

CURRENT TRENDS IN MRI USE IN PROSTATE CANCER

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ACCRA, GHANA

The World Association of Radiopharmaceutical and Molecular Therapy (WARMTH) is excited to invite you to join us for the will be held 18th International Conference on Radiopharmaceutical Therapy (ICRT) Accra, Ghana, 2nd-4th May 2023,



OUTLINE

- OBJECTIVES
- REVIEW OF MRI ANATOMY
- APPROACHES TO DIAGNOSING PROSTATE CANCER
- MULTIPARAMETRIC MRI
- PIRADS CLASSIFICATION
- WHOLE BODY MRI
- CONCLUSION



OBJECTIVES

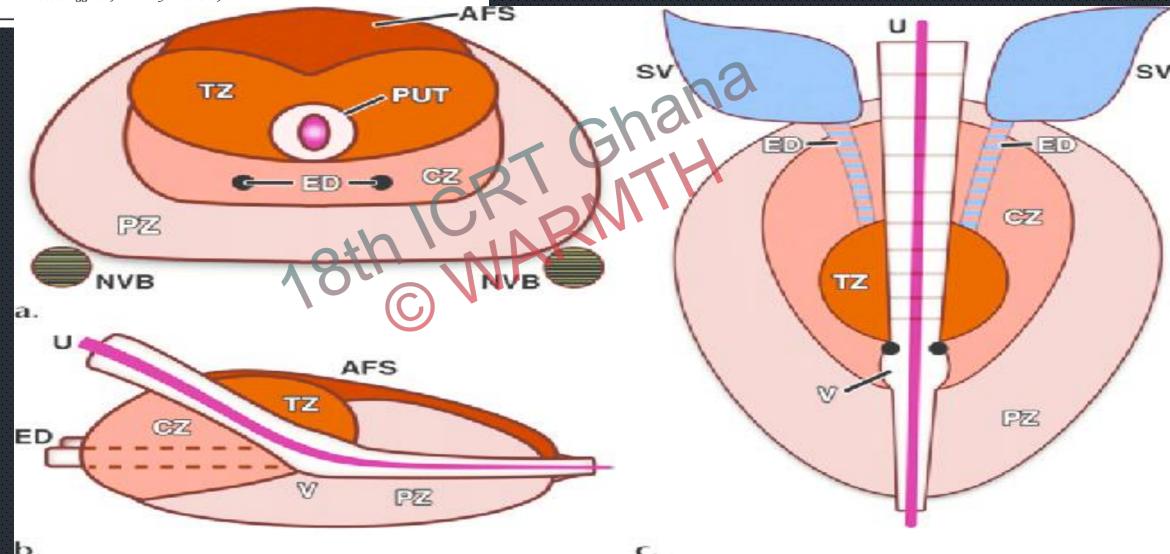
• PARTICIPANTS WILL BE ABLE TO LIST THE STRENGTHS AND WEAKNESSES OF THE MAIN DIAGNOSTIC IMAGING MODALITIES USED IN PROSTATE CANCER MANAGEMENT

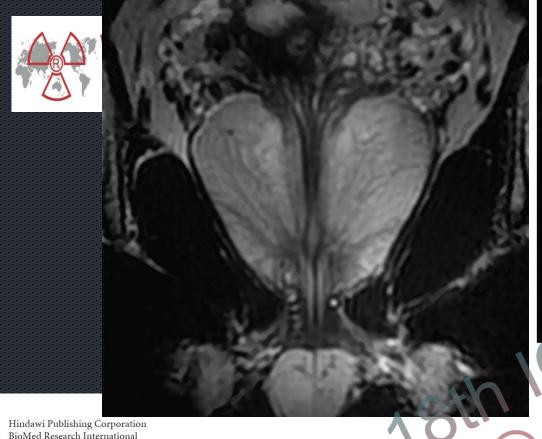
Imaging-guided Prostate Biopsy: Conventional and Emerging Techniques¹

