

ORIGINAL ARTICLE

MRI-Targeted or Standard Biopsy
for Prostate-Cancer Diagnosis

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ABSTRACT

BACKGROUND

Multiparametric magnetic resonance imaging (MRI), with or without targeted biopsy, is an alternative to standard transrectal ultrasonography–guided biopsy for prostate-cancer detection in men with a raised prostate-specific antigen level who have not undergone biopsy. However, comparative evidence is limited.

METHODS

In a multicenter, randomized, noninferiority trial, we assigned men with a clinical suspicion of prostate cancer who had not undergone biopsy previously to undergo MRI, with or without targeted biopsy, or standard transrectal ultrasonography–guided biopsy. Men in the MRI-targeted biopsy group underwent a targeted biopsy (without standard biopsy cores) if the MRI was suggestive of prostate cancer; men whose MRI results were not suggestive of prostate cancer were not offered biopsy. Standard biopsy was a 10-to-12–core, transrectal ultrasonography–guided biopsy. The primary outcome was the proportion of men who received a diagnosis of clinically significant cancer. Secondary outcomes included the proportion of men who received a diagnosis of clinically insignificant cancer.

RESULTS

A total of 500 men underwent randomization. In the MRI-targeted biopsy group, 71 of 252 men (28%) had MRI results that were not suggestive of prostate cancer, so they did not undergo biopsy. Clinically significant cancer was detected in 95 men (38%) in the MRI-targeted biopsy group, as compared with 64 of 248 (26%) in the standard-biopsy group (adjusted difference, 12 percentage points; 95% confidence interval [CI], 4 to 20; $P=0.005$). MRI, with or without targeted biopsy, was noninferior to standard biopsy, and the 95% confidence interval indicated the superiority of this strategy over standard biopsy. Fewer men in the MRI-targeted biopsy group than in the standard-biopsy group received a diagnosis of clinically insignificant cancer (adjusted difference, –13 percentage points; 95% CI, –19 to –7; $P<0.001$).

CONCLUSIONS

The use of risk assessment with MRI before biopsy and MRI-targeted biopsy was superior to standard transrectal ultrasonography–guided biopsy in men at clinical risk for prostate cancer who had not undergone biopsy previously. (Funded by the National Institute for Health Research and the European Association of Urology Research Foundation; PRECISION ClinicalTrials.gov number, NCT02380027.)

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MEN WITH A CLINICAL SUSPICION OF prostate cancer on the basis of an elevated prostate-specific antigen (PSA) level or an abnormal digital rectal examination are typically offered a standard transrectal ultrasonography–guided biopsy of the prostate during which 10 to 12 cores are obtained. This approach is associated with the underdetection of higher-grade (clinically significant) prostate cancers and the overdetection of low-grade (clinically insignificant) cancers.¹ Despite randomized trials showing that men with clinically insignificant cancer do not benefit from treatment,^{2,3} its identification still results in the overtreatment of some men. Some men will receive radical treatment that has side effects,^{4,5} and others will undergo active surveillance with repeated assessment over time that has costs for patients and health care systems.^{6,7}

An alternative diagnostic pathway in men with a clinical suspicion of prostate cancer involves multiparametric magnetic resonance imaging (MRI). With better standardization of the conduct and reporting of multiparametric MRI, the ability to detect clinically significant cancer and to rule it out has improved over the past decade.^{1,8,9} Multiparametric MRI could be used as a triage test to avoid a biopsy if the results were negative,¹ whereas positive results could be used for targeting abnormal areas in the prostate during biopsy.^{10,11}

In single-center studies, the approach of obtaining MRI-targeted biopsy cores alone, without performing standard biopsies, has shown similar or higher rates of detection of clinically significant cancer¹²⁻¹⁵ and lower rates of detection of clinically insignificant cancer¹⁵ than standard biopsy. We compared MRI-targeted biopsy with standard transrectal ultrasonography–guided biopsy in a pragmatic, multicenter, randomized trial. The PRECISION (Prostate Evaluation for Clinically Important Disease: Sampling Using Image Guidance or Not?) trial aimed to evaluate prospectively whether multiparametric MRI, with targeted biopsy in the presence of an abnormal lesion, was noninferior to standard transrectal ultrasonography–guided biopsy in the detection of clinically significant prostate cancer in men with a clinical suspicion of prostate cancer who had not undergone biopsy of the prostate previously.

