
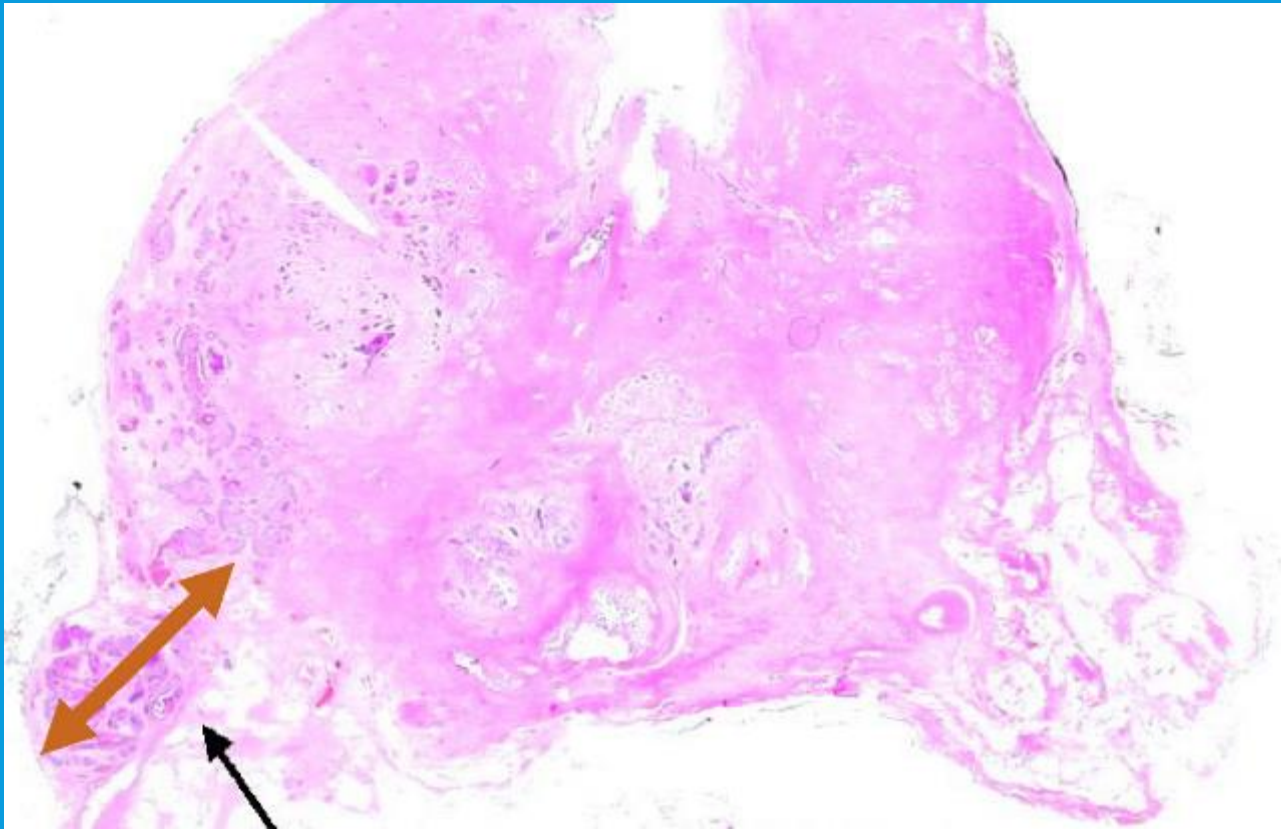


# INTRODUCTION:



- EPE is an unfavourable prognostic factor on PCa progression & survival
- TNM staging system:
  - T<sub>3a</sub>  PCa with extension into periprostatic soft tissue  
(without further subclassification)
- The clinical outcomes of patients with EPE showed variations
- The degree of extraprostatic extension:
  - Associated with the tumor prognosis & biochemical recurrence rates.
  - Various methods have been described for quantifying & subclassifying
  - There was no consensus on the optimal criteria

# INTRODUCTION (RADIAL DISTANCE):



- The radial distance of EPE:
  - The distance of prostate tumor extension that is perpendicular to outer margin of the prostatic stroma
- Pathological radial distance (RD)
  - An independent indicator for prognosis & biochemical recurrence
- Quantifying the degree of EPE with RD is simplest and objective way.

