Original Paper

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Current Role of Multiparametric MRI and MRI Targeted Biopsies for Prostate Cancer Diagnosis in Germany: A Nationwide Survey

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Keywords

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Abstract

Introduction: Multiparametric MRI (mpMRI) and MRI targeted biopsies (MRtb) are a new standard in prostate cancer (PCa) screening and diagnosis. Guidelines already include this approach for patients at risk. We aimed to gather information from German urologists about their knowledge, routine use, and attitude toward mpMRI and consecutive biopsy methods. Materials and Methods: An anonymous online questionnaire was sent via Survey Monkey to the members of the German Society of Urology (DGU). Statistical analyses were performed using SPSS version 25.0. Results: 496 members with a median age of 48.6 years (±11.7) participated in the survey. The majority rated mpMRI of the prostate as a very useful diagnostic tool (72.7%). MRtb of the prostate was considered as very advantageous (71.5%). MpMRI was used by 95.9%, and 83.2% also recommended MRtb predomi-

nantly in clinical institutions. For targeted biopsy, MRI-ultrasound fusion biopsy was clearly favored (75.8%). MpMRI was mostly used in patients with previously negative biopsy (90.9%) and in patients under active surveillance (60.9%). Arguments against the use of prostate mpMRI are costs (84.9%) and/or lack of sufficient radiological infrastructure (17.4%). **Conclusion:** Our data illustrate the meanwhile high acceptance and clinical use of the prostate mpMRI and MRtb in Germany.

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Introduction

While there is an intense debate about the overdetection and overtreatment of low-risk prostate cancer (PCa) and the need for a more sensitive detection of clinically significant PCa, multiparametric MRI (mpMRI) is a min-

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Table 1. Characteristics of the participants

Variables	Total, n = 496
Mean age (SD), years	48.56 (11.66)
Gender, %	
Male	82.5
Female	17.5
Region, %	
Eastern Germany	13.40
Western Germany	86.60
Rural/urban setting, %	
<20,000 citizens	12.02
20,000-100,000 citizens	30.60
>100,000 citizens	54.38
Practice type, %	
University	12.15
Academic	20.85
Communal hospital	12.96
Medical practice	54.04
Fully trained urologist, %	
Yes	90.78
No	9.22
Years in practice, %	
<10 years	24.95
10–20 years	28.83
>20 years	46.22
Trained in PCA diagnosis and treatment, %	
Yes	98.37
No	1.63
Trained in prostate biopsy, %	
Yes	95.10
No	4.90
Average number of prostate biopsies per patie	nt, %
1–5	0.41
6	1.45
6–9	1.24
10–12	72.20
13–20	19.92
>20	4.77
Average number of prostate biopsies monthly	
<5, n (%)	160 (33.54)
5–10, <i>n</i> (%)	187 (39.20)
11–15, <i>n</i> (%)	49 (10.27)
>15, n (%)	81 (16.99)
Do not use prostate MRI	16 (3.52)

imally invasive tool to identify and risk stratify patients prior to prostate biopsy [1]. Recent advances in mpMRI for prostate imaging along with the evolution of MR targeted biopsy techniques have resulted in a paradigm shift in PCa evaluation. In favor to identify more clinically significant (mostly defined as Gleason score $\geq 7a$) PCa cases while avoiding detection of low-risk disease, mpMRI is a very popular and widely used option.

MRI targeted biopsy (MRtb) has a proven utility in several clinical scenarios, such as in patients with prior negative transrectal ultrasound (TRUS)-guided prostate biopsy and in the monitoring of patients under active surveillance (AS) [2, 3]. Especially lesions in areas that are mostly undersampled with TRUS biopsy (e.g., midline, anterior, and distal apical) can be thoroughly assessed by mpMRI and MRtb. Finally, the PRECISION study also forwarded MRtb as the preferable option for diagnostic workup in patients with suspicion of PCa, who had no previous prostate biopsy [4], and this may further change clinical practice guidelines in the near future.

To the best of our knowledge, there is only very little information about the common practice for MRI diagnostics in PCa and the use of MRtb in Germany. Therefore, we assessed the current knowledge, professional attitude, and practice patterns concerning mpMRI and MRtb for PCa diagnosis among urologists in Germany.