METHODS

TRIAL DESIGN

We conducted this multicenter, randomized, noninferiority trial at 25 centers in 11 countries (Table S1 in the Supplementary Appendix, available with the full text of this article at NEJM.org). Men who provided written informed consent were randomly assigned in a 1:1 ratio to either the MRI-targeted biopsy group or the standard-biopsy group (Fig. S1 in the Supplementary Appendix). The assignment sequence used computer-generated, randomly permuted blocks of unequal size, stratified according to center. Group assignments were revealed by the Web-based system once a participant had been assessed as eligible and had provided written informed consent.

The full trial protocol, available at NEJM.org, has been published previously¹⁶ and was approved by the ethics review board at each participating institution. The trial was monitored by an independent trial steering committee and data and safety monitoring committee. The trial was designed by the Standards of Reporting for MRI-Targeted Biopsy Studies (START) working group,¹⁰ and final decisions were made by the first author and the last two authors. Data were gathered by the trial team members who are listed in Section S1 in the Supplementary Appendix. One author analyzed the data, and the analysis was independently verified by another author. The authors assume responsibility for the accuracy and completeness of the data and analyses and for the adherence of the trial to the protocol. The first draft of the manuscript was written by the first author.

No commercial entity was involved in the trial. The trial was funded by the National Institute for Health Research and the European Association of Urology Research Foundation, with trial governance from University College London. The funders had no role in the protocol development, data analysis or interpretation, or manuscript preparation.

PARTICIPANTS

Participants were recruited in outpatient clinics and were eligible for enrollment if they had not undergone biopsy of the prostate previously and had been referred with a clinical suspicion of prostate cancer on the basis of an elevated PSA

level, an abnormal digital rectal examination, or both (Table S2 in the Supplementary Appendix). Participants were required to have a PSA level of 20 ng per milliliter or less, to have results on digital rectal examination that did not suggest extracapsular disease, and to be suitable candidates for biopsy of the prostate and for MRI.

MRI AND MRI-TARGETED BIOPSY

Multiparametric MRI was performed with the use of a 1.5-T or 3.0-T scanner with a pelvic phased-array coil, with or without an endorectal coil (Table S3 in the Supplementary Appendix). T₂-weighted, diffusion-weighted, and dynamic contrast-enhanced sequences were acquired according to minimum standards that have been set by consensus guidelines.⁸ Areas on the multiparametric MRI that were suggestive of prostate cancer were categorized by a local radiologist according to the Prostate Imaging–Reporting and Data System, version 2 (PI-RADS v2),⁹ on a scale from 1 to 5, with higher numbers indicating a greater likelihood of clinically significant cancer. Table S4 in the Supplementary Appendix provides details regarding the experience of the clinicians who took part in the trial.

Men who had a positive result on the multiparametric MRI — that is, in whom an area with a score of 3 (equivocal regarding the likelihood of prostate cancer), 4 (likely to be prostate cancer), or 5 (highly likely to be prostate cancer) was identified — underwent MRI-targeted biopsy with the use of real-time ultrasonographic guidance. A maximum of three areas that were suggestive of prostate cancer were permitted to be chosen for targeted biopsy, with a maximum of 4 biopsy cores obtained per area, resulting in a maximum of 12 biopsy cores obtained per participant. MRI-targeted biopsy registration (i.e., matching of the image of the target on MRI with the real-time image of the prostate during biopsy) could be performed by means of visual registration or software-assisted registration (also known as MRI–ultrasonographic fusion)¹⁰ and could be carried out through the transrectal or transperineal route, according to local expertise (Table S5 in the Supplementary Appendix). In the absence of abnormal areas on the multiparametric MRI (i.e., a negative result, with a score of 1 or 2), the participant was not offered a protocol biopsy.

