

10:15 – 10:45 Integrating Radiology and Oncology: A Multidisciplinary Approach

MRI in prostate cancer

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Thank you



Prof. Henrik S. Thomsen

A Multidisciplinary Approach

- What is MRI of the prostate ?
- Why ?
- How ?
- What is multidisciplinary in this context ?
- How do we do it ?
- What have we learned?
- Is it worth it ?
- Health economists ? Lifequality ? Survival ?
- Ask the patients ?

But you cannot say MRI for the prostata with out saying PI-RADS

- PI-RADS and Jelle Barentsz

Eur Radiol (2012) 22:746–757
DOI 10.1007/s00330-011-2377-y

UROGENITAL

ESUR prostate MR guidelines 2012

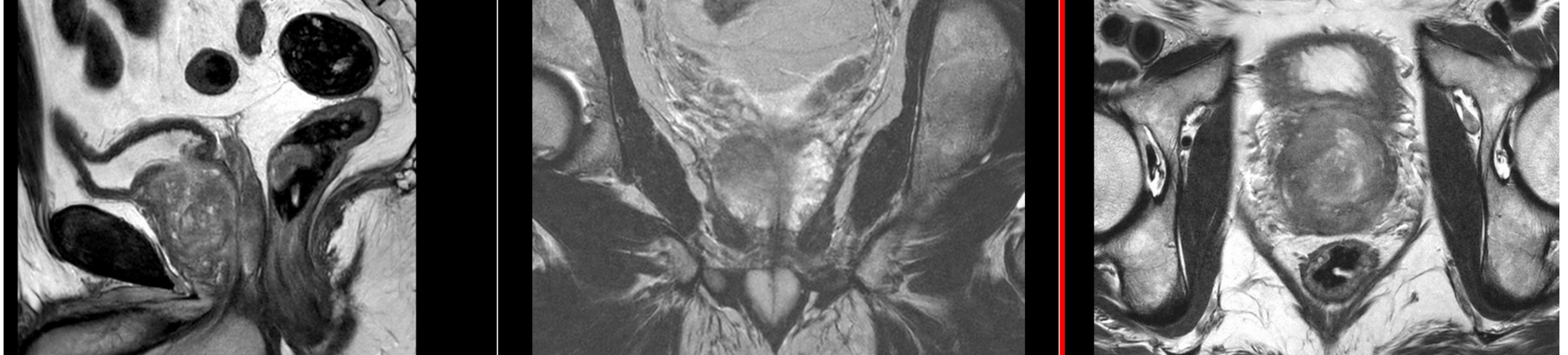
Jelle O. Barentsz • Jonathan Richenberg •
Richard Clements • Peter Choyke • Sadhna Verma •
Geert Villeirs • Olivier Rouviere • Vibeke Logager •
Jurgen J. Fütterer



5-point scoring system used by radiologists and urologists to evaluate prostate MRIs. It acts as a "common language" that scores the likelihood of clinically significant prostate cancer to be present.



This is MRI of the prostate

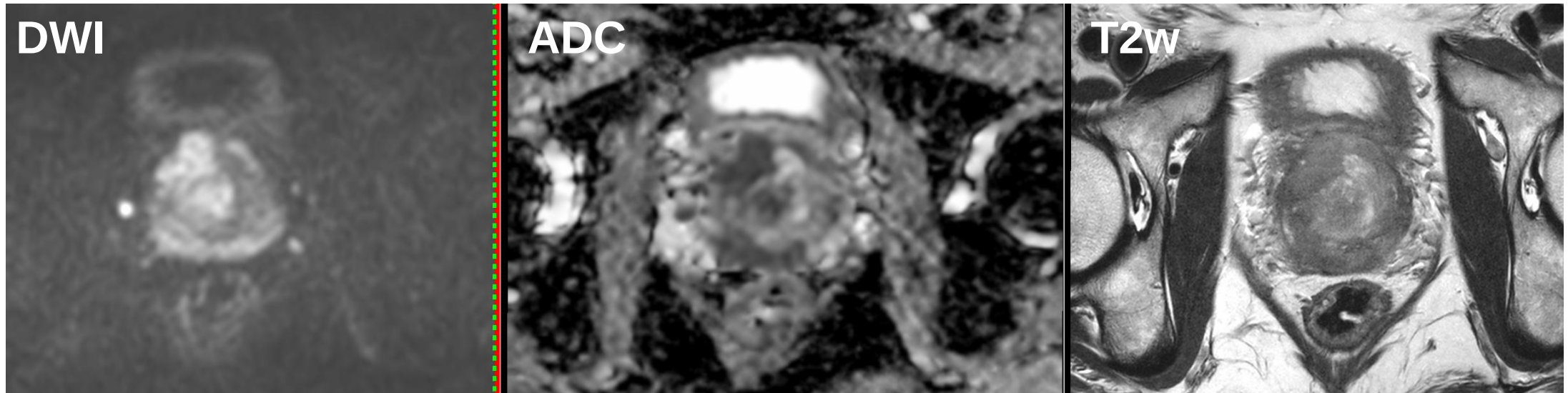


Basic 3-plane T2w, sagittal, coronal and axial. Anatomy is shown.

Patient preparation very important. Avoid rectal gas and ejaculation.

Image quality very important. High signal to noise, high resolution and no movements.

Functional imaging 1. Restricted water movement between cells and inside cells



DWI (Diffusion Weighted Imaging)

ADC (Apparent Diffusion Coefficient)

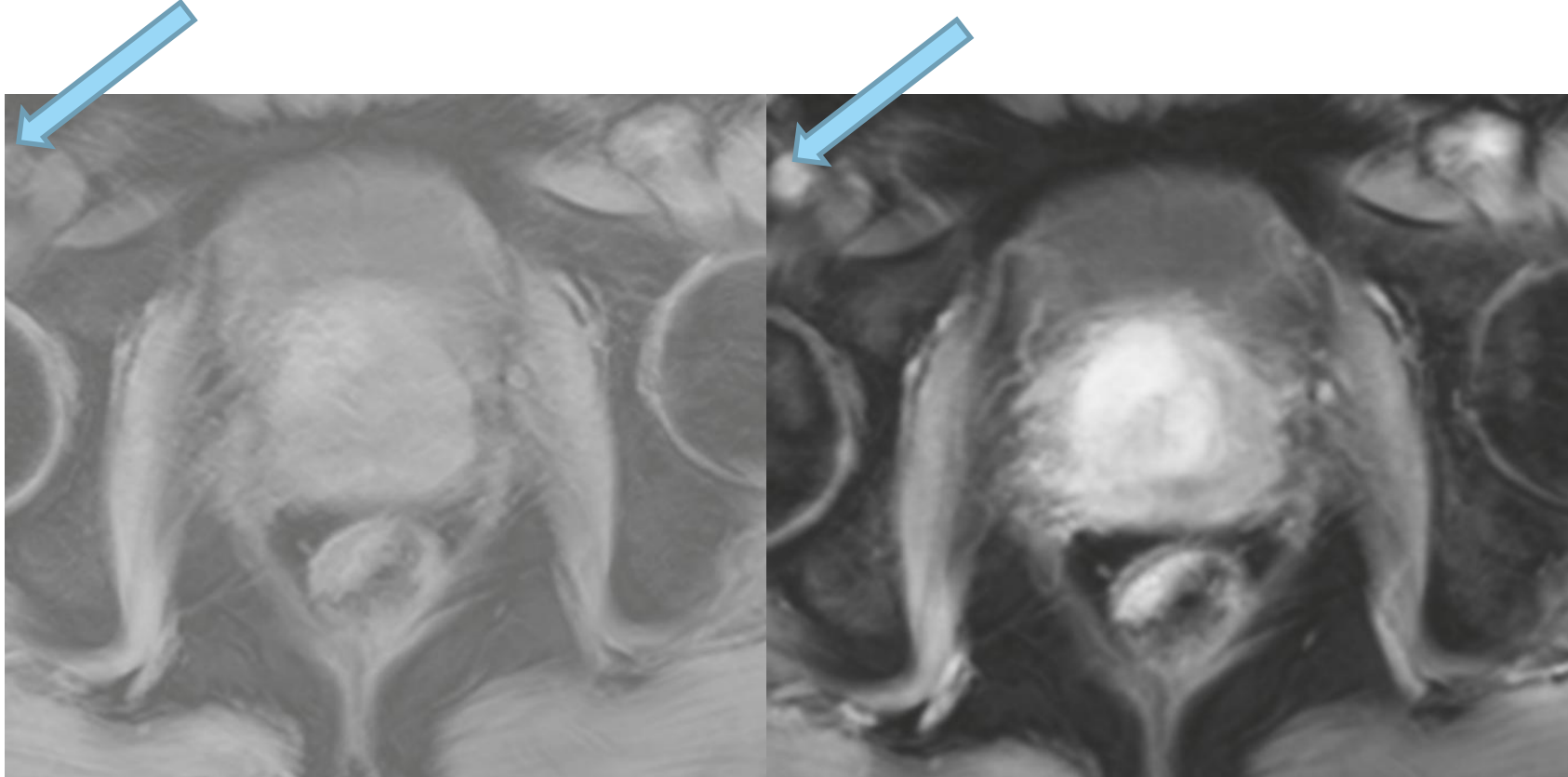
T2w (anatomy but also homogen tumor)