Patients and Methods

Survey

After reviewing the literature, a 30-item questionnaire was designed to collect demographic data and information on urologists' opinion regarding the use of mpMRI and MRtb to detect PCa. The questionnaire contained open questions, multiple choice questions, and certain questions that allowed respondents to "select all that apply." Information was obtained on the respondents' age, gender, practice region, urban area, practice type, level of training, years in practice, number of prostate biopsies performed, and use of prostate mpMRI and MRtb in daily practice.

Study Design

A link to the survey together with a personal invitation from the German Prostate Cancer Consortium (DPKK) was sent through email to all members of the German Society of Urology (DGU). With over 6,500 members, this society is one of the largest medical societies in Germany with a heterogeneous member population of clinicians and urologic practitioners in communal and academic hospitals or in ambulant offices. Approximately 4,900 members received the email invitation. The responses were collected in an SPSS spreadsheet in an anonymous fashion.

Statistics

Statistical analyses were performed using SPSS version 25.0 (IBM, Mount Kisco, NY, USA). Univariate logistic regression analysis was performed to identify factors associated with the use of mpMRI for PCa diagnosis ("Do you use mpMRI for prostate cancer diagnosis?"), the evaluation of the usefulness of MRI targeted biopsy ("How beneficial is MRI targeted biopsy for prostate cancer detection?"), and the recommendation of MRI targeted biopsy for PCa diagnosis ("Would you recommend the use of MRI targeted biopsy for prostate cancer detection?"). Respondent's age, gender, practice region, urban area, practice type, level of training, experience in urology practice, experience in prostate biopsies, and

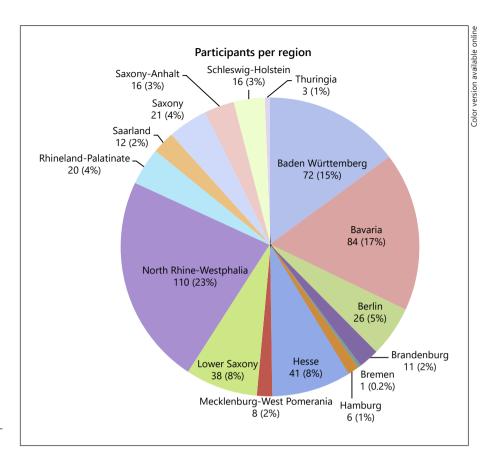


Fig. 1. Distribution of participants throughout Germany.

number of prostate biopsies per patient were used in regression analysis. For significant associations identified in the univariate regression analyses, all factors were identified that changed the calculated odds ratio >10%, and these were included in the respective multivariate model. Statistical significance was defined as p < 0.05.

Results

A total of 496 responses (response rate 496/4,900; 10.1%) were received. Characteristics of the participants are indicated in Table 1. Their distribution throughout Germany is shown in Figure 1. The mean age of the participants was $48.6~(\pm11.7)$ years, 82.5% were male, and 17.5% female. 90.8% were trained urologists with an overall distribution in favor of practicing in a medical practice (54.0%). Selected answers from the participants to the questions of the survey are shown in Figure 2. Table 2 gives an overview on the participants' answers to all survey questions. The majority of participants rated mpMRI of the prostate as very or even extremely beneficial for the evaluation of PCa (72.7%). Consequently, an MRtb of the prostate was usually rated as very or extremely advanta-

geous (71.5%). An mpMRI of the prostate was used by 95.9% of the participants, and 83.2% also recommended MRtb. Concerning the technique for MRtb, software-based MRI-ultrasound fusion biopsy was clearly favored (75.8%) over a cognitive fusion biopsy (25.9%) and "in bore" MRI biopsy (4.5%; Table 2).