STANDARD TRANSRECTAL ULTRASONOGRAPHY–GUIDED BIOPSY

Biopsy was carried out by experienced operators who used a standard transrectal technique. A total of 10 to 12 biopsy cores were obtained from the peripheral zone of the prostate at the base, mid gland, and apex.¹⁷

PARTICIPANT-REPORTED OUTCOME MEASURES

Participant-reported questionnaires were used to collect data about intervention-specific side effects immediately and at 30 days after biopsy and after MRI.^{16,18} Health-related quality of life was assessed with the use of the EuroQol–5 Dimension Self-Report Questionnaire at baseline, 24 hours after the intervention, and 30 days after the intervention.^{19–21}

OUTCOMES

The primary outcome was the proportion of men with clinically significant cancer, defined as the presence of a single biopsy core indicating disease of Gleason score 3+4 (Gleason sum of 7) or greater (the Gleason score is composed of a primary [most predominant] grade plus a secondary [highest nonpredominant] grade; the range for a primary or secondary grade is from 3 to 5, with the Gleason sum ranging from 6 to 10, and with higher scores indicating a more aggressive form of prostate cancer). Secondary outcomes included the proportion of men with clinically insignificant cancer (Gleason score 3+3), the proportion of men in the MRI-targeted biopsy group who did not undergo biopsy, and the proportion of men with adverse events after the intervention. All the secondary outcomes are listed in Table S6 in the Supplementary Appendix. Outcomes were reported according to the START guidelines,¹⁰ which are the consensus criteria for reporting studies of MRI-targeted prostate biopsies.

FOLLOW-UP

Participants were followed until the visit at which their treatment decisions were made or until their 30-day postintervention questionnaires were completed, whichever was later. Participants who underwent further diagnostic tests as a result of the outcome of the treatment-decision visit were additionally followed until after the results of the further investigation were made available and recorded. These participants included men who

had negative test results in either the standard-biopsy group or the MRI-targeted biopsy group and underwent additional testing. Participants who had negative test results in either group at the end of the trial period returned to standard-care monitoring at each center, which typically involved surveillance of the PSA level. Participants who underwent radical prostatectomy on the basis of their treatment decision were also followed until the pathological testing results of their radical prostatectomy were available. Participants provided written informed consent for long-term follow-up as part of future studies involving additional contact from the trial center and linkage to national databases.

QUALITY CONTROL

Uroradiologists and pathologists at the coordinating center, who were unaware of the results of the original reports, reviewed 25% of the multiparametric MRIs and 15% of the original pathological specimens. These MRIs and specimens had been chosen at random from participants at every site.

STATISTICAL ANALYSIS

Using a noninferiority margin of 5 percentage points that was agreed on at an expert consensus group meeting¹⁰ and a one-sided alpha level of 2.5%, we calculated that the randomization of 422 men would provide the trial with 90% power to show the noninferiority of MRI, with or without targeted biopsy, to standard biopsy, assuming a detection rate of clinically significant cancer of 40% in the group that underwent MRI, with or without targeted biopsy, and 30% in the standard-biopsy group. This sample size was increased to 470 to allow for a 10% rate of withdrawal and loss to follow-up. Detailed justification of the sample size is provided in the protocol.¹⁶

The statistical analysis plan was prespecified and approved by the data and safety monitoring committee before the analysis of any data. For the primary outcome, if the lower boundary of the two-sided 95% confidence interval for the difference in the rates of detection of clinically significant cancer in the MRI-targeted biopsy group relative to the standard-biopsy group was greater than -5 percentage points, then MRI, with or without targeted biopsy, would be deemed to be noninferior. Furthermore, if the lower boundary was greater than zero, superior-

ity would be claimed. The difference was estimated with the use of a generalized linear mixed model (with the use of an identity link function with a binomial distribution) that included trial center as a random effect.