# INTRODUCTION (MP-MRI):

- Clinical staging of PCa;
- Digital rectal examination,
- Clinical nanograms (PSA & bx Gleason score) has limited accuracy on preoperative risk assessment of EPE
- Mp-MRI;
- Reference standard imaging modality
- The subjective assessment of EPE using prostate MRI  specificity but  sensitivity with poor interobserver agreement rates.

# INTRODUCTION:

- Prostate MRI might have potential on determining pathological RD in true positive patients for EPE.
- To the best of our knowledge, the value of prostate MRI in determining the degree of EPE has not been established yet.
- The aim of this study is to evaluate the value of prostate MRI in the assessment of pathological RD in prostate cancer.

# MATERIALS & METHODS (PATIENT SELECTION):

- Retrospective,
  - 106 patients with prostate cancer who underwent prostate MRI prior to RP between 2012-2017
- Inclusion criteria:
  - *Patients with satisfactory medical, pathological and imaging records,*
  - *The time interval between prostate Bx & Mp-MRI scanning > 6 weeks,*
  - *The time interval between Mp-MRI scanning & RP > 6 months*
- Exclusion criteria:
  - *Patients with preoperative hormonal or radiation therapy*

# MATERIALS & METHODS (MRI SCANNING):

- 3T MRI scanner (Magnetom Skyra, Siemens Medical Solutions, Erlangen, Germany) & 16-channel phased array surface coil
- The Mp-MRI protocol included; tri-planar T<sub>2</sub>-WI, T<sub>1</sub>-WI, DCE-MRI & DWI with ADC map
- The imaging parameters of tri-planar (in axial, coronal and sagittal planes) T<sub>2</sub>W TSE:

PARAMETER	VALUE
TR/TE	3566–3631/100 msec
Matrix size	512 × 352
FOV	200 mm
Slice thickness	3 mm