In most cases, MRtb was used in patients with previously negative prostate biopsy (90.4%) and in patients with PCa under AS (54.6%). 38.5% of participants would also use MRtb in biopsy-naïve patients. In this context, German urologists stated a long waiting period for a prostate mpMRI of 2–4 weeks (48.1%) or even >4 weeks (25.5%; Table 3). Arguments against MRtb are the costs (77.7%) or a lack of technical infrastructure (18.1%; Table 2). Consequently, most urologists did not recommend MRtb for every patient but rather in particular patients such as those after a previous negative biopsy (Table 2). Furthermore, systematic biopsies in addition to lesiontargeted biopsies were not considered as negligible since most participants agreed in their judgment that MRtb does not eliminate the need for systematic biopsies (91%).

On multivariate logistic regression analysis, practice in a clinic (OR 1.852 [1.221; 2.808]; p = 0.004) and a regular

Table 2. Participants' responses to survey questions

How do you rate the use of mpMRI for prostate cancer diagnosis ($n = 468$)	
Very – extremely beneficial	340 (72.65%)
Disadvantage – moderately beneficial	128 (27.35%)
Do you use mpMRI for prostate cancer diagnosis ($n = 467$)	
Yes	448 (95.93%)
No	19 (4.07%)
Which system is used at your practice/institution to report the suspicion of prostate cancer on prostate $mpMRI$? ^a ($n = 462$)	
Likert	5 (1.08%)
PI-RADS v1	74 (16.02%)
Pi-RADS v2	390 (84.42%)
None of these	20 (4.33%)
Maximal number of lesions indicated in a prostate MRI ($n = 457$)	
<3	67 (14.66%)
3	197 (43.11%)
>3	193 (42.23%)
How many biopsies taken per lesion? $(n = 466)$	
1–2	210 (45.06%)
3–4	232 (49.79%)
≥5	24 (5.15%)
How beneficial is MRI targeted biopsy for prostate cancer detection? ($n = 466$)	
Extremely – very useful	333 (71.46%)
Not – moderately useful	133 (28.54%)
Would you recommend the use of MRI targeted biopsy for prostate cancer detection? $(n = 465)$	
Yes	387 (83.23%)
No	78 (16.77%)
Is MRI targeted biopsy better than purely systemic biopsy to detect significant cancer? ($n = 468$)	(
Yes	266 (56.84%)
No	34 (7.26%)
Sometimes	167 (35.68%)
Do not know	1 (0.22%)
Patient selection for MRI ^a $(n = 496)$	201 (40 520/)
Patients with no prior biopsy	201 (40.52%)
Patients with prior negative biopsy	451 (90.93%)
Patients on active surveillance	302 (60.89%)
Staging before radical prostatectomy	78 (15.73%)
Reasons for not using prostate MRI^a ($n = 304$)	250 (04 070/)
Cost Lack of radiology infrastructure	258 (84.87%)
	53 (17.43%)
Lack of expertise It's no use	20 (6.58%) 39 (12.83%)
Technique you use for MR targeted biopsy ^a ($n = 466$)	37 (12.0370)
Cognitive fusion $(n - 400)$	142 (25.97%)
"In bore" MRI biopsy	21 (4.50%)
MRI-ultrasound fusion biopsy	353 (75.75%)
Not applicable	29 (6.22%)
Patient selection for MR targeted biopsy ^a ($n = 467$)	25 (0.2270)
Patients with no prior biopsy	180 (38.54%)
Patients with prior negative biopsy	422 (90.36%)
Patients on active surveillance	255 (54.60%)
Patients prior to focal therapy	102 (21.84%)
Do not use it at all	34 (7.28%)
Reasons for not using MR targeted biopsy ^a ($n = 282$)	01 (7.2070)
Cost	219 (77.66%)
Lack of radiology infrastructure	51 (18.09%)
Lack of expertise	30 (10.64%)
It's no use	34 (12.06%)
	- (12.0070)

Table 2 (continued)

Believe in sufficient data for MR targeted biopsy use $(n = 459)$	()
Standard for all patients	85 (18.52%)
For particular patients such as prior negative biopsy	362 (78.87%)
Should not be performed	12 (2.61%)
Believe MR targeted biopsy eliminates the need for systematic biopsies? $(n = 465)$	
Yes	38 (8.17%)
No	423 (90.97%)
Do not know	4 (0.86%)
Believe ANNAcTRUS with targeted biopsy of suspicious lesions is a meaningful supplement to systemic biopsies? $(n = 463)$	TRUS
Yes	53 (11.45%)
For particular patients	115 (24.83%)
No	210 (45.36%)
Do not know	85 (18.36%)
Believe elastography is a meaningful supplement to TRUS ultrasound for the detection of suspic lesions? $(n = 463)$	cious
Yes	28 (6.05%)
For particular patients	78 (16.85%)
No	296 (63.93%)
Do not know	61 (13.17%)