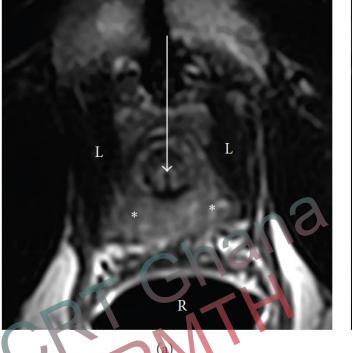
TEACHING POINTS See last page

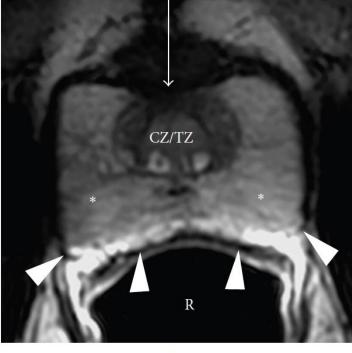
Joseph H.Yacoub, MD • Sadhna Verma, MD • Jonathan S. Moulton, MD
Scott Eggener, MD • Aytekin Oto, MD

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Hindawi

Review Article

Anatomic Imaging of the Prostate

Anil Bhavsar and Sadhna Verma

University of Cincinnati Medical Center, Cincinnati, OH 45267-0762, USA

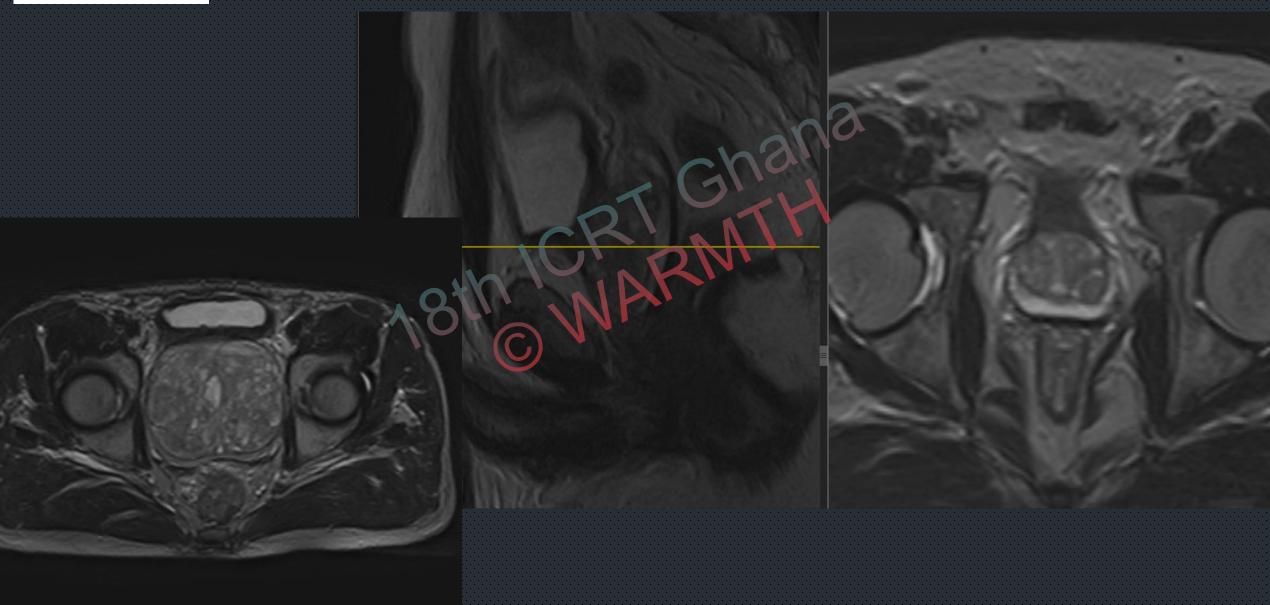
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PROSTATE CANCER DIAGNOSTIC PATHWAYS

- DIGITAL RECTAL EXAMINATION (DRE)
- PROSTATE SPECIFIC ANTIGEN TEST (PSA)
- TRANS RECTAL ULTRASOUND BIOPSY (TRUS)- 27-44%-ROTHWAX ET AL
- MULTIPARAMETRIC-MRI MP-MRI/BIPARAMETRIC MRI -BP-MRI
- WHOLE BODY-MRI WB-MRI