“Disease is White on Image”

“Aware Dark Cancer”

Memo

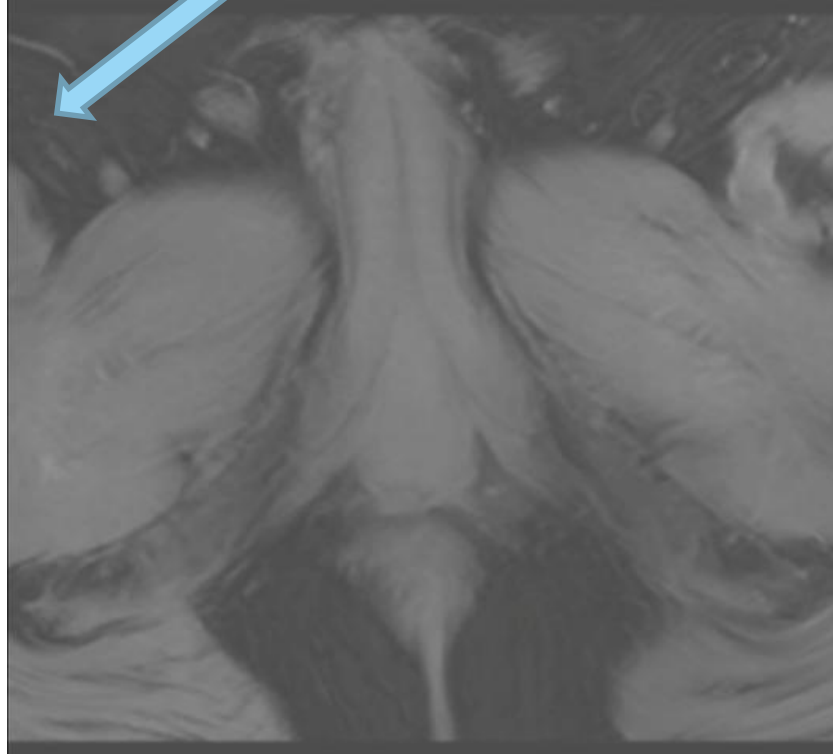
Functional imaging 2. Vascularity



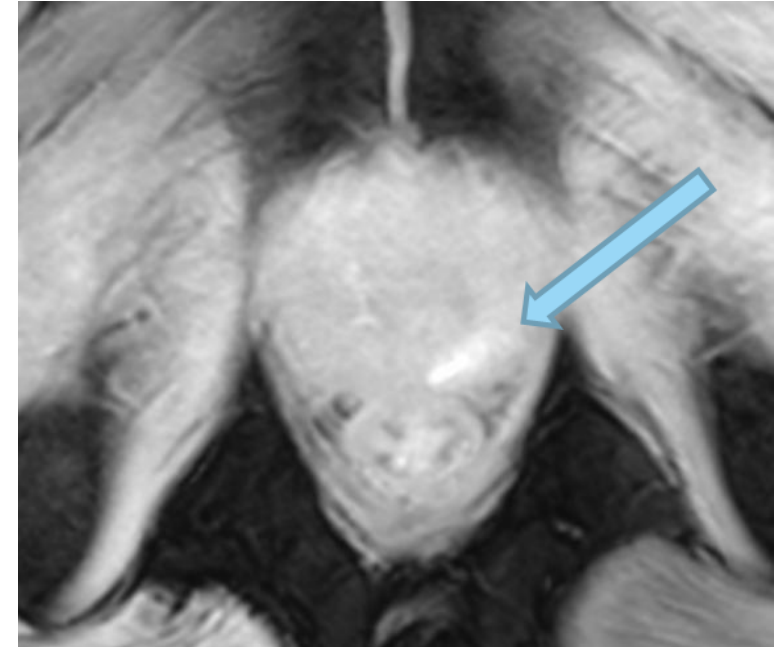
DCE (Dynamic Contrast Enhancement) time resolution $\leq 15\text{sec}$

T1w Fatsaturated or by subtraction

Functional imaging 2. Vascularity



A different patient shortly after biopsy with blood product in the prostate.



DCE (Dynamic Contrast Enhancement) time resolution $\leq 15\text{sec}$

T1w Fatsaturated or by subtraction

But you cannot say MRI for the prostata with out saying PI-QUAL

- **PI-QUAL version 2: an update of a standardised scoring system for the assessment of image quality of prostate MRI**
- Clinical relevance statement High image quality is crucial for prostate MRI, and the updated version of the PI-QUAL score (PI-QUAL v2) aims to address the limitations of version 1. It is now applicable to both multiparametric MRI and MRI without intravenous contrast medium.

Table 2 Essential technical prerequisites per sequence

T2-WI	DWI	DCE
3 mm Slice thickness	≤ 4 mm Slice thickness High <i>b</i> value sequence (≥ 1400 s/mm ²), calculated or acquired ADC map using at least two <i>b</i> values up to 1000 s/mm ²	3 mm Slice thickness Temporal resolution ≤ 15 s Fat suppression (or include post-processing, e.g. subtraction/heat maps) 3D sequences (preferred)

T2-WI T2-weighted imaging, DWI diffusion-weighted imaging, DCE dynamic contrast-enhancement, ADC apparent diffusion coefficient

This is multiparametric MRI of the prostate

- Images are evaluated according to PI-RADS guidelines v.2.1
- PI-RADS 1 – Very low (clinically significant cancer is highly unlikely to be present)
- PI-RADS 2 – Low (clinically significant cancer is unlikely to be present)
- PI-RADS 3 – Intermediate (the presence of clinically significant cancer is equivocal)
- PI-RADS 4 – High (clinically significant cancer is likely to be present)
- PI-RADS 5 – Very high (clinically significant cancer is highly likely to be present)



The diagram shows a prostate cross-section with two zones: the Peripheral Zone (indicated by a blue arrow) and the Transition Zone (indicated by a red arrow). A yellow circle labeled 'PI-RADS v 2.1' is positioned between the two zones.

Peripheral Zone		PI-RADS v 2.1	Transition Zone	
ADC / DWI			T2W	
1 Normal		PI-RADS 1	Normal appearing TZ (rare) or round, completely encapsulated nodule	1
2 ADC: Linear/wedge shaped hypointense and/or DWI: linear/wedge shaped hyperintense		PI-RADS 2	Mostly encapsulated nodule or Homogeneous circumscribed nodule without capsule or Homogeneous mildly hypointense area between nodules. DWI ≤ 3	2
3 ADC: Focal hypointense and/or DWI: focal hyperintense May be markedly hypointense on ADC or markedly hyperintense on high b-value DWI, but not both. DCE -		PI-RADS 3	Same as above but DWI ≥ 4	2
3 Same as above but DCE +			Heterogeneous signal intensity with obscured margins. Includes others that do not qualify as 2, 4, or 5. DWI ≤ 4	3
4 ADC: Focal markedly hypointense DWI: markedly hyperintense Diameter < 1.5cm		PI-RADS 4	Same as above but DWI = 5	3
4 ADC: Focal markedly hypointense DWI: markedly hyperintense Diameter < 1.5cm			Lenticular or non-circumscribed, homogeneous, moderately hypointense, < 1.5cm. any DWI	4
5 Same as 4, but ≥ 1.5cm or extraprostatic extension		PI-RADS 5	Same as 4, but ≥ 1.5cm or extraprostatic extension. any DWI	5

Why ?