# MATERIALS & METHODS (PATHOLOGICAL ANALYSIS):

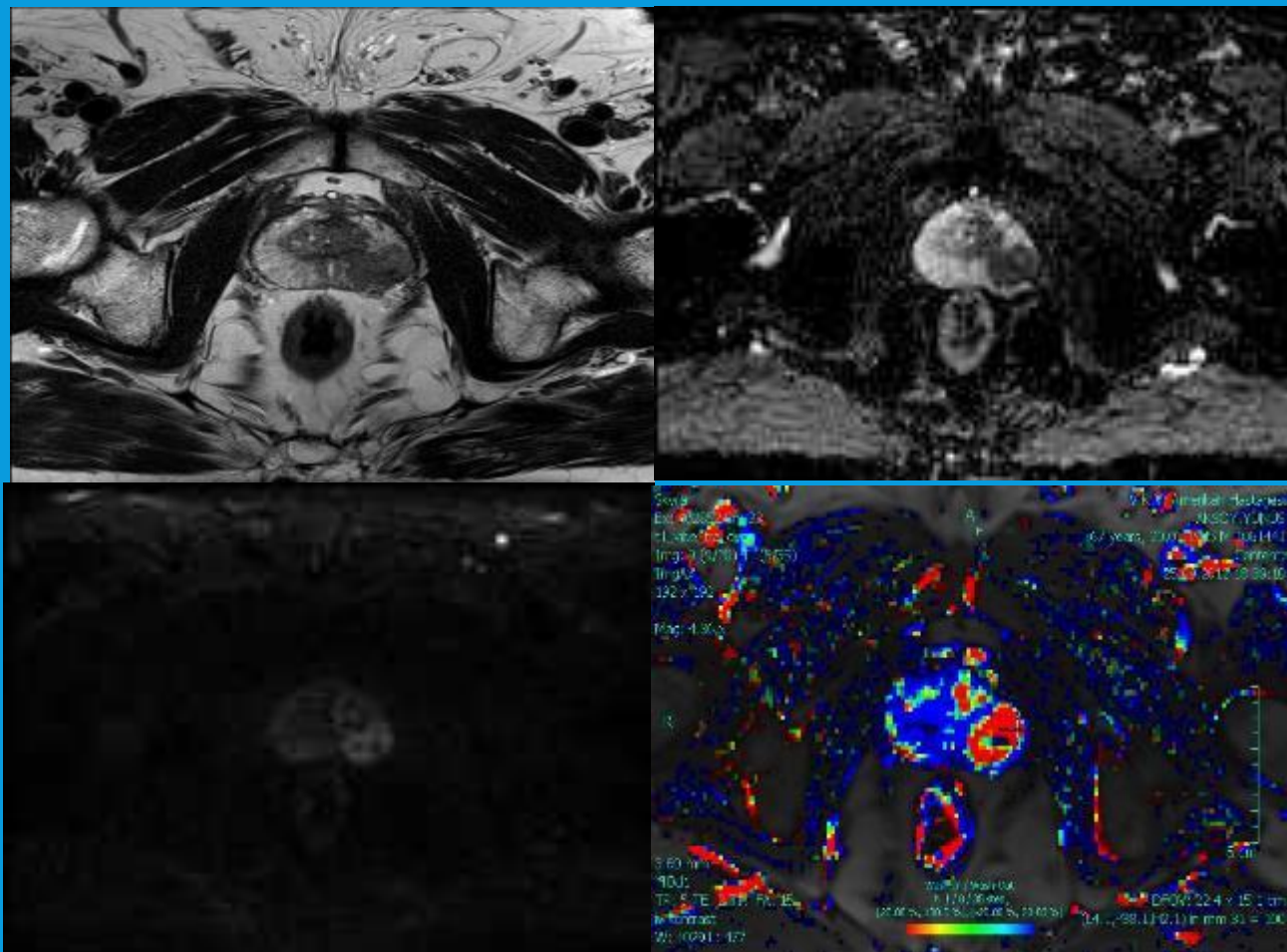
- The RP specimens were processed by an experienced genitourinary pathologists in accordance with the standard procedures recommended by the International Society of Urologic Pathology
- The index lesions were marked on 16-sector divided standardized prostate diagram by the urogenital pathologist who is blinded to any MRI interpretations.
- The following criteria have been used to describe index lesion;
  - 1- Prostate tumors with extension to outer margin of prostatic stroma
  - 2- If, none of the tumor foci had extraprostatic extension, the tumor foci with the highest Gleason score
  - 3- If the tumor foci have the same Gleason score, the lesion with maximum size

# MATERIALS & METHODS (PATHOLOGICAL ANALYSIS):

- The volume, GS & the presence / absence of EPE
- The pathological RD of EPE was measured by using an ocular micrometer from the focus with maximum extension if multiple foci of EPE were exist on RP specimens.



