

Current practices and future directions in prostate biopsy techniques: insights from a meta-analysis and european multicenter survey

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Dr. Wu's recent systematic review and network meta-analysis—encompassing 211 studies and over 74,000 MRI-positive patients—establishes an evidence-based benchmark for prostate biopsy strategies.¹ Utilizing Bayesian methodologies, the study demonstrated that ipsilateral systematic biopsy combined with targeted biopsy (ips-SB+TB) or saturated targeted biopsy (saturated TB) achieves comparable detection rates of clinically significant prostate cancer (csPCa; Gleason $\geq 3 + 4$) to traditional combined biopsy (SB+TB), while significantly reducing detection of clinically insignificant disease (ciPCa; Gleason $\leq 3 + 3$). These findings challenge the conventional paradigm equating higher biopsy core numbers with improved diagnostic yield, proving that sampling within a 10–15 mm radius around MRI-suspicious lesions optimizes csPCa detection while mitigating overdiagnosis. Our group's 2024 prospective RCT validated this approach, demonstrating that 9-core regional saturation biopsy (RSB) using brachytherapy grid guidance improved csPCa detection (44.1%) while reducing biopsy cores by 40% compared to traditional methods, thus providing robust support for the EAU 2024 guideline strategy: “When MRI is positive (PI-RADS

≥ 4), combine targeted biopsy with perilesional sampling”* (Strength rating: Weak).²

Paradoxically, despite guideline recommendations, a European cross-sectional survey reveals a substantial implementation gap: 84% of clinicians continue to perform “MRI-targeted + bilateral systematic biopsy.” This discrepancy reflects persistent concerns regarding MRI's false-negative rate for csPCa (15–20%) and the indispensable role of systematic biopsy in multidisciplinary care—particularly for baseline data in active surveillance and neurovascular bundle mapping during surgery.³ This clinical reliance on systematic sampling underscores the tension between current technical limitations of MRI-TRUS fusion (registration errors >1.5 mm in 40% of grassroots institutions) and complex real-world diagnostic needs. Although targeted biopsy with perilesional sampling represents a precision-sampling advancement, its adoption faces significant barriers: ambiguous anatomical definitions of “surrounding areas” (e.g., 10 mm vs. 15 mm radii), absence of standardized fusion training, and limited evidence in high-risk subgroups (e.g., bilateral lesions, repeat biopsies).

Optimizing biopsy strategies requires addressing three critical barriers: Firstly, inconsistent precision and accessibility of fusion technology (grassroots freehand techniques yield 12–15% lower csPCa detection vs. grid-guided methods);⁴ Secondly, lack of consensus on zone-specific sampling protocols (inter-clinician diagnostic consistency: 68–75%); Thirdly, insufficient long-term pathological validation (grade

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upgrade rates: 18% regional vs. 22% traditional biopsy). Future efforts should prioritize multicenter collaborations to: Develop standardized biopsy templates integrating PI-RADS v2.1 and PSA density; Combine PSMA-PET/CT with mpMRI to enhance detection of small lesions (>3 mm; accuracy $\geq 90\%$); Leverage AI-driven needle trajectory planning for complex anatomies (e.g., prostate volume >80 mL).⁵ Achieving end-to-end standardization across imaging, procedural, and pathological domains is essential to transition from “experience-driven systematic sampling” to “evidence-driven precision diagnosis.” This evolution will minimize overdiagnosis while providing reliable pathological foundations for focal therapy and active surveillance.

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Author Contributions

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Availability of Data and Materials

Not applicable.

Ethics Approval

Not applicable.

Conflicts of Interest

The authors declare no conflicts of interest to report regarding the present study.

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