All the participants who underwent randomization were included in the primary intention-to-treat analysis. Analyses were repeated in the modified intention-to-treat population and the per-protocol population as sensitivity analyses (Table S7A, S7B, and S7C in the Supplementary Appendix). The modified intention-to-treat analysis excluded participants who did not complete a diagnostic test strategy; this analysis was carried out to prevent unequal withdrawal in the two groups from contributing to a difference between the groups. The per-protocol analysis included only men who underwent the randomly assigned testing procedure as specified in the protocol; this analysis was carried out because this was a noninferiority trial, so a per-protocol analysis would reduce the chance of biasing the result toward the null. If, after participants had undergone the trial test procedures, further tests provided different information about the presence of cancer, no adjustment was made to the analyses of the primary and secondary outcomes. A post hoc Bonferroni correction was used to adjust for three secondary outcomes (proportion of men with clinically insignificant cancer, maximum cancer core length, and health-related quality of life), with a two-sided P value of less than 0.017 considered to indicate statistical significance. The methods of analysis of the other outcomes are described in Section S2 in the Supplementary Appendix.

RESULTS

TRIAL POPULATION

From February 2016 through August 2017, a total of 500 participants underwent randomization at 23 of the 25 sites, with 252 participants being assigned to the MRI-targeted biopsy group and 248 to the standard-biopsy group (Fig. 1, and Table S1 in the Supplementary Appendix). The characteristics of the participants at baseline were similar in the two groups (Table 1).

A total of 71 of 252 participants (28%) in the MRI-targeted biopsy group had a result on multiparametric MRI that was not suggestive of prostate cancer (PI-RADS v2 score, ≤ 2), and so they did not undergo biopsy. Among the partici-

pants with a positive result on multiparametric MRI, 51 of 175 (29%) had a PI-RADS v2 score of 3, 70 (40%) had a score of 4, and 54 (31%) had a score of 5 (Table S8 in the Supplementary Appendix). The remaining 6 men did not complete the MRI assessment (Fig. 1). Among the participants who underwent biopsy, a median of 4 biopsy cores were obtained in the MRI-targeted biopsy group, as compared with a median of 12 cores in the standard-biopsy group (Table S9 in the Supplementary Appendix).

OUTCOMES

Clinically significant cancer was detected in 95 men (38%) in the MRI-targeted biopsy group, as compared with 64 (26%) in the standard-biopsy group (adjusted difference, 12 percentage points; 95% confidence interval [CI], 4 to 20; $P=0.005$) (Table 2). The lower boundary of the 95% confidence interval for the difference was greater than -5 percentage points, so MRI, with or without targeted biopsy, was deemed to be non-inferior to standard transrectal ultrasonography-guided biopsy in the detection of clinically significant cancer. Furthermore, the 95% confidence interval showed the superiority of MRI, with or without targeted biopsy, over transrectal ultrasonography-guided biopsy. The results were consistent in the modified intention-to-treat and per-protocol populations (Fig. 2).

Fewer participants received a diagnosis of clinically insignificant cancer in the MRI-targeted biopsy group than in the standard-biopsy group (23 men [9%] vs. 55 [22%]; adjusted difference, -13 percentage points; 95% CI, -19 to -7 ; $P<0.001$). In men with cancer, the mean maximum cancer core length was 7.8 mm in the MRI-targeted biopsy group and 6.5 mm in the standard-biopsy group (adjusted mean difference, 1.0 mm; 95% CI, 0.0 to 2.1; $P=0.053$). Interpretations of the results for these secondary outcomes were unchanged by post hoc Bonferroni correction (see Section S3 in the Supplementary Appendix).