# IMAGE ANALYSIS AND MEASUREMENT OF MRI-DETERMINED RADIAL DISTANCE:

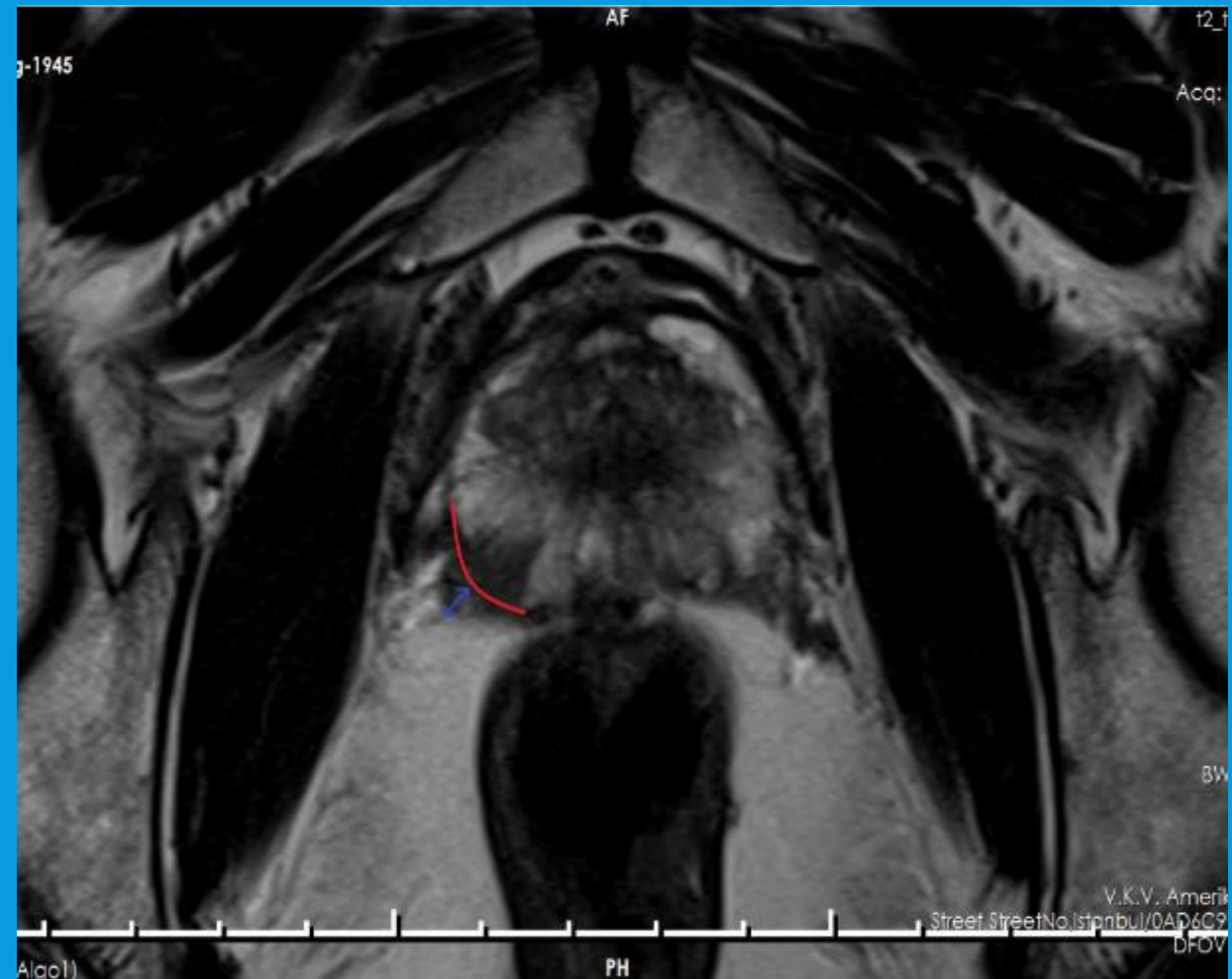
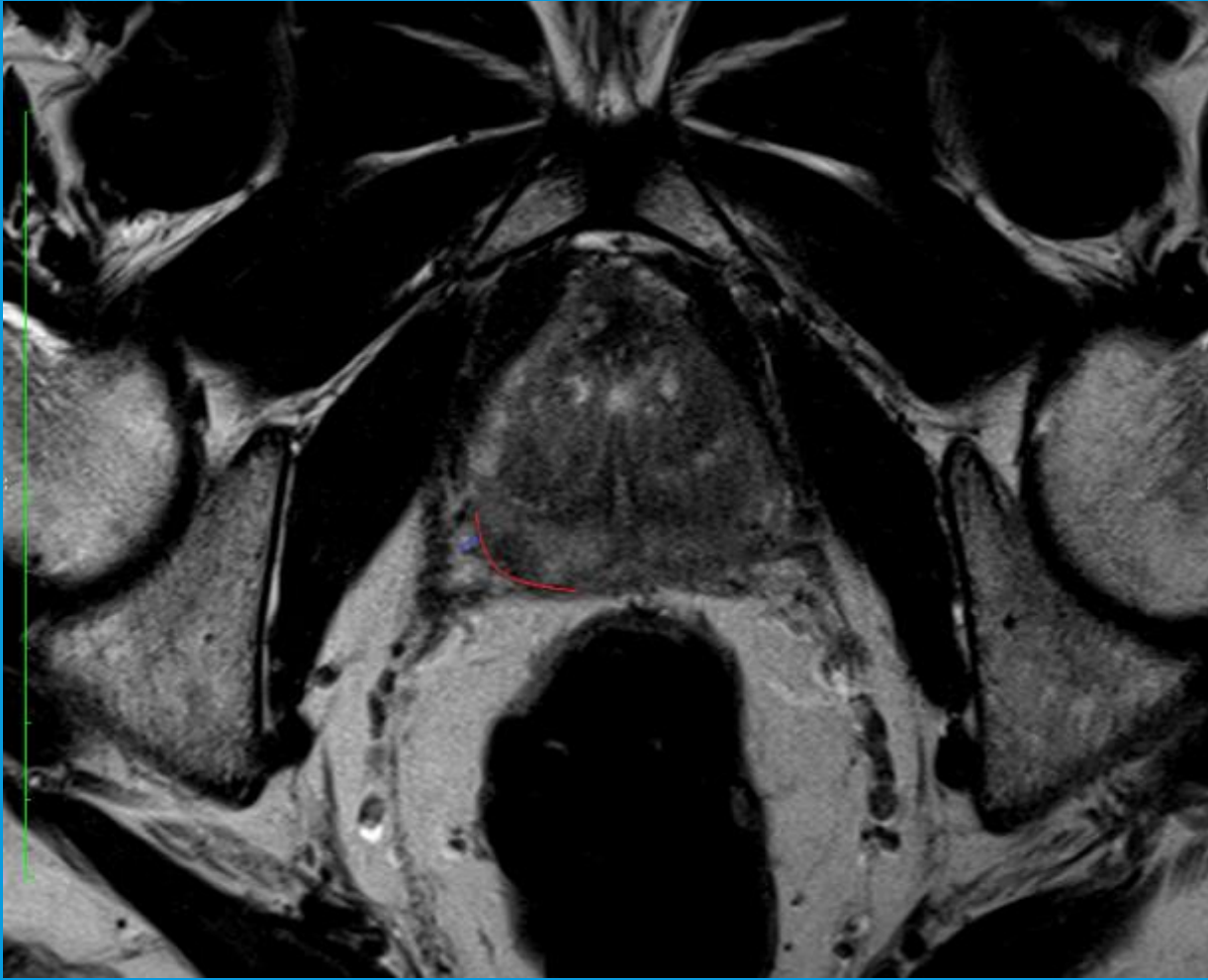


