauser A, Steiner C, Harris DR, Coffey RM. Comorbidity measures for use with administrative data. *Med Care*. 1998;36:8–27.
18. Davidoff AJ, Gardner LD, Zuckerman IH, Hendrick F, Ke X, Edelman MJ. Validation of disability status, a claims-based measure of functional status for cancer treatment and outcomes studies. *Med Care*. 2014;52:500–510.
19. Fewell Z, Davey Smith G, Sterne JA. The impact of residual and unmeasured confounding in epidemiologic studies: a simulation study. *Am J Epidemiol*. 2007;166:646–655.
20. Austin PC. Optimal caliper widths for propensity-score matching when estimating differences in means and differences in proportions in observational studies. *Pharma Stat*. 2011;10:150–161.
21. Klotz L, Vesprini D, Sethukavalan P, et al. Long-term follow-up of a large active surveillance cohort of patients with prostate cancer. *J Clin Oncol*. 2015;33:272–277.
22. Tosoian JJ, Mamawala M, Epstein JI, et al. Intermediate and longer-term outcomes from a prospective active-surveillance program for favorable-risk prostate cancer. *J Clin Oncol*. 2015;33:3379–3385.
23. Leapman MS, Cowan JE, Nguyen HG, et al. Active surveillance in younger men with prostate cancer. *J Clin Oncol*. 2017;35:1898–1904.
24. Ouzzane A, Renard-Penna R, Marliere F, et al. Magnetic resonance imaging targeted biopsy improves selection of patients considered for active surveillance for clinically low risk prostate cancer based on systematic biopsies. *J Urol*. 2015;194:350–356.
25. Muthigi A, Sidana A, George AK, et al. Current beliefs and practice patterns among urologists regarding prostate magnetic resonance imaging and magnetic resonance-targeted biopsy. *Urol Oncol*. 2017;35. 32.e31-32.e37.
26. Jemal A, Fedewa SA, Ma J, et al. Prostate cancer incidence and PSA testing patterns in relation to USPSTF screening recommendations. *JAMA*. 2015;314:2054–2061.
27. Audit NPC. Results of the NPCA prospective audit in England and Wales for men diagnosed from 1 April 2015–31 March 2016.
28. Schmid M, Meyer CP, Reznor G, et al. Racial differences in the surgical care of medicare beneficiaries with localized prostate cancer. *JAMA Oncol*. 2016;2:85–93.
29. Davis BA, Aminawung JA, Abu-Khalaf MM, et al. Racial and ethnic disparities in oncotype DX test receipt in a statewide population-based study. *J Natl Compr Cancer Network*. 2017;15:346–354.
30. Gordon LG, James R, Tuffaha HW, Lowe A, Yaxley J. Cost-effectiveness analysis of multiparametric MRI with increased active surveillance for low-risk prostate cancer in Australia. *JMRI*. 2017;45:1304–1315.

EDITORIAL COMMENT



Active surveillance and prostate magnetic resonance imaging (MRI) have revolutionized the management of prostate cancer in recent years. This study evaluates the association between the use of prostate MRI and observation for 8144 men with newly diagnosed low-risk prostate cancer using data from the Surveillance, Epidemiology, and End Results-Medicare database. From 2010–2013, the use of MRI increased 3-fold and the use of observation increased from 30.8%–48.1% in these patients. After propensity-score matching to control for potential confounders captured in the data, the authors demonstrate that men who received a prostate MRI surrounding the diagnosis of prostate cancer were significantly more likely to undergo observation. The authors conclude that prostate MRI may increase confidence in the assignment of low-risk classification and thereby facilitate the use of active surveillance for prostate cancer.

This study offers real-world data that may support the beliefs of many urologists: a prostate MRI without suspicious lesions can make a strong case for active surveillance in an otherwise appropriate candidate. However, there are alternative explanations for the association between MRI and observation noted in this study and the authors are appropriately cautious about making an argument for a causal relationship. Prostate MRIs captured in this analysis may have been used in men who were already being managed with active surveillance. In that context, the use of observation “caused” the MRI more than the converse. Alternatively, there may be unmeasured confounders that are associated with both the use of observation and the use of prostate MRI. One such possibility is a characteristic of the treating physician. Physicians who are more likely to order a prostate MRI for their patients may also be more likely to recommend observation for men with low-risk disease. Particularly in the study period (2010–2013), physicians who were earlier adopters of prostate MRI may also have been more likely to recommend active surveillance to their low-risk prostate cancer patients. Propensity-score matching, which the authors used to generate matched cohorts of patients with and without prostate MRI, can only account for covariates that are captured in the data and cannot control for physician- or hospital-factors that are not available in administrative claims.¹

Understanding the use of active surveillance on a national level is critically important for the field of urology. Despite long-term data from several centers supporting the safety of active surveillance, its use still varies considerably from physician to physician² and the optimal protocol remains unknown.^{3,4} Innovations such as prostate MRI and biomarker tests offer us the potential to further refine our patient selection, but we do not know exactly how these tools are being used. Analyses such as this one, using nationally representative data, may help us better understand how these pieces fit

together and how we might continue to improve the management of men with prostate cancer.