PROSTATE CANCER (PCA) DIAGNOSTIC GOALS

- LOCALISE THE LESION WITH THE GLAND
- REDUCE DIAGNOSIS OF CLINICALLY INSIGNIFICANT PCA (CIPA)
- INCREASE DIAGNOSIS OF CLINICALLY SIGNIFICANT PCA (CSPA)
- ASSESS THE AGGRESSIVE NATURE OF THE LESION

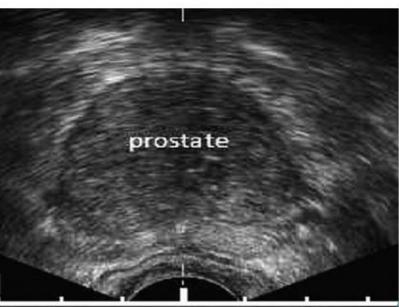


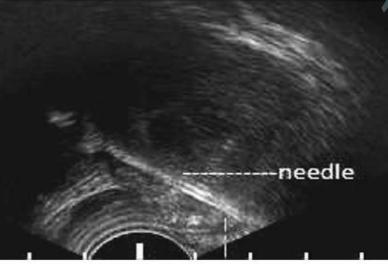
TRANS RECTAL ULTRASOUND BIOPSY (TRUS)

- ACCEPTED MODE OF BIOPSYING THE PROSTATE
- MOST COMMON METHOD OF BIOPSYING THE PROSTATE
- LOW COST COMPARED TO MP-MRI
- MAY BE RANDOM OR MRI-ASSISTED
- OFTEN 10-12 CORES ARE USUALLY TAKEN



TRANS RECTAL ULTRASOUND BIOPSY (TRUS)





Systematic Biopsy Schemes

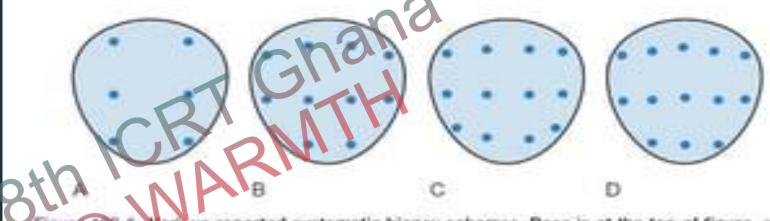
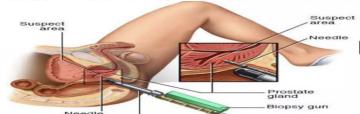


Figure 40%-5. Various reported systematic biopsy schemes. Base is at the top of figure, apex is at bottom. A, Sextant biopsy scheme originally proposed by Hodge and associates (Hodge et al, 1964). B, The 10-core biopsy (Presti et al, 2000). C, The 12-core, or double-sextant, biopsy. This is the currently recommended sequence endorsed by the American Urological Association (Bjurlin et al, 2013). D, The 13-core, 5-region biopsy (Eskew et al, 1997).



PROSTATE BIOPSY

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PMCID: PMC6828624

PMID: 31172275

The primacy of multiparametric MRI in men with suspected prostate cancer

Jonathan Richenberg,^{™1} Vibeke Løgager,² Valeria Panebianco,³ Olivier Rouviere,^{4,5} Geert Villeirs,⁶ and Ivo G. Schoots⁷