Two radiologists were independently evaluated the images.

The radiologists identified the dominant tumor as defined a mass like lesion that showed low signal intensity on T2WI and ADC map with or without early contrast enhancement

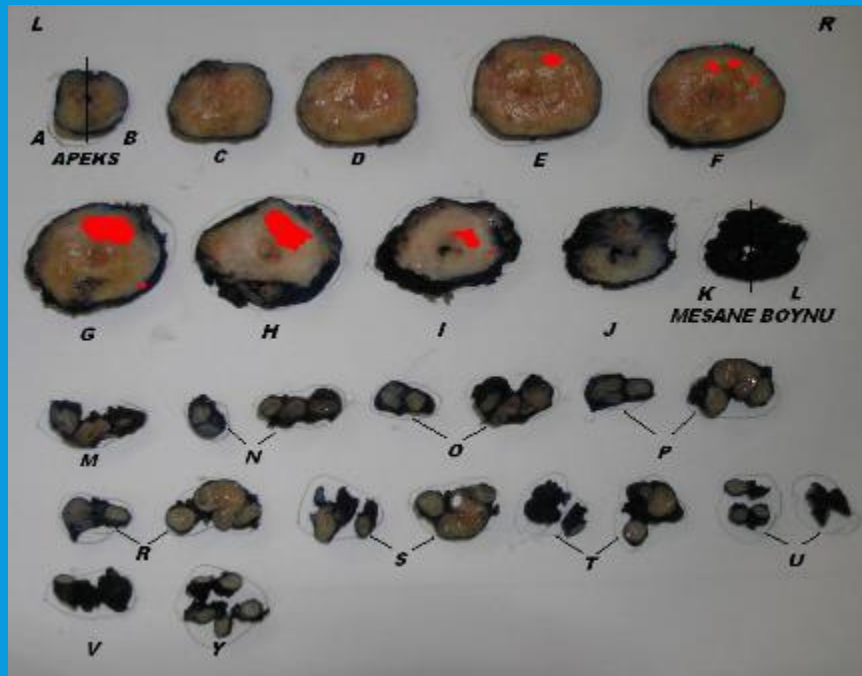
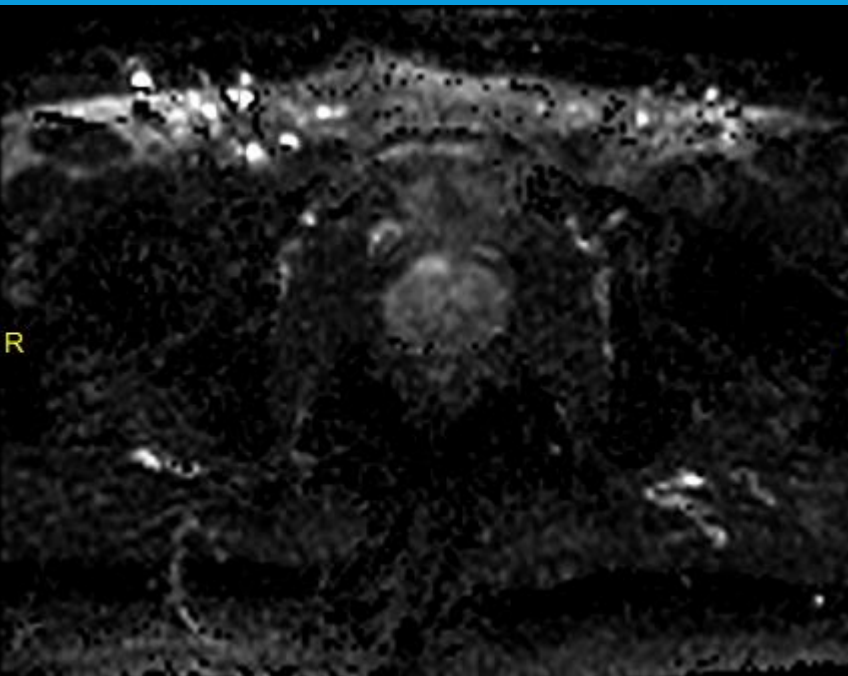
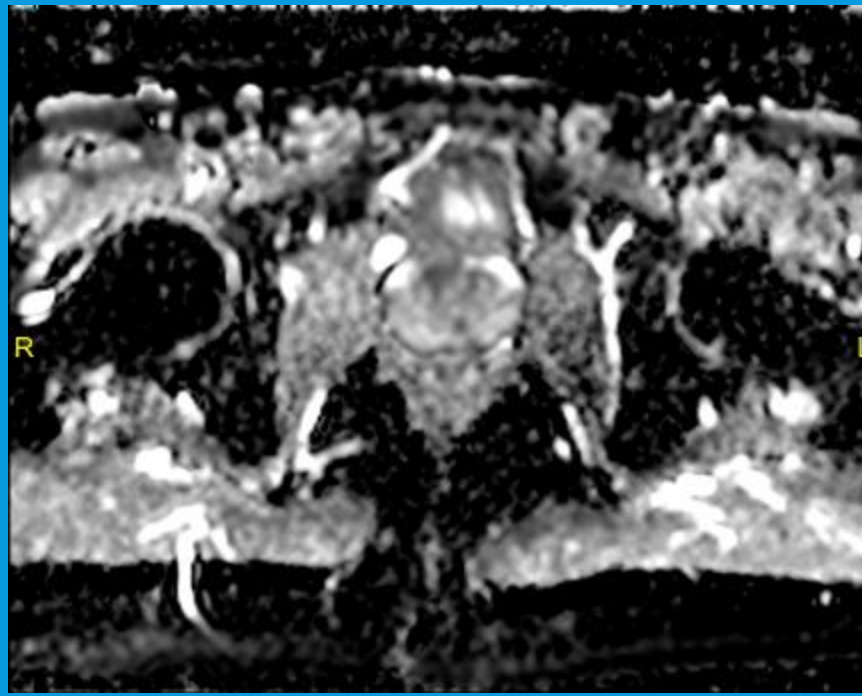
Dominant lesions were subjectively evaluated with a 5-point probability scale by using the modified PI-RADS criteria for assessment of EPE.