A greater percentage of cores were positive for cancer in the MRI-targeted biopsy group (422 of 967 cores [44%]) than in the standard-biopsy group (515 of 2788 [18%]). Among men with a positive result on MRI, the percentage of men with clinically significant cancer was highest among participants with a PI-RADS v2 score of 5 (83%), followed by those with a score of 4 (60%) and those with a score of 3 (12%). Con-

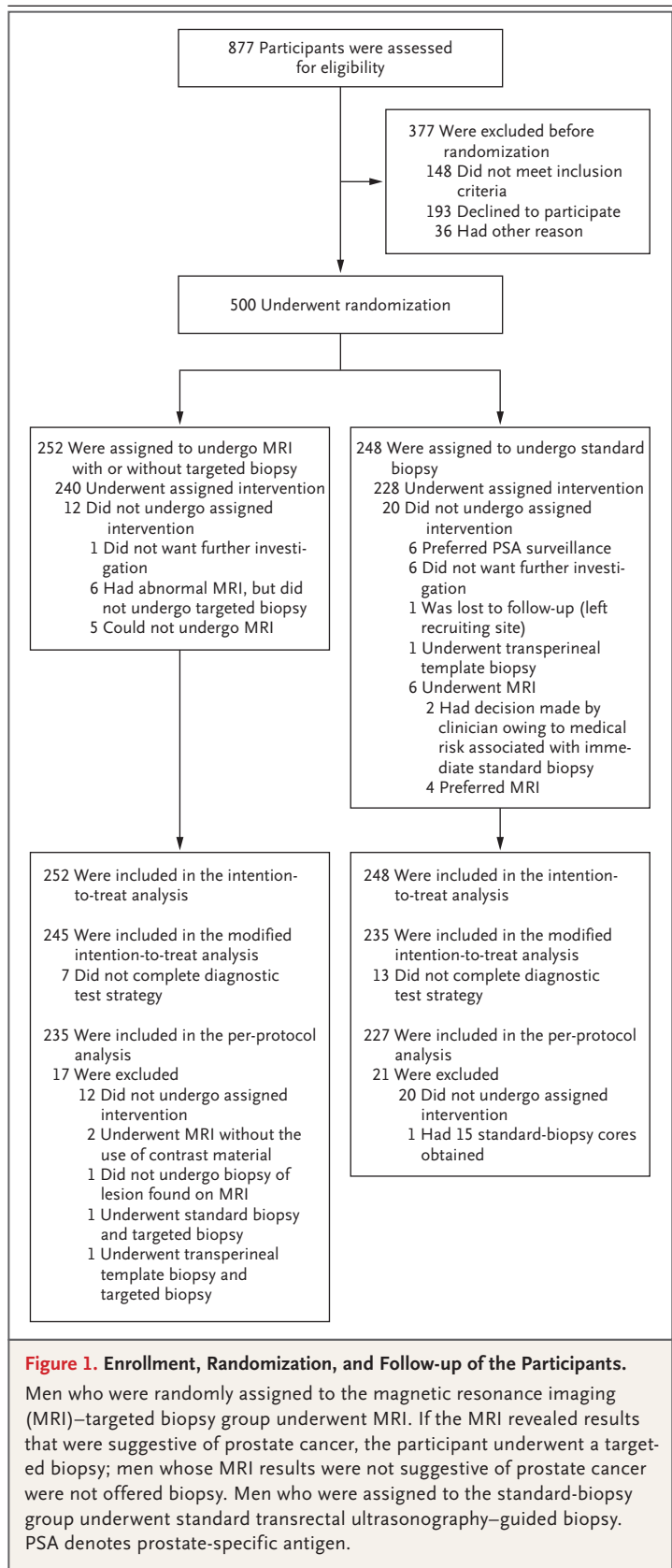


Table 1. Characteristics of the Participants at Baseline.*

Characteristic	MRI-Targeted Biopsy Group (N = 252)	Standard-Biopsy Group (N = 248)
Age — yr	64.4±7.5	64.5±8.0
PSA level — ng/ml		
Median	6.75	6.50
Interquartile range	5.16–9.35	5.14–8.65
Family history of prostate cancer — no. (%)	48 (19)	40 (16)
Abnormal digital rectal examination — no. (%)	36 (14)	38 (15)

* Plus–minus values are means ±SD. Men who were randomly assigned to the MRI-targeted biopsy group underwent MRI. If the MRI revealed results that were suggestive of prostate cancer, the participant underwent a targeted biopsy; men whose MRI results were not suggestive of prostate cancer were not offered biopsy. Men who were assigned to the standard-biopsy group underwent standard transrectal ultrasonography–guided biopsy. The characteristics of the participants at baseline were similar in the two groups. PSA denotes prostate-specific antigen.

versely, the percentage of men without cancer was highest among participants with a PI-RADS v2 score of 3 (67%), followed by those with a score of 4 (31%) and those with a score of 5 (6%) (Fig. 3).