Introduction

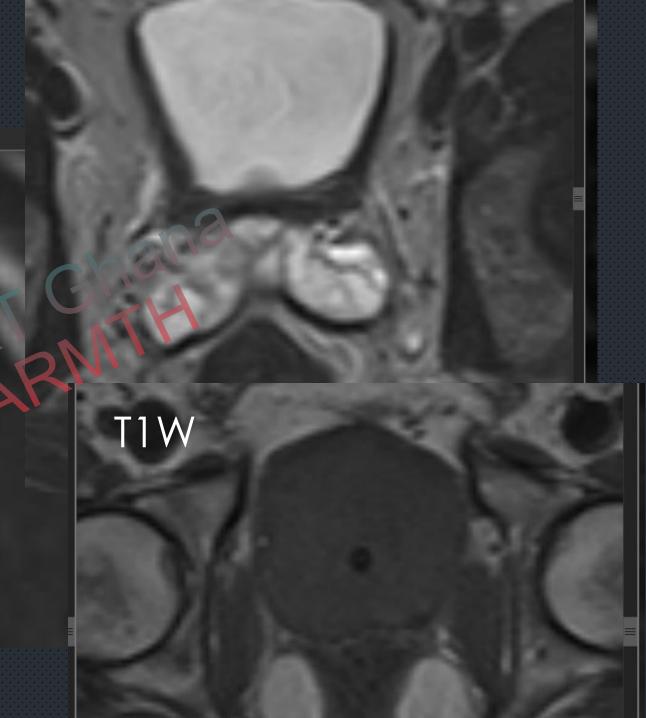
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In 2012, the European Society of Urogenital Radiology (ESUR) prostate committee promoted the use of multiparametric MRI (mpMRI) in the routine management of men with suspected or confirmed prostate cancer [1]. That proposal has gained widespread acceptance. The debate has now moved to when mpMRI should be used.



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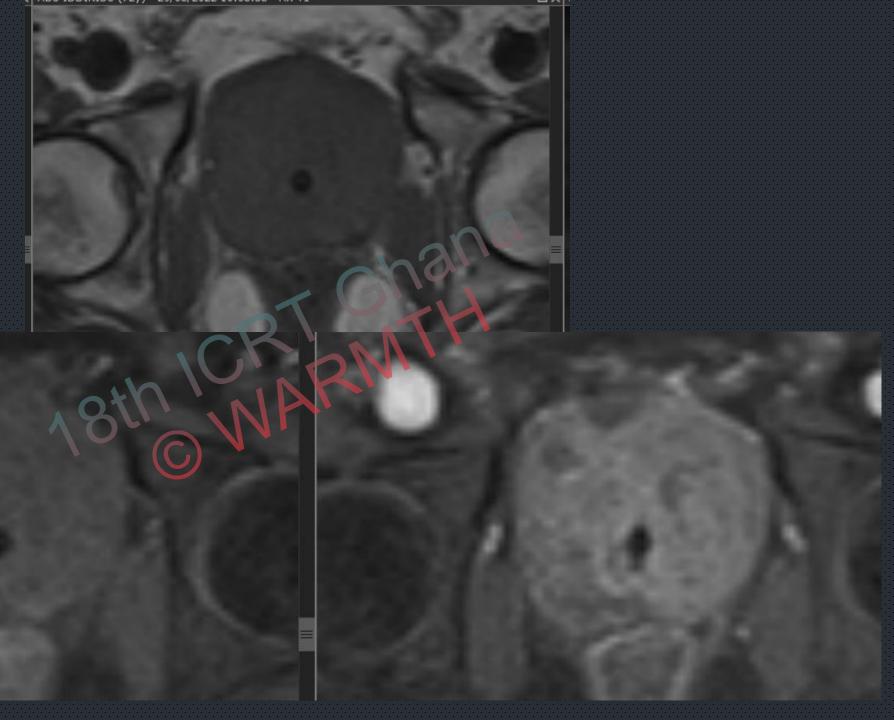




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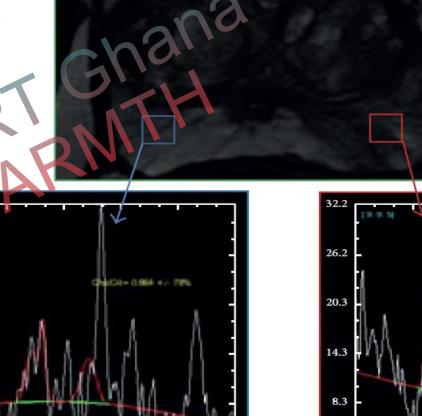
MRI SPECTROSCOPY

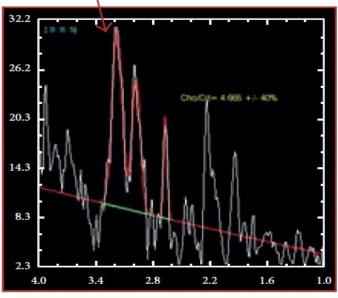
21.4

Review Article

Functional MRI in Prostate Cancer Detection

Sandeep Sankineni, Murat Osman, and Peter L. Choyke







PIRADS V2.1

PI-RADS™ v2.1 assessment uses a 5-point scale based on the likelihood (probability) that a combination of mpMRI findings on T2W, DWI, and DCE correlates with the presence of a clinically significant cancer for each lesion in the prostate gland.