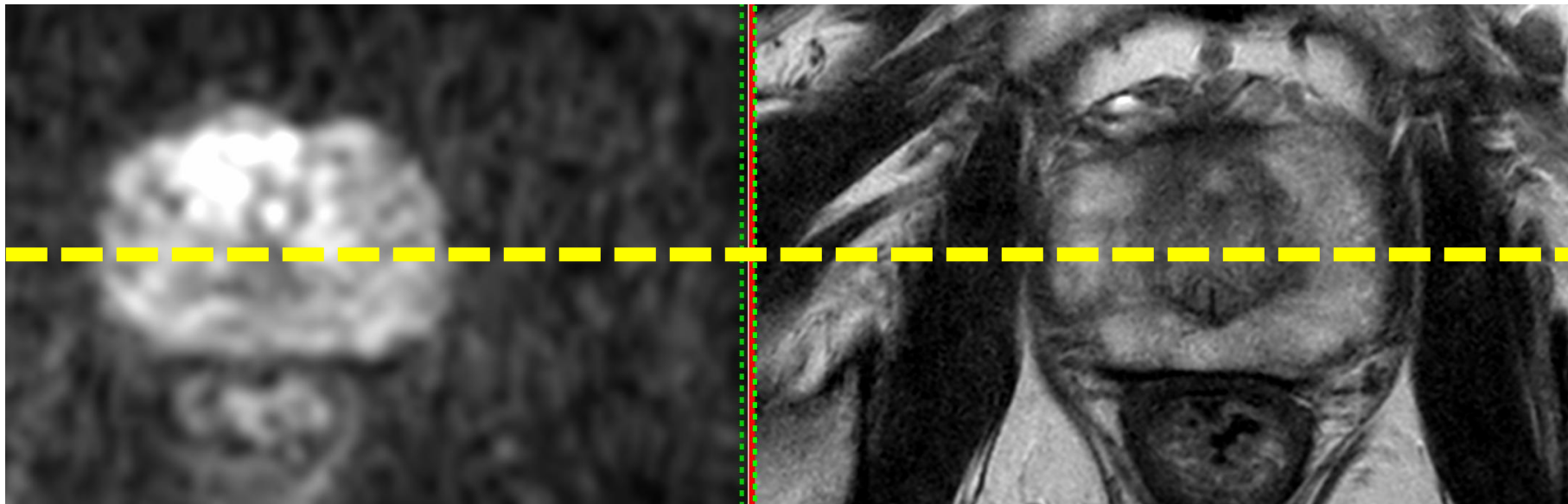
Numbers from 2022 World Cancer Research Fund

		<u>Incidents</u>	<u>ASR/100,000</u>	<u>deaths/ASR100,000</u>
Japan	125M	104,318	50,1	14,095/4.4
Denmark	5.9M	4,443	?	aprox.1370/?

- PROSTATE CANCER STATISTICS
- **Prostate cancer is the 4th most common cancer worldwide.**
- **It is the 2nd most common cancer in men.**
- **Global Numbers at a Glance**
- **New Cases:** ~1.47 million per year
- **Deaths:** ~397,000 per year
- **Global Incidence Rate:** 29.4 cases per 100,000 men
- **Lifetime Risk:** Roughly 1 in 8 men will be diagnosed
- Incidence Japan 167/100.000 men
- Incidence Denmark 151/100.000 men

Before the MRI era:

Biopsy was done transrectal, ultrasound guided or without image guidance



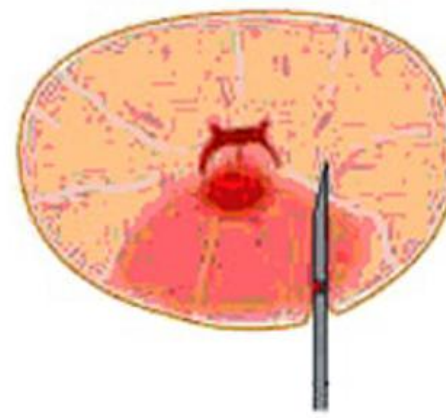
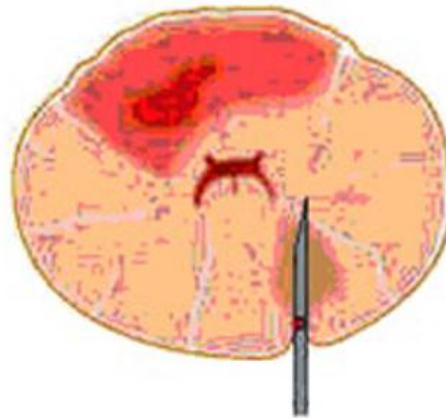
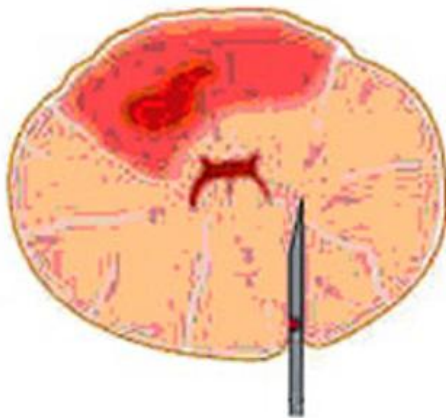
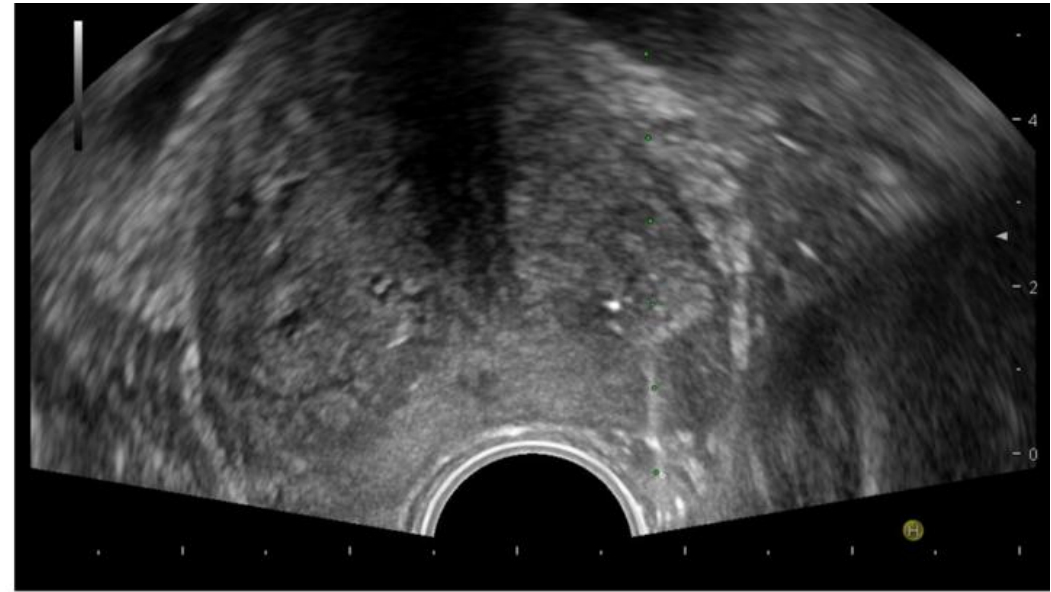
only 1,8 cm



TRUS biopsy transrectal

All men suspected on having PCa
with expected survival of >10years,

Sepsis 5%



How?

- Patient with non specific symptoms: PSA? Hæmospermia? Hamaturia? Problems voiding? Erectil dysfunction? Other?
- Visit GP
- Referral to specialist/hospital
- Referral to MRI (the whole team and information letter to the patient)..
- Imaging (the whole team), reading, reporting, lesion marking..
- Refere back to urologist, biopsy, (the whole team) send to..
- Pathologist (the whole team) slide analysis and reporting back to the
- Urologist.. Now informing the patient of treatment posiblities....
- Whats next?????

1. What is the role of the radiologist...

... in the MDT PCa approach?



That's me in the corner

MDT at Herlev hospital (DK) :

When: 2x per week, 1 hour.

Who: radiologist, nuclear-medicine physician,
oncologist, urologist, gynecologist, staff,
“flow”-nurses

How: prior, a list of 25pt w/ relevant
clinical information and questions.

I review images and the reports to speed
discussion

Everybody prepares to speed discussion !