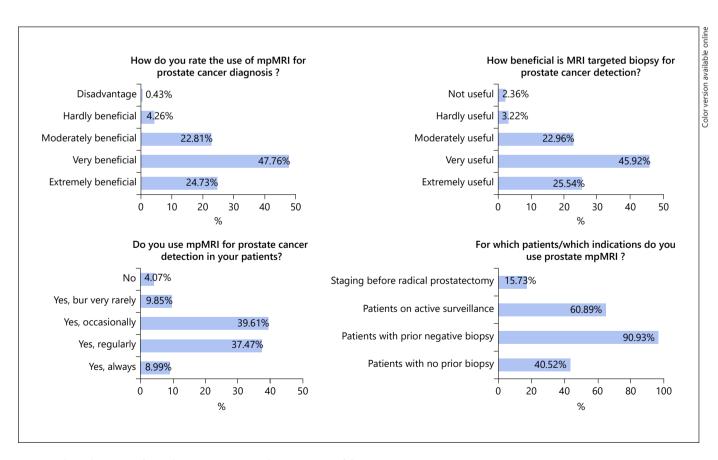


Fig. 2. Selected answers from the participants to the questions of the survey.

Table 3. Participants' turnaround time for prostate biopsy and MRI

Waiting period for pathological pr	ostate biopsy results ($n = 466$)				
1–3 days					
1 week	180 (38.63%)				
1–2 weeks	55 (11.80%)				
>2 weeks	2 (0.43%)				
Waiting period for a prostate MRI	appointment ($n = 455$)				
<1 week	16 (3.52%)				
1–2 weeks	88 (19.34%)				
2-4 weeks	219 (48.13%)				
>4 weeks	116 (25.49%)				
Do not use MRI	16 (3.52%)				

biopsy core number >12 cores (OR 2.219 [1.411; 3.488]; p = 0.001) were significantly associated with frequent utilization of prostate mpMRI for PCa diagnosis (Table 4). While an urban setting with >100,000 citizens was significant in univariate analysis (OR 1.561 [1.079; 2.258]; p = 0.018), it was not found to be an independent predictor on multivariate analysis (1.388 [0.949; 2.029]; p =0.091). Furthermore, practice in a clinic (OR 1.820 [1.138; 2.912]; p = 0.012) was independently associated with considering MRtb useful in practice (Table 4), and the recommendation for MRtb significantly emerged from practice in a clinic (OR 1.775 [1.069; 2.948]; p = 0.027). Interestingly, other techniques such as elastography or ANNAcTRUS were mostly not considered as a meaningful supplement to TRUS ultrasound with ANNAcTRUS as the most likely alternative for particular patients. Noteworthy, most participants waited <1 week for pathological results (87.8%; Table 3).

Discussion

There is a large discrepancy between the incidence and mortality of PCa [5]. Prostate biopsy has always aimed to detect clinically significant PCa, while "clinically significant" has been defined as a cancer that is likely to progress and that will therefore affect the life expectancy of a particular patient. For decades, systematic ultrasound-guided transrectal prostate biopsy has been the gold standard for a further diagnostic workup of patients suspected to have PCa based on a positive digital rectal exam or an elevated prostate-specific antigen. However, systematic TRUS-guided biopsies suffer from overdiagnosis and overtreatment on one hand, but also from underdiagnosis and undertreatment owing to the poor visibility of PCa

in b-mode ultrasound images as well as the poor accessibility of certain prostate areas (e.g., anterior) through the transrectal route on the other hand. At this point, mpMRI with its detailed morphological and functional information has changed our clinical algorithm. Since 2012, the PI-RADS standard of imaging and reporting prostate MRIs was introduced and improved the detection of clinically significant PCa with a further update in 2019 [6]. Of note, the majority of participants in our survey (84.4%) already use the PI-RADS v2 classification to report prostate MRIs, which is in line with another recent survey among urologists and radiologists [7].