PI-RADS™ v2.1 Assessment Categories

PIRADS 1 – Very low (clinically significant cancer is highly unlikely to be present)

PIRADS 2 – Low (clinically significant cancer is unlikely to be present)

PIRADS 3 – Intermediate (the presence of clinically significant cancer is equivocal)

PIRADS 4 – High (clinically significant cancer is likely to be present)

PIRADS 5 – Very high (clinically significant cancer is highly likely to be present)

PI-RADS®

Prostate Imaging – Reportinand Data System

2019 Version 2.1







2. PI-RADS Assessment for T2W

Score	Peripheral Zone (PZ)
1	Uniform hyperintense signal intensity (normal)
2	Linear or wedge-shaped hypointensity or diffuse mild hypointensity, usually indistinct margin
3	Heterogeneous signal intensity or non-circumscribed, rounded, moderate hypointensity Includes others that do not qualify as 2 4, or 5
4	Circumscribed, homogenous moderate hypointense focus/mass confined to prostate and <1.5 cm in greatest dimension
5	Same as 4 but ≥1.5cm in greatest dimension or definite extraprostatic extension/invasive behavior
Score	Transition Zone (TZ)
1	Normal appearing TZ (rare) or a round, completely encapsulated nodule. ("typical nodule")
2	A mostly encapsulated nodule OR a homogeneous circumscribed nodule without encapsulation. ("atypical nodule") OR a homogeneous mildly hypointense area between nodules
3	Heterogeneous signal intensity with obscured margins
	Includes others that do not qualify as 2, 4, or 5
4	Lenticular or non-circumscribed, homogeneous, moderately hypointense, and <1.5 cm in greatest dimension
5	Same as 4, but ≥1.5cm in greatest dimension or definite extraprostatic extension/invasive behavior



2. PI-RADS Assessment of DWI

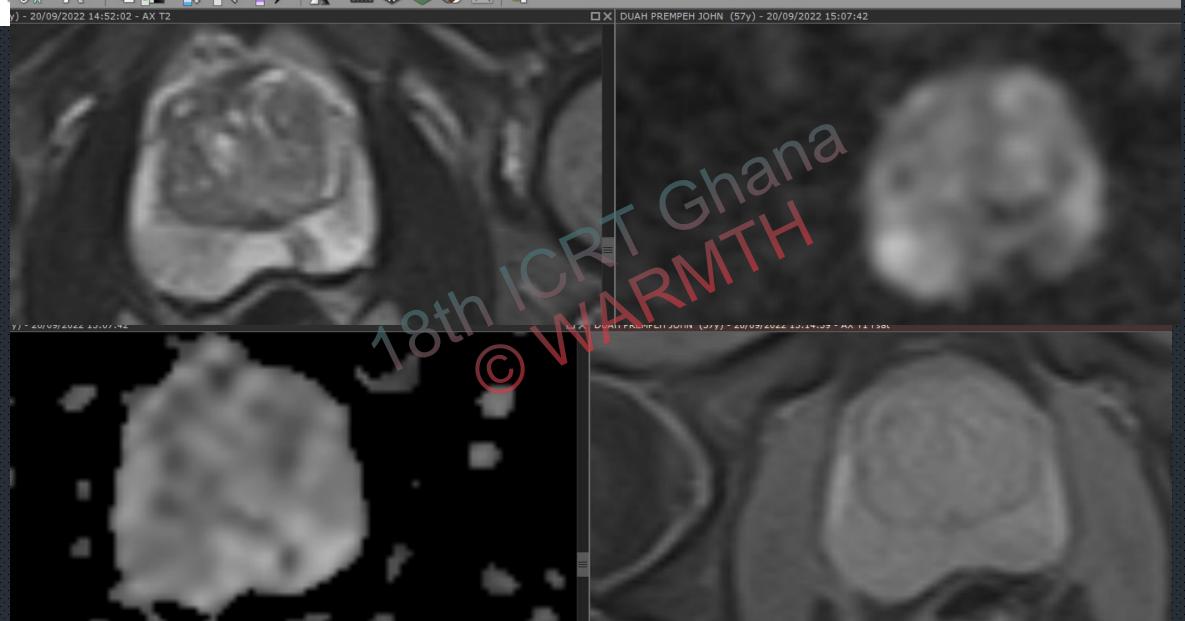
Signal intensity in a lesion should be visually compared to the average signal of "normal" prostate tissue in the histologic zone in which it is located.