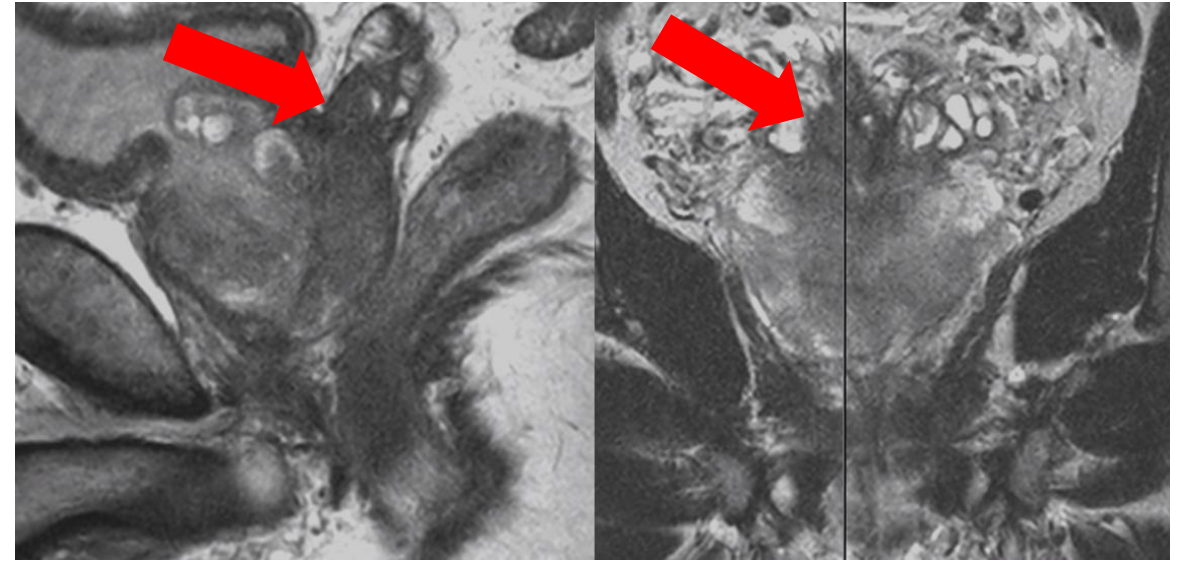
**MDT at Herlev hospital:
We discuss, one at the time.....**

- Co-morbidity etc.
- Where and how many metastasis?
- Is it necessary and/or possible to biopsy them?
- Is other imaging needed ? Which modalities?
- Treatment decision



And most important

- MRI evaluation of Prostate:
 - staging, seminal vesicles, urethra, EPE (“capsule”) and lymph nodes (local?) other findings? ..operability ...
- Clinical info:
 - urologist->DRE, mobility of prostate-> operability.
- MRI targeted biopsies:
 - Were they correct or a mismatch?
 - Good → feedback on own performance.



Evaluation...

• Mismatch - was it

- 1. bad MRI acquisition,
- 2. bad MRI reporting,
- 3. bad biopsies/bad contouring or alignment/targeting error
 - - many aspects
- Collect data to improve.

Evaluation...

- Compare prostatectomies with MRI
• Index lesions, secondaries
• Tumor size, location,
• spread..seminal vesicles
- INVISIBLE lesions!!!
- And learn from mistakes.....

EDUCATION
AND
EXPERIENCE

When treatment decisions are based on MRI findings
You must know your own performance statistics

But you cannot say Education and Experince with out saying..

➤ Eur Radiol. 2020 Oct;30(10):5404-5416. doi: 10.1007/s00330-020-06929-z. Epub 2020 May 19.

ESUR/ESUI consensus statements on multi-parametric MRI for the detection of clinically significant prostate cancer: quality requirements for image acquisition, interpretation and radiologists' training

Consensus-based criteria 'basic' versus 'expert' radiologists. N/A not applicable

Basic	Criterion	Expert
100	Minimum number of supervised cases before independent reporting	N/A
400	Minimum number of cases read	1000
150	Minimum number of cases/year	200*
1	Examination interval (year(s))	4
80	Agreement in double reads with expert centre (%)	≥ 90

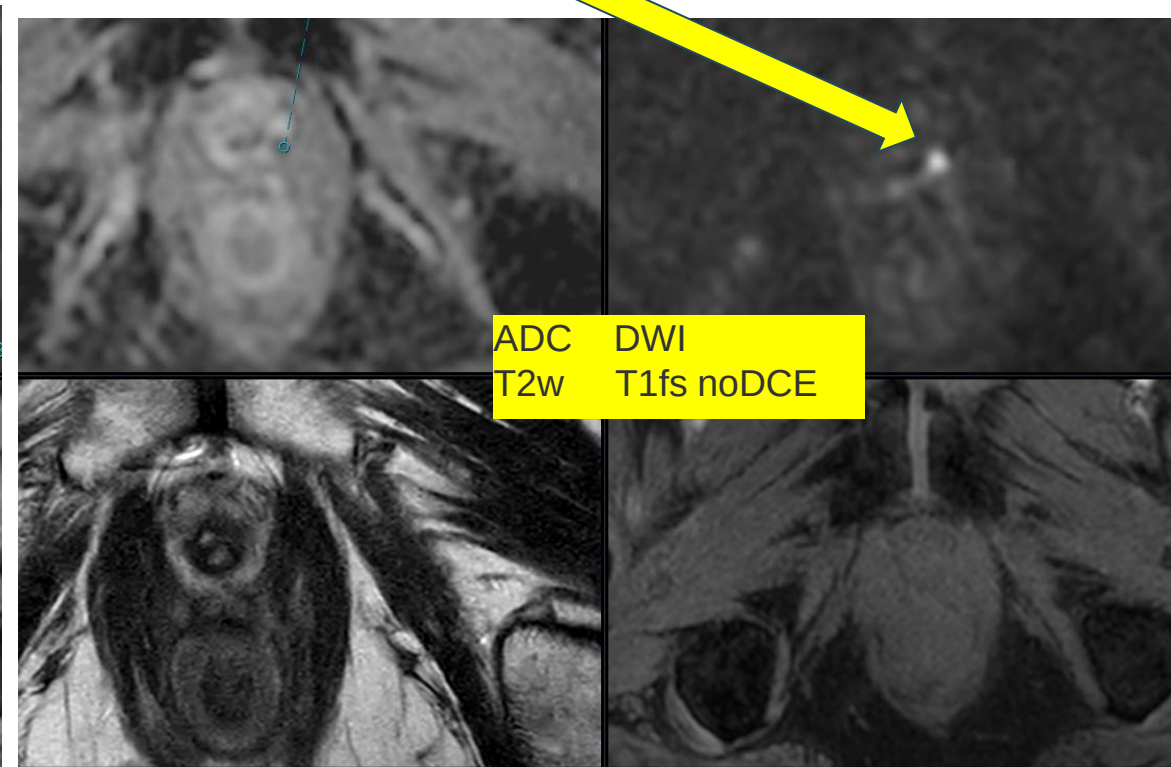
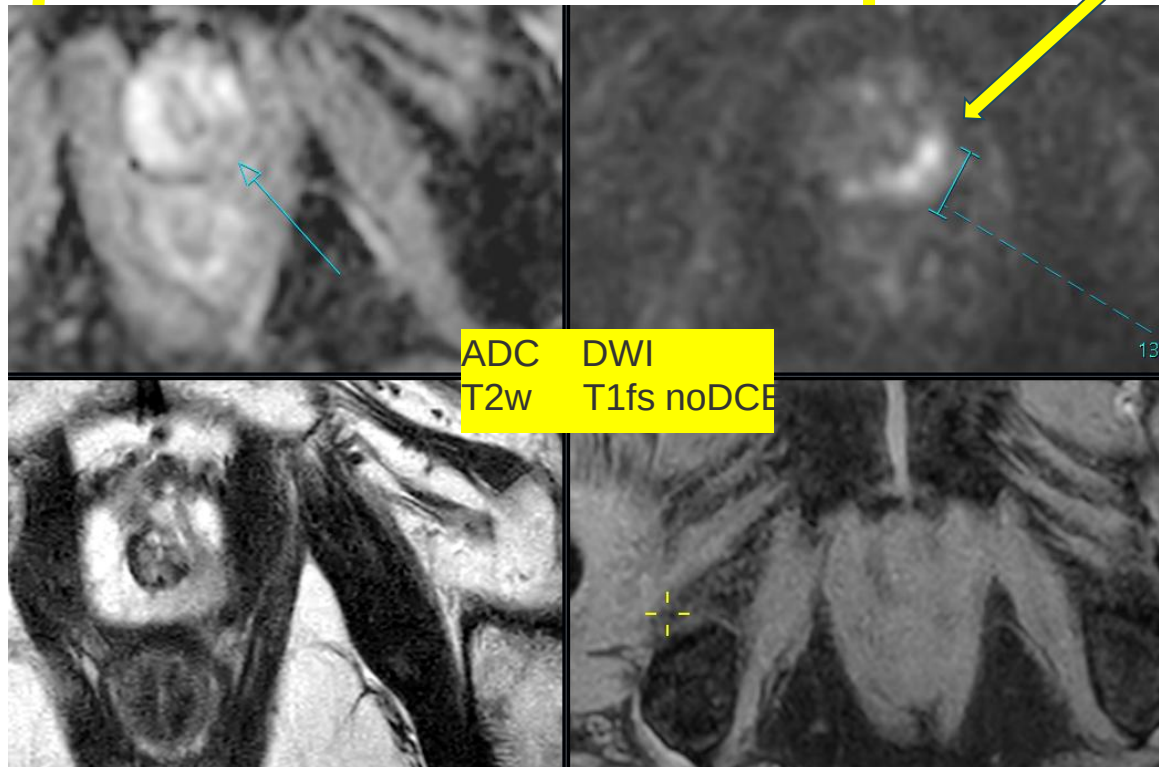
Three headings summarize the outcomes: (1) image quality assessment, (2) interpretation and reporting and (3) experience and training centres.