Our survey was designed to assess the use of mpMRI and MRtb and more specifically their implementation in urologists' daily practice for the diagnosis and staging of PCa in Germany. Not surprisingly, mpMRI was proved to be a valuable, widely used tool as 95.9% of the participants use mpMRI of the prostate at some point and 83.2% also recommend MRtb. There have been only few surveys, predominantly in the US, indicating a similarly high number of urologists to utilize prostate mpMRI imaging in their daily practice (85.7–86%) but a lower number of urologists considering MRtb (60-63%) [8, 9]. Our study might report a higher acceptance for and application of MRtb since this technique needs some sophisticated equipment and infrastructure and therefore its broader application takes time. A 2013 national survey showed a use of only 34% for cognitive or ultrasound fusion guided biopsies indicating a strong trend in favor of this technique over time.

The widespread use of mpMRI and MRtb in the diagnosis and management of PCa is also supported by its 60.9 and 54.6% use in AS patients in our survey. While evidence has mostly been generated retrospectively, an initial analysis concluded that stable findings on mpMRI were predictive of stable Gleason score resulting in fewer biopsies for patients on AS [10]. Furthermore, mpMRI at the time of AS enrollment reduces the misclassification rate of csPCa, and therefore MRtb should be performed in patients with PI-RADS \geq 3 [11]. Consequently, the German S3 guideline recommends the use of mpMRI for AS patients [12], and therefrom, application of mpMRI might further increase in Germany for these particular patients. Adding to this, our study again confirms an increasing use of mpMRI with MRtb to improve urologists' confidence recommending AS [13].

Interestingly, several environmental factors show a significant impact on use of mpMRI for PCa diagnosis. Not only the majority of participants lives in cities with >100,000 citizens (54.4%), but such a setting is also pre-

Table 4. Uni- and multivariate logistic regression analysis

Independent characteristics	Univariate regression as	Univariate regression analysis		Multivariate regression analysis	
	OR [upper; lower KI]	p value	OR [upper; lower KI]	p value	
Use of mpMRI for prostate cancer diagnosis (frequent	tly or always versus occasionally or	never)			
Age (≥50 vs. <50 years)	1.158 [0.804; 1.667]	0.431			
Gender (F vs. M)	0.881 [0.543; 1.432]	0.610			
Region (West vs. East)	0.910 [0.515; 1.606]	0.744			
Setting (≥100,000 vs. <100,000 citizens) ^a	1.561 [1.079; 2.258]	0.018	1.388 [0.949; 2.029]	0.091	
Practice type (clinic vs. medical practice) ^b	1.932 [1.335; 2.797]	< 0.001	1.852 [1.221; 2.808]	0.004	
Trained urologist (yes vs. no)	1.014 [0.533; 1.928]	0.967			
Experience (>20 vs. ≤20 years)	1.258 [0.873; 1.815]	0.218			
Self-biopsy (yes vs. no)	1.032 [0.452; 2.353]	0.941			
Core numbers per patient (>12 vs. ≤12) ^c	2.468 [1.594; 3.821]	< 0.001	2.219 [1.411; 3.488]	0.001	
How beneficial is MR targeted biopsy for the diagnosi	is of prostate cancer (extremely – v	ery useful vs.	not – moderately useful)		
Age $(\geq 50 \text{ vs.} < 50 \text{ years})^d$	0.592 [0.386; 0.909]	0.017	0.587 [0.278; 1.240]	0.163	
Gender (F vs. M)	1.729 [0.928; 3.222]	0.084			
Region (West vs. East)	1.192 [0.627; 2.264]	0.592			
Setting (≥100,000 vs. <100,000 citizens)	1.363 [0.894; 2.081]	0.150			
Practice type (clinic vs. medical practice) ^e	2.168 [1.393; 3.375]	0.001	1.820 [1.138; 2.912]	0.012	
Trained urologist (yes vs. no) ^f	0.298 [0.104; 0.859]	0.025	0.633 [0.196; 2.046]	0.445	
Experience (>20 vs. ≤20 years)	0.695 [0.455; 1.060]	0.091			
Self-biopsy (yes vs. no)	1.005 [0.386; 2.612]	0.992			
Core numbers per patient (>12 vs. ≤12) ^c	1.559 [0.930; 2.613]	0.092			
Would you recommend MR targeted biopsy for prosta	ate cancer diagnosis? (yes vs. no)				
Age (≥ 50 vs. < 50 years)	0.646 [0.394; 1.060]	0.084			
Gender (F vs. M)	2.019 [0.930; 4.383]	0.076			
Region (West vs. East)	1.136 [0.545; 2.376]	0.733			
Setting (≥100,000 vs. <100,000 citizens)	1.207 [0.739; 1.969]	0.452			
Practice type (clinic vs. medical practice) ^g	1.775 [1.069; 2.948]	0.027	1.775 [1.069; 2.948]	0.027	
Trained urologist (yes vs. no)	0.642 [0.244; 1.690]	0.370			
Experience (>20 vs. ≤20 years)	0.759 [0.466; 1.236]	0.268			
Self-biopsy (yes vs. no)	0.457 [0.105; 1.992]	0.297			
Core numbers per patient (>12 vs. ≤12)	1.490 [0.811; 2.737]	0.198			