Score	Peripheral Zone (PZ) or Transition Zone (TZ)
1	No abnormality (i.e., normal) on ADC and high b-value DWI
2	Linear/wedge shaped hypointense on ADC and/or linear/wedge shaped hyperintense on high b-value DWI
3	Focal (discrete and different from the background) hypointense on ADC and/or focal hyperintense on high b-value DWI; may be markedly hypointense on ADC or markedly hyperintense on high b-value DWI, but not both.
4	Focal markedly hypontense on ADC and markedly hyperintense on high b-value DWI; <1.5cm in greatest dimension
5	Same as 4 but ≥1.5cm in greatest dimension or definite extraprostatic extension/invasive behavior



2. PI-RADS Assessment for DCE

Score	Peripheral Zone (PZ) or Transition Zone (TZ)
(-)	no early or contemporaneous enhancement; or diffuse multifocal enhancement NOT corresponding to a focal finding on T2W and/or DWI or focal enhancement corresponding to a lesion demonstrating features of BPH on T2WI (including features of extruded BPH in the PZ)
(+)	focal, and; earlier than or contemporaneously with enhancement of adjacent normal prostatic tissues, and; corresponds to suspicious finding on T2W and/or DWI

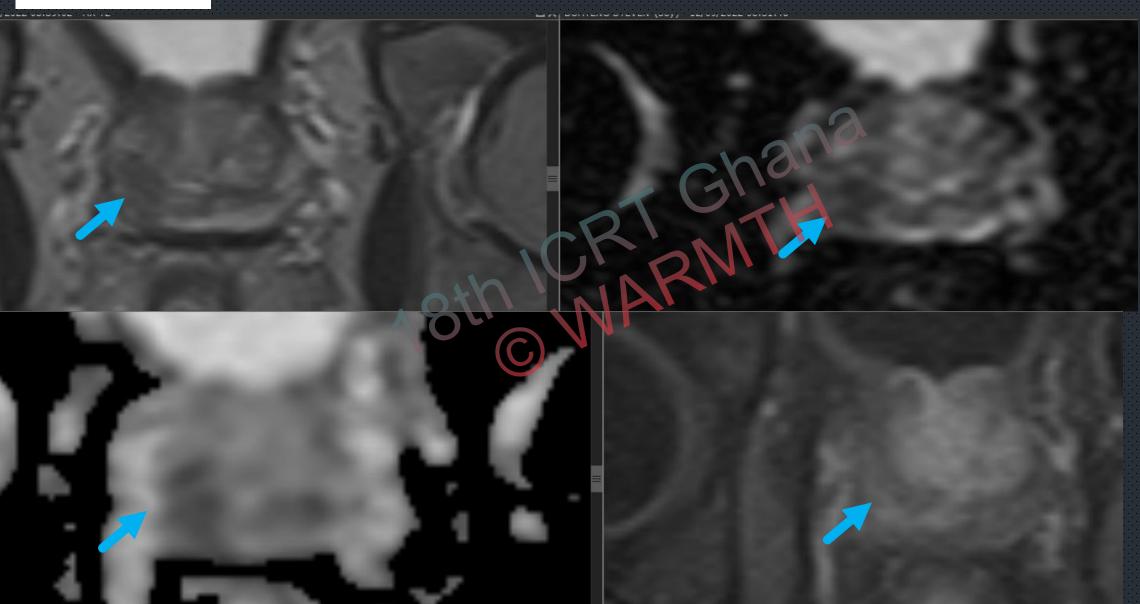




WARMTH WORLD ASSOCIATION OF RADIOPHARMACEUTICAL AND MOLECULAR THERAPY PIRA 5









BP-MRI

3. Commentary on bi-parametric MRI

Despite the limited role of DCE in determining the overall PI-RADS assessment category experience has shown that, in some instances, DCE may assist in detection of csPCa in both the PZ and TZ, and in clinical practice some have viewed DCE as a 'safety-net' or 'back-up' sequence, especially when DWI is degraded by artifacts or inadequate signal-to-noise ratio (SNR).