Let us have a case
62Y PSA 4.3 vol 39ml
PSAD 0,11.

- 1. MRI showed 2 lesions**

01:PIRADS 4 LMPZpm 13 mm - ADC 0,85

02:PIRADS 3-LAPZa 6 mm - ADC 0,63



01: GS 7 (3+4) i 5 / 15 mm, 2 / 11 mm, 1 / 13 mm og mikrofokus / 10 mm.

02: GS7 (3+4) i 3 / 11 mm, 0 / 13 mm, 0 / 13 mm og 0 / 4 mm.

Second MRI showed

nr 01 Il defined VM PZpm but location is marked

nr 02 PI-RADS 4 LA PZ a, 8 mm ADC 0,65

01:. Adenokarcinoma, Gleason score 7 (4+3) i 6 / 11 mm, 3 / 14 mm og 2 / 9 mm

02: Adenokarcinoma, Gleason score 9 (4+5) i 7 / 14 mm, 1 / 12 mm, 0 / 11 mm og 0 / 13 mm.

03: Benign biopsy

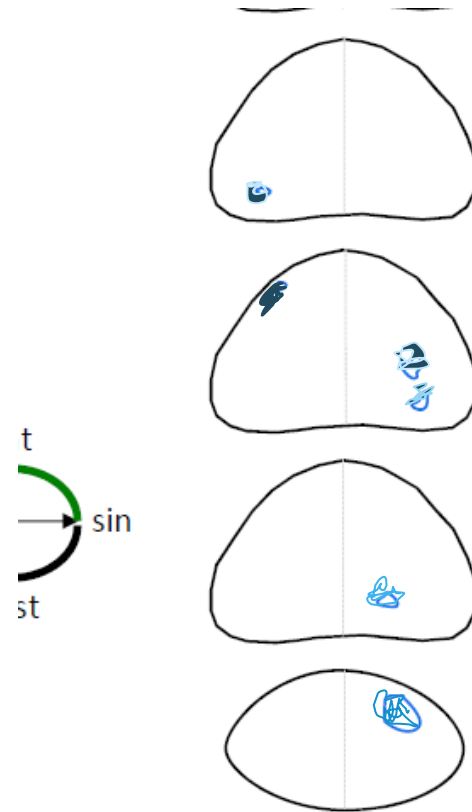
.

MDT meeting descision

- Biopsies showed adenokarcinoma Gleason Score 9 (4+5) in 5 of 10 biopsier from of (2/3 MR-targes).
- TNM: cT1c N0, M0, PSA 5,3.
- **Final conclusion**
- TNM: cT1c N0, M0, PSA 5,3. No metastasis shown on PSMA-PET-CT. No findings suspicious of seminal vescles involvement or EPE.
- High risk patient
- Prostatectomy is offered.

**We see the prostatectomy results at MDT
this ensures educational
feedback and a possibility
for improvement and research.**

Prostatectomy drawing
pT2 pNx pR0



But you cannot say Reporting and Local staging with out saying..

Agrotis et al. *European Radiology*
<https://doi.org/10.1007/s00330-026-12397-8>



INVITED REVIEW

Open Access

ESR Essentials: MRI-based T-staging in prostate cancer—practice recommendations by the European Society of Urogenital Radiology



Key Points

Question Wide variation in prostate MRI acquisition, image quality, and reporting undermines diagnostic accuracy. A structured roadmap is needed to ensure consistent quality and reproducible practice.

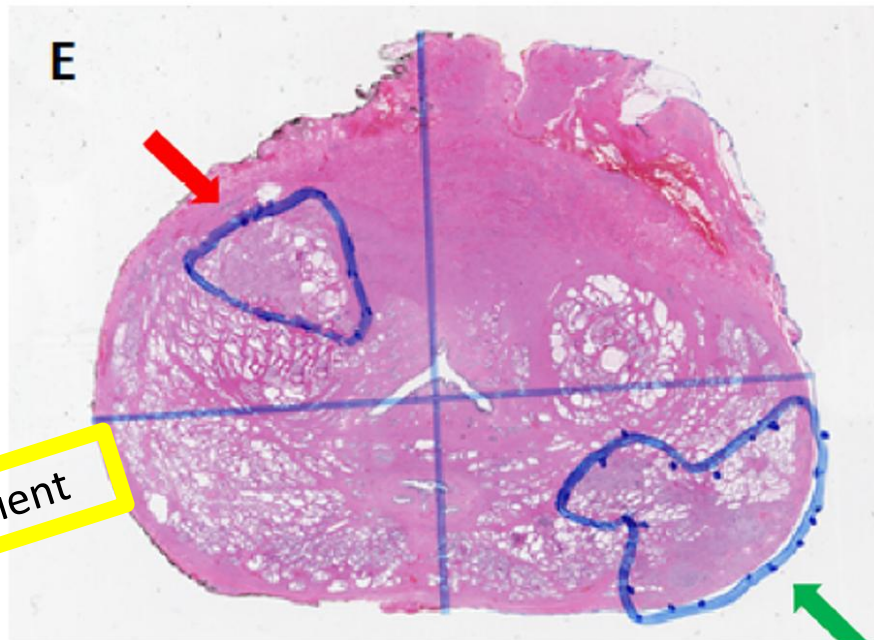
Findings The ESUR Prostate MRI Working Group outlines a three-step framework —‘Build it right’, ‘See it right’, ‘Improve and innovate’—to standardise acquisition, interpretation, and quality assurance.

Clinical relevance Applying this roadmap in clinical practice aims to enhance diagnostic confidence and promote consistent, high-quality prostate cancer care across diverse healthcare settings.

- What does your pathology report look like?
- Do you get histopathological feedback?

- Whole mount sectioning?
- Angulation of cuts of the tissue .
- Able to compare with MRI?
- And which surgical procedure ?

(Red arrow) with Gleason score 3 + 3.



(Green arrow) with Gleason score 3 + 4

What have we learned?

Good quality



Bad quality



Does the radiologist perform good independent of image quality?

- No of course not
- **Results: (only some)**
- High-quality images higher csPCa detection rate (56.5% to 64.3%)
- High-quality images had a higher PPV (72.5% to 78.8)
- PI-RADS 3 (24.0% to 52.9%) contained more low-quality images compared to
- PI-RADS 4-5 (20.6% to 39.3%),

Impact of prostate MRI image quality on diagnostic performance for clinically significant prostate cancer (csPCa)

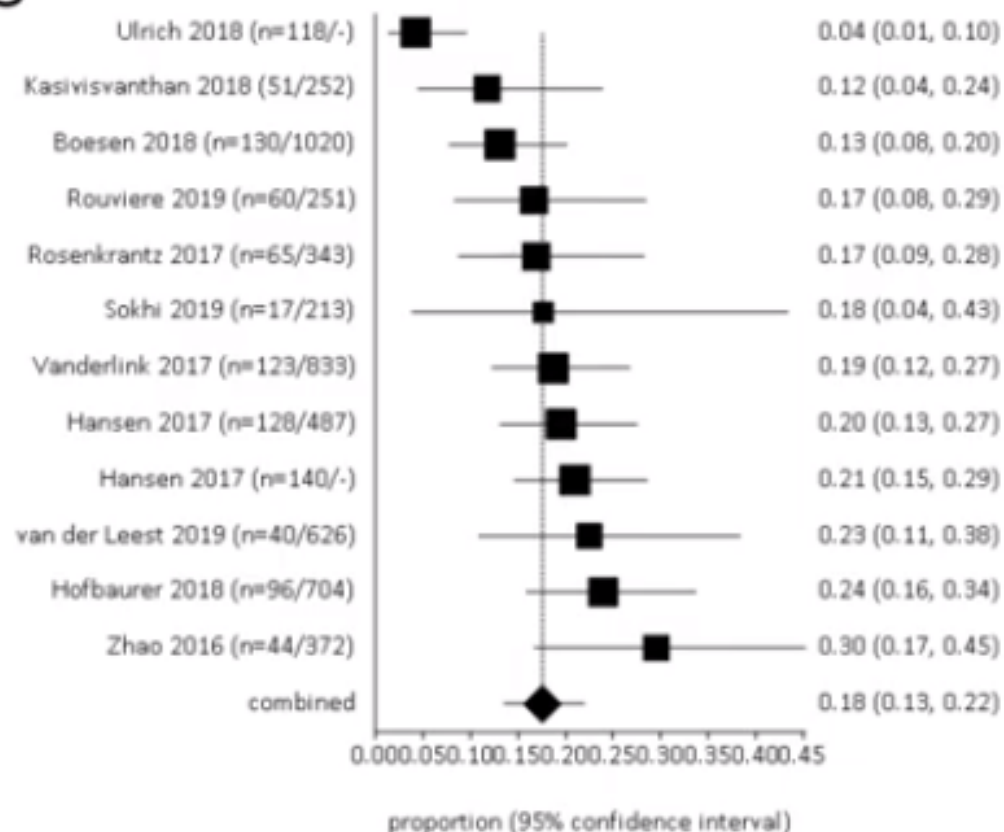
Abdom Radiol (NY) 2024 Nov;49(11):4113-4124. doi: 10.1007/s00261-024-04458-7. Epub 2024 Jun 27.