QUALITY OF LIFE AND SAFETY

Health-related quality of life at 24 hours and at 30 days after the intervention did not differ significantly between the MRI-targeted biopsy group and the standard-biopsy group. The intervention was associated with similar results regarding immediate postintervention discomfort and pain in the two groups. The participant-reported complications at 30 days were less frequent in the MRI-targeted biopsy group than in the standard-biopsy group, including events of blood in the urine (30% vs. 63%), blood in the semen (32% vs. 60%), pain at the site of the procedure (13% vs. 23%), rectal bleeding (14% vs. 22%), and erectile dysfunction (11% vs. 16%). These findings reflected the lower percentage of men undergoing biopsy and fewer biopsy cores obtained in the MRI-targeted biopsy group than in the standard-biopsy group. A total of 2% of the men in the MRI-targeted biopsy group and 2% in the standard-biopsy group had serious adverse events. Details regarding health-related quality-of-life scores, participant-reported complications, and adverse events are provided in Tables S10 through S12 in the Supplementary Appendix.

FURTHER DIAGNOSTIC TESTING

After the discussion of the test results with each participant, more men in the standard-biopsy group (39 men [16%]) than in the MRI-targeted

biopsy group (7 [3%]) underwent further diagnostic tests. Of the 39 further diagnostic tests that were performed in the standard-biopsy group, 38 (in 15% of the participants in the group) were diagnostic multiparametric MRIs that were carried out in men with negative results on the transrectal ultrasonography–guided biopsy. In the MRI-targeted biopsy group, only 3 participants (1%) with negative results on MRI subsequently underwent standard transrectal ultrasonography–guided biopsy. More men who underwent MRI-targeted biopsy (104 men [41%]) than men who underwent standard transrectal ultrasonography–guided biopsy (74 [30%]) adopted a strategy of monitoring of the PSA level, although the percentage of men undergoing active surveillance or radical treatment was similar in the two groups.

Among the participants who underwent further biopsy, clinically significant cancer was detected in none of the 4 men in the MRI-targeted biopsy group and in 3 of 9 men (33%) in the standard-biopsy group. Of the 71 men with negative results on MRI and no biopsy, 3 (4%) were discharged, 62 (87%) were referred for monitoring of the PSA level, 3 (4%) underwent further prostate biopsy (all had negative results), 1 (1%) underwent an additional multiparametric MRI, and 2 (3%) had missing information. The percentage of men whose Gleason score was upgraded (i.e., found to be higher) after radical prostatectomy was similar in the MRI-targeted biopsy group (5 of 30 men [17%]) and the standard-biopsy group (4 of 27 [15%]). Details are provided in Tables S13 through S15 in the Supplementary Appendix.

Table 2. Comparison of Cancer Detection between Groups.*

Outcome	MRI-Targeted Biopsy Group (N=252)	Standard-Biopsy Group (N=248)	Difference†	P Value
Biopsy outcome — no. (%)			—	—
No biopsy because of negative result on MRI	71 (28)	0		
Benign tissue	52 (21)	98 (40)		
Atypical small acinar proliferation	0	5 (2)		
High-grade prostatic intraepithelial neoplasia	4 (2)	10 (4)		
Gleason score				
3+3	23 (9)	55 (22)		
3+4	52 (21)	35 (14)		
3+5	2 (1)	1 (<1)		
4+3	18 (7)	19 (8)		
4+4	13 (5)	6 (2)		
4+5	7 (3)	2 (1)		
5+5	3 (1)	1 (<1)		
No biopsy‡	4 (2)	3 (1)		
Withdrawal from trial§	3 (1)	13 (5)		
Clinically significant cancer¶				
Intention-to-treat analysis — no. (%)	95 (38)	64 (26)	12 (4 to 20)	0.005
Modified intention-to-treat analysis — no./total no. (%)	95/245 (39)	64/235 (27)	12 (3 to 20)	0.007
Per-protocol analysis — no./total no. (%)	92/235 (39)	62/227 (27)	12 (3 to 20)	0.007
Clinically insignificant cancer — no. (%)	23 (9)	55 (22)	-13 (-19 to -7)	<0.001
Maximum cancer core length — mm	7.8±4.1	6.5±4.5	1.0 (0.0 to 2.1)	0.053
Core positive for cancer — no./total no. of cores (%)	422/967 (44)	515/2788 (18)	—	—
Men who did not undergo biopsy — no. (%)	78 (31)	16 (6)	—	—