^a Multivariate model adjusted for practice type. ^b Multivariate model adjusted for age and core numbers. ^c Multivariate model adjusted for practice type, trained urologist, and experience. ^e Multivariate model adjusted for age and trained urologist. ^f Multivariate model adjusted for age, gender, practice type, and experience. ^g No confounders to be adjusted for.

dictive for an increased use of prostate mpMRI in a univariate analysis (p = 0.018). This may be due to a greater availability and higher technical standard in urban agglomerations and might also reflect the technical interest of respondents from such areas. Appropriately, urologists from clinical centers demonstrate a greater use of mpMRI and MRtb compared to urologists from medical practices (p = 0.004). These results suggest that short access routes to an mpMRI and greater resources influence decision making in practice since only 36.9% of those in medical practice routinely perform mpMRI/MRtb compared with 52.2% in clinical centers and 67.8% at a university hospi-

tal. Furthermore, those in academic centers tend to perform more biopsies and radical prostatectomies and therefore more often utilize MRtb in their practice [8].

Environmental and institutional factors are also reflected as the most common barriers for implementation of MRtb, namely, costs (77.7%) and linkage with radiology (18.1%). Interestingly, costs are a nearly as large constraint in the clinic (44.7%) as in urology practice (58.2%) probably reflecting the insufficient cost-recovery for mpMRI in daily practice by public as well as private health insurances. Additionally, users have to overcome the higher upfront costs to purchase a fusion prostate biopsy

device or implement advanced imaging in their daily practice, which additionally requires availability of an expert radiologist or the confidence in reading of prostate mpMRI in community offices. Certainly, such constraints still affect attitudes toward the implementation and use of this technology.

While mpMRI/MRtb is seen as supportive and helpful for the diagnosis of PCa and attitudes toward its use are generally positive, there are some reservations about a universal use for all patients. The majority of respondents believe that mpMRI should be reserved for selected men such as those with a prior negative biopsy (90.9%) or those under AS (60.9%). This recommendation is advised in many guidelines [12, 14] and reflects the proven effectiveness of prostate mpMRI and MRtb in these clinical situations [2, 15]. Therefrom, our data suggest that urologists support mpMRI and MRtb for complex patients.