Cancers in PI-RADS 3 cases

Studies, Cases, Prevalence & Cancer Detection Rates

- 12 studies with 5362 patients (mixed)
- **PI-RADS 3 prevalence: 16%** (CI: 11, 20; range: 6-26), n=1015
- All cancers detection: 38% (CI: 34, 40; range 11-53)
- **ISUP $\geq$ 2: 18%** (13, 22; range, 4-30)

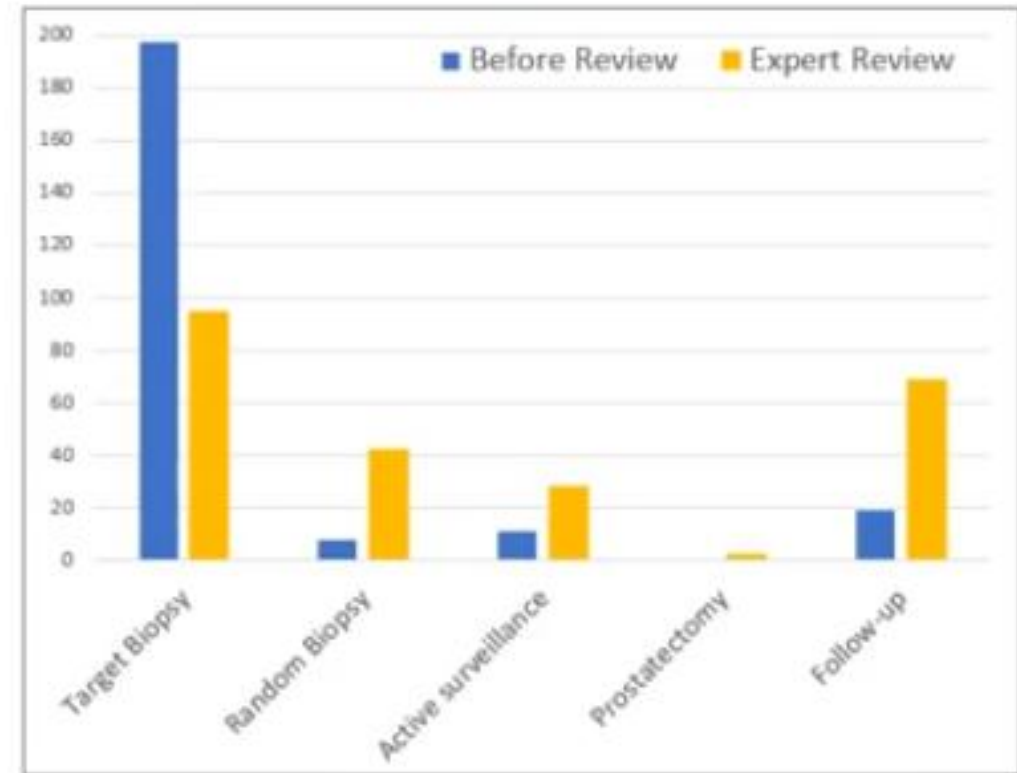
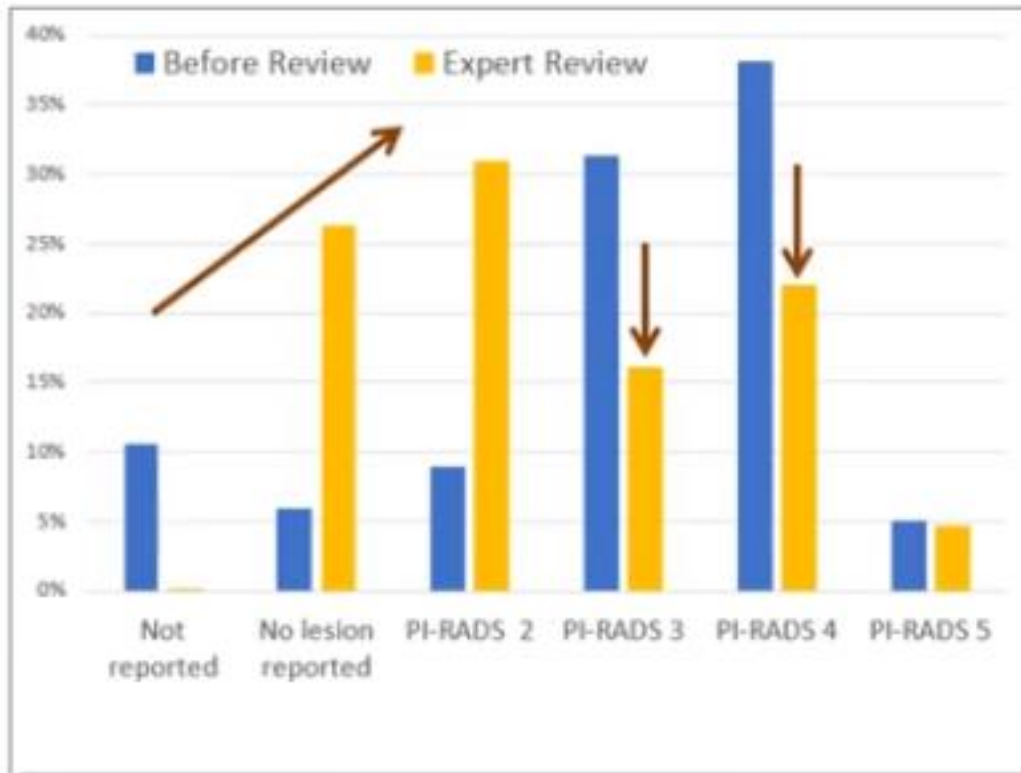
Proportion of PI-RADS 3 cases with ISUP $\geq$ 2



All PI-RADS v2
No Active Surveillance
Systematic & MR-directed
biopsies - all

Patient level analysis
5362 patients
1015 PI-RADS 3 cases
ISUP $\geq$ 2 18% (CI: 13, 22)

Expert radiologic-urological review reduces PI-RADS 3-4 cases and need for/type of biopsy



Significant discordance between PI-RADS reports in peripheral and tertiary subspecialized centers ($k=0.23$)

51% patient referred for targeted biopsies could be avoided

35% of men could skip prostate sampling all together

Luzzago S, Petralia G, et al. Clin Genitourin Cancer. 2018 [Epub]

Vibeke Løgager

THE LANCET

Diagnostic accuracy of multi-parametric MRI and TRUS biopsy in prostate cancer (PROMIS): a paired validating confirmatory study

Hashim U Ahmed^a, Ahmed El-Shater Bosaily^a, Louise C Brown^a, Rhian Gabe, Richard Kaplan, Mahesh K Parmar, Yolanda Collaco-Mor, Katie Ward, Richard G Hindley, Alex Freeman, Alex P Kirkham, Robert Oldroyd, Chris Parker, Mark Emberton, and the PROMIS study group

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

MRI-Targeted or Standard Biopsy for Prostate-Cancer Diagnosis

V. Kasivisvanathan, A.S. Rannikko, M. Borghi, V. Panebianco, L.A. Mynderse, M.H. Vaarala, A. Briganti, L. Budäus, G. Hellawell, R.G. Hindley, M.J. Roobol, S. Eggener, M. Ghei, A. Villers, F. Bladou, G.M. Villeirs, J. Viridi, S. Boxler, G. Robert, P.B. Singh, W. Venderink, B.A. Hadaschik, A. Ruffion, J.C. Hu, D. Margolis, S. Crouzet, L. Klotz, S.S. Taneja, P. Pinto, I. Gill, C. Allen, F. Giganti, A. Freeman, S. Morris, S. Punwani, N.R. Williams, C. Brew-Graves, J. Deeks, Y. Takwoingi, M. Emberton, and C.M. Moore, for the PRECISION Study Group Collaborators*

available at www.sciencedirect.com
journal homepage: www.europeanurology.com



Platinum Priority – Prostate Cancer
Editorial by XXX on pp. x–y of this issue

Head-to-head Comparison of Transrectal Ultrasound-guided Prostate Biopsy Versus Multiparametric Prostate Resonance Imaging with Subsequent Magnetic Resonance-guided Biopsy in Biopsy-naïve Men with Elevated Prostate-specific Antigen: A Large Prospective Multicenter Clinical Study