* Clinically significant cancer was defined as the presence of a single biopsy core indicating disease of Gleason score 3+4 (Gleason sum of 7) or greater, and clinically insignificant cancer as a biopsy sample with a Gleason score of 3+3 (Gleason sum of 6). The Gleason score is composed of a primary (most predominant) grade plus a secondary (highest nonpredominant) grade; the range for a primary or secondary grade is from 3 to 5, with the Gleason sum ranging from 6 to 10, and with higher scores indicating a more aggressive form of prostate cancer.

† Differences between rates are shown in percentage points, and the difference in maximum cancer core length is shown in millimeters. Differences in the percentages of men with clinically significant cancer detected and men with clinically insignificant cancer were calculated with a generalized linear mixed model (with the use of an identity link function with a binomial distribution) that included trial center as a random effect. The between-center variance estimates for the intention-to-treat analysis of the proportion of men with clinically significant cancer was 0.002 and for the proportion of men with clinically insignificant cancer was 0; the 95% prediction intervals for the detection rates of clinically significant and clinically insignificant cancer, incorporating between-center variation, were 14 to 39% and 17 to 28%, respectively, for standard biopsy, and 26 to 51% and 4 to 11%, respectively, for MRI-targeted biopsy. The difference in the maximum cancer core length was calculated with the use of a linear mixed model with trial center as a random effect. The between-center estimate of variance was 2.14; the 95% prediction interval for the maximum cancer core length, incorporating between-center variation, was 3.3 to 9.8 mm for standard biopsy and 4.4 to 10.8 mm for MRI-targeted biopsy.

‡ In four participants in the MRI-targeted biopsy group, MRI identified at least one area with a score on the Prostate Imaging–Reporting and Data System, version 2, of 3 or greater (on a scale from 1 to 5, with higher numbers indicating a greater likelihood of clinically significant cancers), but targeted biopsy was not performed. In the standard-biopsy group, three participants declined transrectal ultrasonography–guided biopsy and underwent an MRI. The MRI revealed no areas that were suggestive of prostate cancer, and the participants did not undergo biopsy.

§ These participants did not complete any diagnostic test.

¶ The intention-to-treat analysis included all the participants who underwent randomization, the modified intention-to-treat analysis excluded participants who did not complete a diagnostic test strategy, and the per-protocol analysis included only participants who underwent the randomly assigned testing procedure as specified in the protocol.

|| Data include men who did not undergo biopsy because they withdrew before undergoing any diagnostic test or because they did not complete the diagnostic strategy.