Given the limited role of DCE, there is growing interest in performing prostate MRI without DCE, a procedure termed "biparametric MRI" (bpMRI). A number of studies have reported data that supports the value of bpMRI for detection of csPCa in biopsy-naïve men and those with a prior negative biopsy.

The PI-RADS Steering Committee supports continued research concerning the performance of bpMRI in various clinical scenarios and acknowledges the potential benefits, including: (1) elimination of adverse events and gadolinium retention that have been associated with some gadolinium based contrast agents (GBCAs), (2) shortened examination times, and (3) reduced costs, possibly resulting in increased accessibility and utilization of MRI for biopsy-naïve men with suspected prostate cancer.



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Platinum Priority – Prostate Cancer

Editorial by Jelle O. Barentsz, Peter Mulders, Winald Cerritsen and Jurgen J. Fütterer on pp. 93–95 of this issue

METastasis Reporting and Data System for Prostate Cancer: Practical Guidelines for Acquisition, Interpretation, and Reporting of Whole-body Magnetic Resonance Imaging-based Evaluations of Multiorgan Involvement in Advanced Prostate Cancer

Anwar R. Padhani^{a,*}, Frederic E. Lecouvet^b, Nina Tunariu^c, Dow-Mu Koh^c, Frederik De Keyzer^d, David J. Collins^c, Evis Sala^e, Heinz Peter Schlemmer^f, Giuseppe Petralia^g, H. Alberto Vargas^e, Stefano Fanti^h, H. Bertrand Tombalⁱ, Johann de Bono^j



Pricolo et al. Cancer Imaging (2020) 20:77 https://doi.org/10.1186/s40644-020-00350-x

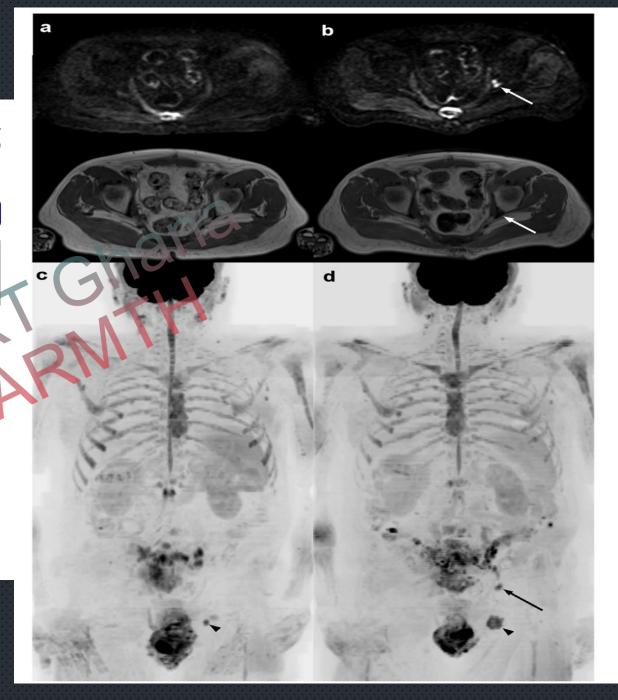
Cancer Imaging

RESEARCH ARTICLE

Open Access

Whole-body magnetic resonance imaging (WB-MRI) reporting with the METastasis Reporting and Data System for Prostate Cancer (MET-RADS-P): inter-observer agreement between readers of different expertise levels