Marloes van der Leest^a, Erik Cornel^b, Bas Israël^a, Rianne Hendriks^c, Anwar R. Padhani^d, Martijn Hoogenboom^e, Patrik Zamecnik^f, Dirk Bakker^g, Anglita Yanti Setiasti^h, Jeroen Veltmanⁱ, Huib van den Hout^j, Hans van der Lelij^k, Inge van Oort^l, Sjoerd Klaver^m, Frans Debruyneⁿ, Michiel Sedelaar^o, Gerjon Hannink^p, Maroeska Rovers^q, Christina Hulsbergen-van de Kaa^{r,s}, Jelle O. Barentsz^{t,u,v}



BIDOC

PRECISION

MRI-FIRST

4M

COCHRANE

JAMA
Network | Open



Original Investigation | Urology

Assessment of the Diagnostic Accuracy of Biparametric Magnetic Resonance Imaging for Prostate Cancer in Biopsy-Naive Men The Biparametric MRI for Detection of Prostate Cancer (BIDOC) Study

Lars Boesen, MD, PhD; Nis Nergaard, MD; Vibeke Løgager, MD; Ingegerd Balslev, MD; Rasmus Bisbjerg, MD; Karen-Cecilie Thstrup, MD; Mads D. Winther; Henrik Jakobsen, MD; Henrik S. Thomsen, DMC



Use of prostate systematic and targeted biopsy on the basis of multiparametric MRI in biopsy-naive patients (MRI-FIRST): a prospective, multicentre, paired diagnostic study

Olivier Rouvière, Philippe Puech, Raphaële Renard-Penna, Michel Claudon, Catherine Roy, Florence Mège-Lechevallier, Myriam Decaussin-Petrucci, Marine Dubreuil-Chambardel, Laurent Magaud, Laurent Remontet, Alain Ruffion, Marc Colombel, Sébastien Crouzet, Anne-Marie Schott, Laurent Lemaître, Muriel Rabilloud, Nicolas Grenier, for the MRI-FIRST Investigators*

Cochrane Database of Systematic Reviews

Prostate MRI, with or without MRI-targeted biopsy, and systematic biopsy for detecting prostate cancer

Cochrane Systematic Review - Diagnostic | Version published: 25 April 2019
<https://doi.org/10.1002/14651858.CD012663.pub2>

Am score 33 View article information

Frank-Jan H Drost | Daniël F Osses | Daan Nieboer | Ewout W Steyerberg | Chris H Bangma | Monique J Roobol
| Ivo G Schoots
View authors' declarations of interest

Level 1 evidence – Clear advantage of an MRI-based diagnostic strategy

Multiple factors influencing biopsy-outcome

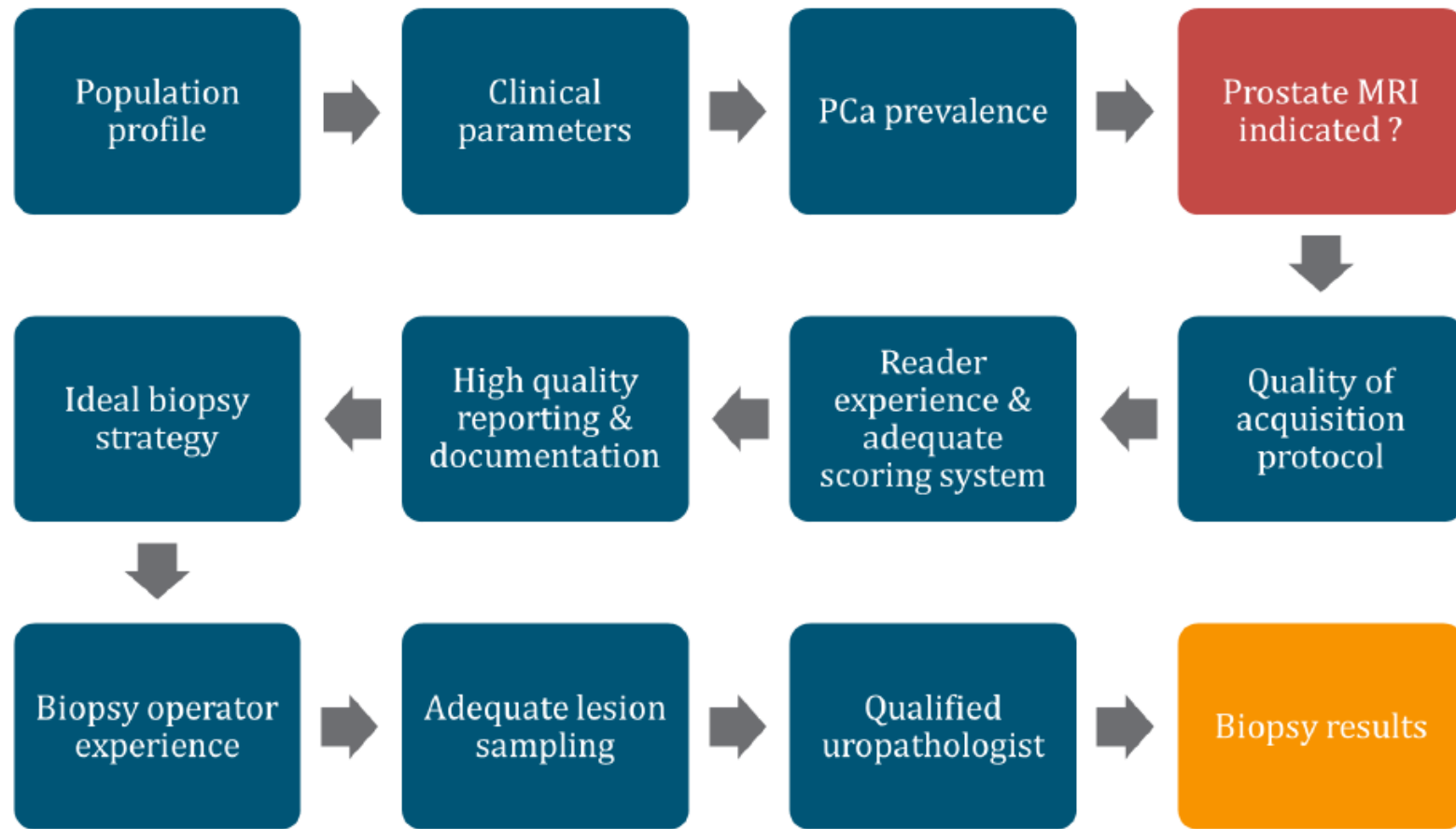


Figure 7: Factors that can influence the biopsy outcome of an MRI-guided diagnostic pathway.

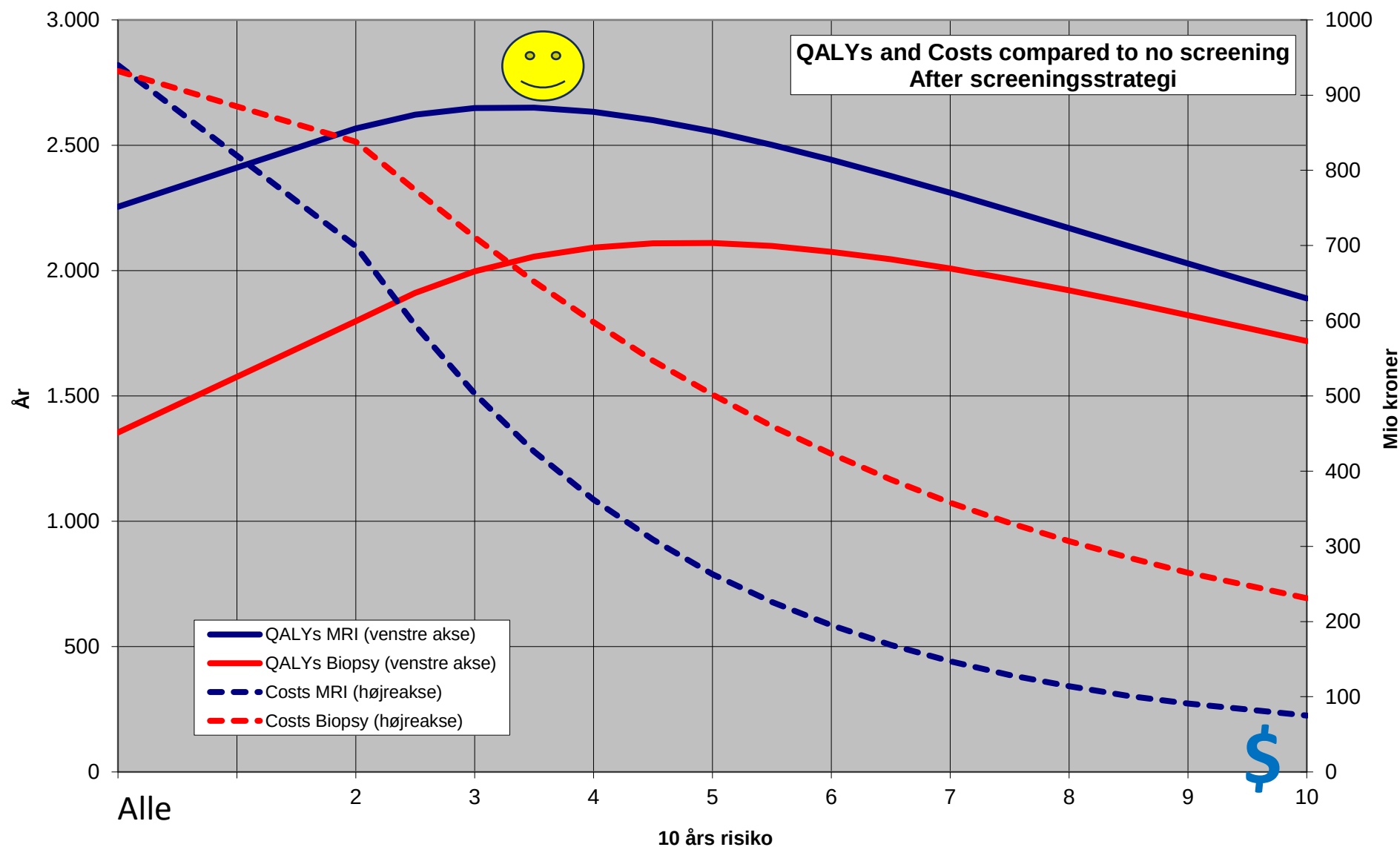
Is it worth it?



