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Relationship of prostate cancer topography and tumour conspicuity on multiparametric magnetic resonance imaging: a systematic review and meta-analysis

Pranav Satish,^{1,2} Alex Freeman,³ Daniel Kelly,⁴ Alex Kirkham,⁵ Clement Orczyk,^{2,6} Benjamin S. Simpson,⁷ Francesco Giganti^{2,5}, Hayley C. Whitaker,² Mark Emberton,^{2,6*} Joseph M. Norris,^{2,6*}

¹ UCL School of Medicine, University College London, London, UK

² UCL Division of Surgery & Interventional Science, University College London, London, UK

³Department of Pathology, University College London Hospitals NHS Foundation Trust, London, UK

⁴ School of Healthcare Sciences, Cardiff University, Wales, UK

⁵Department of Radiology, University College London Hospitals NHS Foundation Trust, London, UK

⁶Department of Urology, University College London Hospitals NHS Foundation Trust, London, UK

⁷ UCL Cancer Institute, University College London, London, UK

* joint-senior authors

Introduction & Objectives

Multiparametric magnetic resonance imaging (mpMRI) has improved the triage of men with suspected prostate cancer. Multiple factors influence cancer conspicuity on mpMRI, however, the implications of tumour location are not fully understood. Identifying topographical correlates that influence mpMRI conspicuity may improve outcomes. Here, we present the first systematic review and meta-analysis describing the effect of tumour location on prostate cancer conspicuity on mpMRI.

Materials & Methods

We systematically searched Medline, PubMed, Embase and Cochrane databases up to January 2021, and results were assessed as per the PRISMA statement. Tumour conspicuity on mpMRI was compared between cancers in the peripheral zone (PZ), transitional zone (TZ), base, apex, anterior and posterior. Meta-analysis was conducted to compare the diagnostic odds ratios (DOR) of detection for tumours in the PZ and TZ. Five studies had suitable data, so a random-effects model was fit and DOR compared. Risk of bias was measured via a modified Newcastle-Ottawa score (NOS). The review was prospectively registered with PROSPERO: CRD42021228087.

Results

We screened 960 articles, 17 of which fulfilled the inclusion criteria. Our NOS deemed the quality of inclusions was generally high. Thematic synthesis showed apical and basal tumours had reduced conspicuity on mpMRI, whilst mid-gland tumours had higher detection rates. TZ cancer displayed increased conspicuity on T2-weighted imaging, whilst PZ cancers were higher on diffusion-weighted imaging and dynamic contrast enhancement. With few exceptions, mpMRI had better overall diagnostic accuracy for PZ lesions, albeit higher specificity for TZ lesions. Meta-analysis showed an

increased DOR for PZ tumours on mpMRI (OR:9.7 [95% CI:4.0-23], compared to TZ (OR:5.2 [95% CI:2.6-10]). However, no significant differences between subgroups were observed (p=0.27). The model showed 38% heterogeneity (I2 =38%, p=0.10).

Study	Experin	nental Total	Events	ontrol Total	Odds Ratio	OR	95%-CI	Weight (fixed)	Weight (random)
Subaroup = D7					1 6				
Affi	22	43	0	11	· · · · ·	72 20	12 09- 1252 951	0.2%	2 204
Cosma	53	65	17	51		8.83	[3.30, 1333.03]	5.5%	12.5%
Reisaeter	171	311	151	823	ri.	5 44	[4 09 7 23]	58 5%	16.6%
Ito	66	132	3	79		25.33	17 61 84 371	2.9%	9.8%
Schimoeller	91	95	98	119		4.88	[1.61: 14.74]	5.8%	10.5%
Fixed effect model		646		1083	•	6.72	[5.23: 8.65]	73.1%	
Random effects mode	1					9.66	[3.99; 23.43]		52.7%
Heterogeneity: $l^2 = 59\%$, t	2 = 0.6746	s, p = 0	.05						
Subgroup = TZ									
Afifi	6	8	0	4		23.40	[0.89; 612.98]	0.3%	2.6%
Cosma	44	57	38	113		6.68	[3.21; 13.88]	9.1%	13.5%
Reisaeter	10	54	30	512		3.65	[1.67; 7.96]	7.3%	13.1%
Ito	37	52	2	11	- <u></u>	11.10	[2.14; 57.53]	1.5%	7.2%
Schimoeller	54	58	244	294	- * 1	2.77	[0.96; 7.99]	8.7%	10.9%
Fixed effect model		229		934	-	5.01	[3.14; 7.99]	26.9%	
Random effects mode Heterogeneity: $l^2 = 4\%$, τ^2	= 0.2678,	p = 0.3	38		-	5.16	[2.57; 10.33]		47.3%
Fixed offect model		075		2017		6.26	15.02 7.021	100.0%	
Pandom offecte mode		015		2017	1 1	7 16	[4 03: 12 72]	100.0%	100.0%
Hataraganahu /2 = 20%	2 - 0 4076	0 - 0	10			1.10	[4.03, 12.72]	-	100.0%
z = 16.19 (p < 0.001) z = 6.71 (p < 0.001)	- 0.4875	o, p = 0	. 10	0.	001 0.1 1 10 10	00			

Conclusions

Tumour location appears to influence prostate cancer conspicuity on mpMRI. Apical or basal tumours may be less conspicuous on mpMRI than mid-gland tumours. Similarly, TZ cancer appears to have reduced conspicuity compared to PZ cancer, however, meta-analysis did not show a significant difference in DOR. Future studies with robust, prospective data-sets are required to clarify the relationship between tumour position and conspicuity.