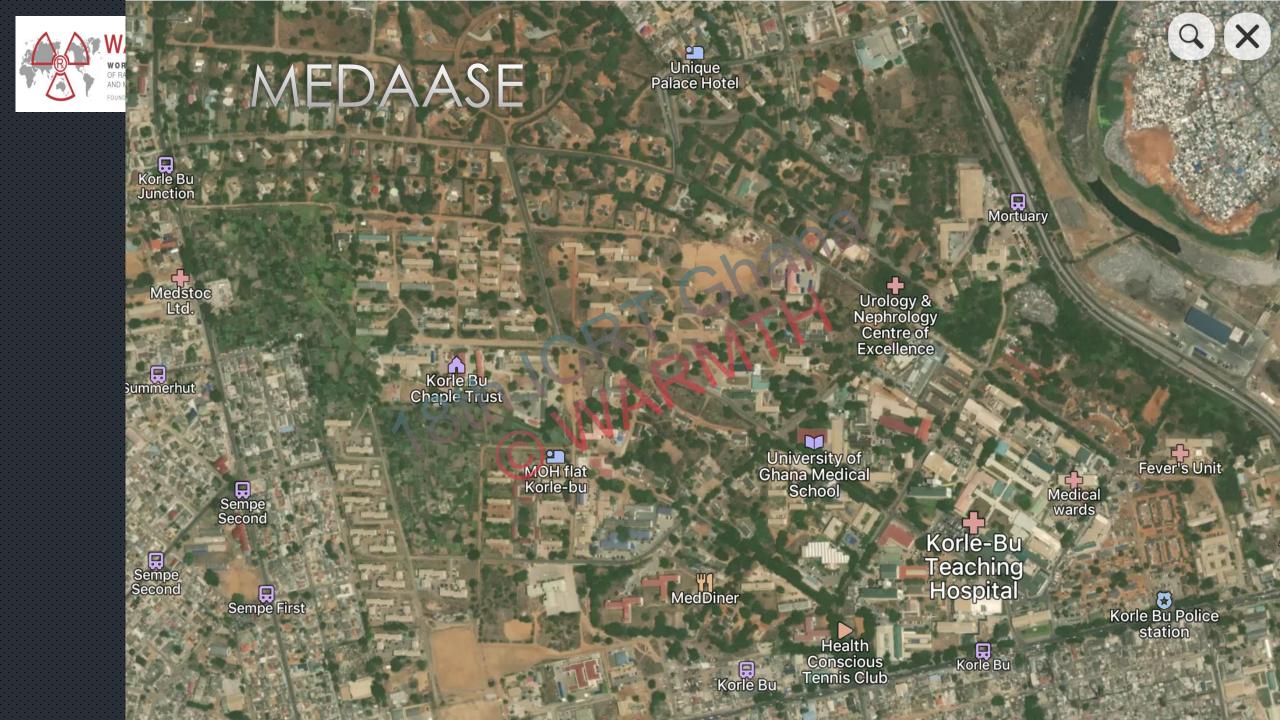
Paola Pricolo¹, Eleonora Ancona^{1,2}, Paul Summers¹, Jorge Abreu-Gomez³, Sarah Alessi¹, Barbara Alicja Jereczek-Fossa^{4,5}, Ottavio De Cobelli^{5,6}, Franco Nolè⁷, Giuseppe Renne⁸, Massimo Bellomi^{1,5}, Anwar Roshanali Padhani⁹ and Giuseppe Petralia^{5,10*}





CONCLUSION

- IMAGING HAS HAD A SIGNIFICANT POSITIVE INFLUENCE ON THE MANAGEMENT OF PROSTATE CANCER—TRUS, MP-MRI
- THE ADDITION OF MP-MRI HAS SIGNIFICANTLY IMPROVED THE SENSITIVITY OF PCA DIAGNOSIS ALBEIT WITH VARYING DEGREES OF LIMITATIONS
- WB-MRI IS GRADUALLY PROVING TO BE VERY SENSITIVE TO METASTATIC LESIONS





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 PMCID: PMC6828624.