CanTest frameworks for evaluation of diagnostic strategies for early and timely detection of cancers

Phases of evaluation	Phase 1 Selection of test and initial measures of single test performance	Study objectives	Analytical validity	Diagnostic accuracy	Refinement of the test					
		Study designs	Assay performance					Case series	Case-control	
	Phase 2 Measures of clinical test performance	Study objectives	Diagnostic accuracy	Test feasibility in intended settings: staffing, sampling, sample processing				Analytical validity/test reproducibility in intended setting		
		Study designs	Case series				Case-control	Cohort	Qualitative	Assay performance
	Phase 3 Impact on clinical decision-making and health outcomes	Study objectives	Diagnostic accuracy	Effects on patients and clinicians: acceptability, feasibility, benefits and harms				Effects on diagnostic triage	Incorporation of test into diagnostic strategies	
		Study designs	Natural experiments				Cohort	Randomized controlled trial	Qualitative	Health economic modelling
	Phase 4 Effectiveness of new diagnostic strategy on clinical outcomes	Study objectives	Effectiveness and cost-effectiveness of diagnostic strategy compared to existing strategy				Patient safety and quality: appropriate test use, accurate clinical interpretation, test follow-up			
		Study designs	Natural experiments				Cohort	Randomized controlled trial	Step-wedge design	Qualitative
	Phase 5 Implementation and effects at health-care and population level	Study objectives	Effects on health system: utilization, referral patterns, costs			Effects on population: stage, mortality, inequalities				
		Study designs	Natural experiments			Analysis of routine data		Qualitative	Health economic impact studies	

Results III: Screening minus no screening



Ekstra QALYs
Ekstra Costs

Blue for MRI first strategy
Red for Biopsi first strategy

QALYs are
consequently higher
in MRI first strategies
Costs is constantly
lower in MRI first
strategies

QALY max at 10 yrs
risk 3-4% for MRI
first strategy
At price 350-500 mio
kr over several yrs

Results IV: Screening minus no screening

Screeningsstrategier

JAMA
Network | **Open**

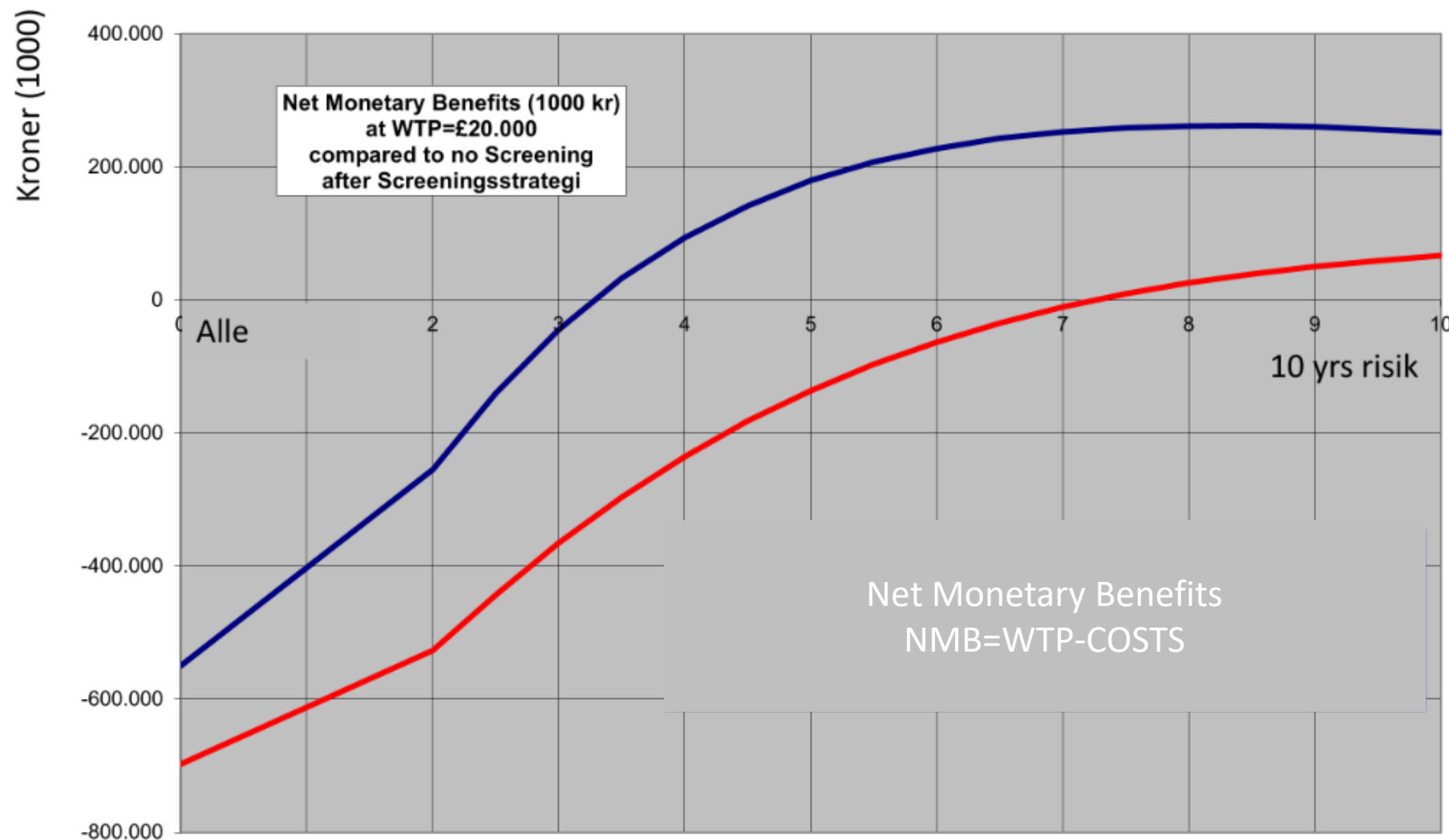
Callender et al, 2021

Net Monetary
Benefits

Blue for MRI first strategy
Red for Biopsi first strategy

NMB er
consequently higher
for MRI first strategy
than for Biopsi first
strategy

NMB reaches
plateau at 10 yrs
risk of 7% for MRI
first strategy, at the
price of 150 mio kr
over several yrs



Summa Summarum

- MRI first strategies exceeds Biopsy first strategy on:
- Benefit-harm profiles
 - On QALYs and
 - On Costs
- **Quite convincing**
- Pointing towards a risk based strategy
 - Preferable 7% risk over 10 yrs – depending on willingness to pay
 - Minimum 3-4% over 10 år – otherwise the gain of QALYs is lost
- This could be input for a new RCT in Europa – or at least a prospektiv evaluation
- This is what the Health Econom-y (-mist) says
- *But they say alot of things...*



JES SØGAARD

Putting bits and pieces
together



Thank you

- With a lot of help from all our friends...



Thank you

- With a lot of help from all our friends...



Any questions?