Parth K. Modi, MD, MS, Department of Urology, Dow Division of Health Services Research, Michigan Medicine, Ann Arbor, MI

References

1. Sheetz KH, Derstine B, Englesbe MJ. Propensity scores for comparative effectiveness research: finding the right match. *Surgery*. 2016;160:1425–1426.
2. Auffenberg GB, Lane BR, Linsell S, Cher ML, Miller DC. Practice- vs physician-level variation in use of active surveillance for men with low-risk prostate cancer: implications for collaborative quality improvement. *JAMA Surg*. 2017;152:978–980.
3. Loeb S, Walter D, Curnyn C, Gold HT, Lepor H, Makarov DV. How active is active surveillance? Intensity of followup during active surveillance for prostate cancer in the United States. *J Urol*. 2016;196:721–726.
4. Luckenbaugh AN, Auffenberg GB, Hawken SR, et al. Variation in guideline concordant active surveillance followup in diverse urology practices. *J Urol*. 2017;197:621–626.

<https://doi.org/10.1016/j.urology.2018.07.042>
UROLOGY 124: 105–106, 2019. Published by Elsevier Inc.

AUTHOR REPLY



The ascendance of prostate magnetic resonance imaging (MRI) coincided with a popular reckoning about years of overdiagnosis and overtreatment of low-grade prostate cancer.¹ Prostate MRI has been heralded as a solution to improve accuracy when diagnosing and staging prostate cancer with 2 primary advantages: (1) detecting occult, high-grade cancer in men who would otherwise be missed, allowing timely treatment, and (2) ruling out aggressive disease in men with ostensibly low-risk cancers allowing greater confidence in avoiding treatment. Studies supporting the performance of prostate MRI in identifying clinically significant cancers have been performed under best-case circumstances—largely in high-volume centers of imaging excellence, and by experts using state-of-the-art equipment.² Therefore, it is important to begin to evaluate the assumption that MRI will lead to better clinical outcomes in the “real world.”

In this context we appreciate the thoughtful editorial addressing our study which examined the association of prostate MRI and initial management among Medicare beneficiaries with low-risk prostate cancer in Surveillance, Epidemiology, and End Results. We found that men who received prostate MRI in the period surrounding their diagnosis were more likely to be initially observed for their disease. As well-stated by the author(s), there are several alternative explanations that are important to consider in the study period where prostate MRI was in its infancy. As a methodological point, we first wish to clarify that patients in this study were included on the basis of a new diagnosis of prostate cancer, limiting the possibility that MRI was undertaken in the setting of prior active surveillance. Nonetheless, it is possible that

physicians who used MRI in the early period were more likely to recommend observation as management, particularly in light of known associations of academic institutions and observation for low-risk cancers.³ Further, we agree with the commentary that administrative claims lack clinical granularity, limiting our understanding of how MRI data was used when making decisions. For these reasons, we took care to not assert a causal relationship between prostate MRI and observation.

Notwithstanding the possibility that the use of prostate MRI is explained by provider-level variation in the use of observational management or other confounders, there are several notable findings from our study.⁴ If a causal association is validated in other studies, the utility of MRI in the management of localized prostate cancer will further support its use. In addition, we found regional, racial, and socioeconomic differences in the use of prostate MRI. In light of recent data showing the growing use of MRI in the contemporary period, there is a timely need to determine how new technologies affect entrenched disparities in prostate cancer care and outcome.⁵ Continued expansion of prostate MRI into routine care is likely. Anticipating such changes in the use of MRI and other tools, we fully agree that additional study is needed to understand the benefit of these innovations once put into practice.

Michael S. Leapman, MD, Rong Wang, PhD, Henry S. Park, MD, MPH, James B. Yu, MD, MHS, Jeffrey C. Weinreb, MD, Cary P. Gross, MD, Xiaomei Ma, PhD, Department of Urology, Yale School of Medicine, New Haven, CT; Yale Cancer Outcomes, Public Policy, and Effectiveness Research Center, New Haven, CT; Department of Chronic Disease Epidemiology, Yale School of Public Health, New Haven, CT; Department of Therapeutic Radiology, Yale School of Medicine, New Haven, CT; Department of Radiology and Biomedical Imaging, Yale School of Medicine, New Haven, CT; Department of Internal Medicine, Yale School of Medicine, New Haven, CT

References

1. Bhindi B, Mamdani M, Kulkarni GS, et al. Impact of the U.S. Preventive Services Task Force recommendations against prostate specific antigen screening on prostate biopsy and cancer detection rates. *J Urol*. 2015;193:1519–1524.
2. Siddiqui MM, Rais-Bahrami S, Turkbey B, et al. Comparison of MR/ultrasound fusion-guided biopsy with ultrasound-guided biopsy for the diagnosis of prostate cancer. *JAMA*. 2015;313:390–397.
3. Lester-Coll NH, Park HS, Rutter CE, et al. The association between evaluation at academic centers and the likelihood of expectant management in low-risk prostate cancer. *Urology*. 2016;96:128–135.
4. Tyson MD, Graves AJ, O’Neil B, et al. Urologist-level correlation in the use of observation for low- and high-risk prostate cancer. *JAMA Surg*. 2017;152:27–34.
5. Liu W, Patil D, Howard DH, et al. Adoption of prebiopsy magnetic resonance imaging for men undergoing prostate biopsy in the United States. *Urology*. 2018;117:57–63.

<https://doi.org/10.1016/j.urology.2018.07.043>
UROLOGY 124: 106, 2019. Published by Elsevier Inc.