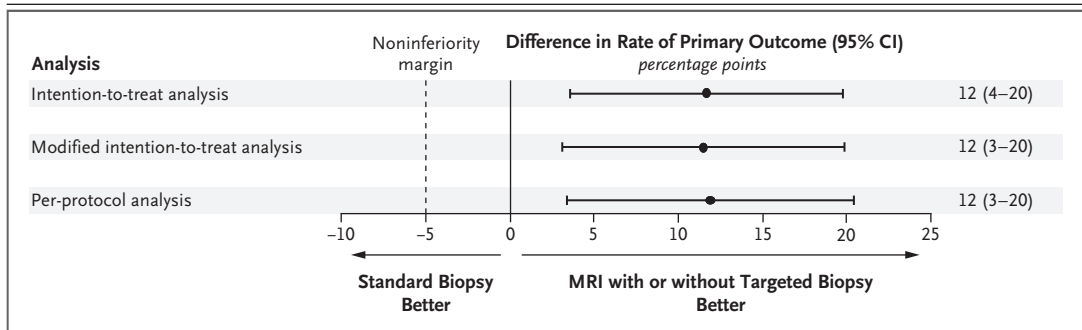


Figure 2. Intention-to-Treat, Modified Intention-to-Treat, and Per-Protocol Analyses of the Primary Outcome for the Detection of Clinically Significant Prostate Cancer.

Shown are the absolute differences between the MRI-targeted biopsy group and the standard-biopsy group in the rates of detection of clinically significant cancer. The intention-to-treat analysis included all the participants who underwent randomization, the modified intention-to-treat analysis excluded participants who did not complete a diagnostic test strategy, and the per-protocol analysis included only participants who underwent the randomly assigned testing procedure as specified in the protocol. If the lower boundary of the two-sided 95% confidence interval for the difference (MRI-targeted biopsy group minus standard-biopsy group) was greater than -5 percentage points (dashed line), then MRI, with or without targeted biopsy, would be deemed to be noninferior. If the lower boundary was greater than zero (solid line), superiority would be claimed.

QUALITY CONTROL

Results of the quality-control review of multiparametric MRI showed that the percentage of cases that were scored with agreement for concordant biopsy decision by the central radiology team and the site radiologist was 78% (50 of 64 cases). The percentage of cases that were scored with agreement on the Gleason score by the central pathologists and the site pathologist was 88% (53 of 60 cases). Details are provided in Table S16A, S16B, and S16C in the Supplementary Appendix.

DISCUSSION

The ideal test for prostate cancer would be minimally invasive, have few side effects, identify a high proportion of men who would benefit from treatment, and minimize the identification of men with clinically insignificant cancer in order to prevent overtreatment. In men with a clinical suspicion of prostate cancer who had not undergone biopsy of the prostate previously, the PRECISION trial showed that MRI, with or without targeted biopsy, appeared to achieve these goals better than the traditional standard of care, transrectal ultrasonography-guided biopsy. MRI, with or without targeted biopsy, led to fewer men undergoing biopsy, more clinically significant cancers being identified, less over-detection of clinically insignificant cancer, and

fewer biopsy cores being obtained than did standard transrectal ultrasonography-guided biopsy. Slightly more than one quarter of the men avoided a biopsy altogether, and the 30-day participant-reported side-effect profile appeared to be more favorable in the MRI-targeted biopsy group than in the standard-biopsy group. The MRI-targeted biopsy approach was also well adhered to by the participants and clinicians, with only 7 of 252 men (3%) not completing the diagnostic test strategy (Fig. 1).

The results of single-center studies have been mixed. Some studies have not shown the superiority of an MRI-based pathway over transrectal ultrasonography-guided biopsy, although these comparisons were likely to have been underpowered.^{12,22} Other single-center studies have shown advantages of an MRI-based diagnostic pathway over transrectal ultrasonography-guided biopsy,^{13,14,23} and a meta-analysis of published studies had findings concordant with those of our trial.¹⁵ These single-center studies have limitations in their lack of generalizability, and the majority of the studies were small and nonrandomized.

The PRECISION trial was an international trial, and key strengths included its size and pragmatism.²⁴ We did not limit the performance of MRI-targeted biopsy to highly experienced operators, and most of the participating investigators had modest experience with MRI-targeted biopsy, particularly as compared with standard

Ghent University Hospital, Ghent, Belgium (G.M.V.); University Hospital Bern, Bern, Switzerland (S.B.); Weill Cornell Medicine, New York–Presbyterian Hospital (J.C.H., D.M.), and New York University Langone Medical Center (S.S.T.), New York; National Institutes of Health, Bethesda, MD (P.P.); and the University of Southern California Institute of Urology, Keck School of Medicine, Los Angeles (I.G.).

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