However, the question arises if mpMRI should be performed for all men prior to biopsy to reduce unnecessary interventions, to reduce overdiagnosis of clinically insignificant PCa, and to improve the detection of clinically significant PCa. Only 40.5% of our respondents would recommend a prostate mpMRI for patients with no prior biopsy, but this is already a large matter of debate. The PROMIS trial included >500 biopsy-naïve men and was able to show an avoidance of primary biopsy in 27% of the patients with the use of mpMRI when used as a triage test before first biopsy [16]. Furthermore, the PRECISION trial reported that prostate mpMRI prior to first biopsy was superior to standard transrectal ultrasound-guided prostate biopsy in men at clinical risk of PCa [4]. Additionally, the question if such an mpMRI first strategy is cost effective has been answered in the PROMIS trial, in which this workflow was shown to be clinically effective and cost effective for the diagnosis of significant PCa [17]. In a position paper on population-based PCa screening, the European Association of Urology (EAU) recommends mpMRI as a test to select biopsy candidates in men at risk of significant PCa according to prostate-specific antigen levels [18]. However, there is at the moment only a weak recommendation in the EAU guidelines to perform mpMRI before first prostate biopsy [14]. The negative predictive value of mpMRI for PCa remains an important factor and is rated - depending on the patient selection – between 63 and 98% [19–21]. Overall, mpMRI as a triage test could detect more significant PCa in 40% of men after prior negative biopsy and in 5% of biopsynaïve men compared to TRUS-guided biopsy only. Nevertheless, there is a strong recommendation to combine

targeted and systematic biopsies as soon as an mpMRI is positive with a PI-RADS \geq 3 lesion.

Even though systematic biopsies detect more insignificant and less significant cancers and targeted biopsies detect more significant and less insignificant PCa [2, 22], the combination of targeted and systematic biopsies detects more clinically significant PCa than either modality alone irrespective of a first line approach or in patients with previously negative biopsy [23–25]. The majority of our respondents kept that data in mind and consequently indicated that MRtb does not eliminate the need for systematic biopsies (91%). However, concerning lesion-targeted biopsies, there seems to be a clear limit for the number of cores per lesion. Most respondents would take a maximum of 4 biopsies per lesion (94.9%), which is in line with the average number of biopsies per lesion as well as the recommended number of biopsies in larger studies [4, 26].

As expected, MRI-ultrasound fusion biopsy is used from the majority of respondents (75.8%), while cognitive fusion is still appropriate for 26% and in bore MRI biopsies are rarely used (4.5%). While all these 3 techniques enhance detection rates for clinically significant PCas compared to standard TRUS-guided biopsies [27], none of these is clearly superior to the others in their detection rate.

There are several limitations in this study. First, the survey design that addressed the members of the German association of urology via email has to be mentioned. This has some reporting bias with a lack of information on candidates that did not respond. A majority of respondents might be more interested in mpMRI and new techniques, therefore being appealed by the technological topic of the survey and its electronic design. Therefrom, we cannot totally generalize the results to the overall urologic community and all members of the German Society of Urology. Also, prostate MRI remains an evolving field, with frequent publication of new trials. Our survey asked urologists' opinions at a single point in time and results could be subject to change with newer results and expert opinions to be published. Finally, this survey represents an assessment of urologists' preferences not necessarily including other disciplines.

Conclusion

Our data are the first to show the meanwhile high acceptance and clinical use of the prostate mpMRI and MRtb in Germany. Urologists regard mpMRI and MRtb as improving risk stratification for localized PCa and increasing confidence in using AS. Clinic centers report a

greater utilization of mpMRI and MRtb than medical practice. However, mpMRI is still not available for all urologists with an excellent image quality and a dedication and experience of radiologic centers. Hence, close collaboration is desirable and will ensure direct interdisciplinary communication between radiologists and urologists while modified mpMRI protocols might be necessary to further reduce costs and increase availability.

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Statement of Ethics

All data were collected according to the General Data Protection regulation of the EU. This study was conducted in accordance with the World Medical Association Declaration of Helsinki. The

survey does not involve studies of human or animal subjects. Analyzed data were completely anonymized with rigorous data protection measures.

Conflict of Interest Statement

The authors declare that they have no conflicts of interests to declare

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No funding was received for this study.

Author Contributions

M.S.: project development, data collection, data analysis and interpretation, manuscript writing, and critical revision. J.L.: project development, data collection, data analysis and interpretation, and critical revision. G.K. and B.W: project development, data analysis and interpretation, manuscript writing, critical revision, and supervision. A.B., S.F., K.U., and L.B.: project development, data interpretation, language revision, and critical revision.

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