	Probability Scale	
LIKERT SCALE-1	EPE absent	If the normal tissue can be visualized between intact prostate capsule & tumor
LIKERT SCALE-2	EPE probably not present	If the tumor abuts prostate capsule
LIKERT SCALE-3	Equivocal for EPE	If the tumor abuts and cause irregularity on prostate capsule
LIKERT SCALE-4	EPE probably present	If tumor bulges, deforms and obscures the prostate capsule
LIKERT SCALE-5	EPE absolutely present	If gross, measurable tumor is detected



- Likert scale 4 and 5 presumed as positive for EPE.
- The radiologists measured the RD of EPE on T2WI for the patients considered as EPE (+)
- The RD measured perpendicular to outer margin of prostate.





- After the image evaluation, the location of index lesions and subjective EPE assesment were examined by using the RP specimes as referance standart.

- The results were classified under 4 categories: true-positive results, true-negative result, false positive results and false negative results

True Positive	<p>The dominant tumor focus AND The presence of EPE have been correctly determined</p>	
True Negative	<p>The dominant tumor focus AND The absence of EPE have been correctly determined</p>	
False Positive	<p>The dominant tumor focus has been correctly determined AND Incorrectly suspected from presence of EPE</p>	<p>Incorrectly suspected from dominant tumor focus with the presence of EPE</p>
False Negative	<p>The dominant tumor focus have been correctly determined AND The presence of EPE could not be determined</p>	<p>The dominant tumor focus with the presence of EPE could not be determined</p>

# MATERIALS & METHODS (STATISTICAL ANALYSIS):

- Sensitivity, specificity, PPV & NPP (prostate MRI for assessment of EPE according to subjective analysis)
- The relationship between pathological RD & MRI-determined RD were calculated separately for patients with Likert scale of 4 and 5 by using ICC analysis.
- The relationship between pathological RD&MRI determined RD were calculated for the patients with true positive results by using ICC analysis.
- Cohen's kappa statistics was used to calculate the inter-rater reliability of the subjective EPE assesment with modified Likert scoring between readers.
- The inter-reader relability of RD were calculated between readers by using ICC

# RESULTS (HISTOPATHOLOGICAL ANALYSIS)

- EPE (+) = 23.6%
- EPE (-) = 76.4 %
- The mean pathological RD of index lesion= 1.8 mm (ranged between 0.2 mm to 7 mm)
- The median pathological RD of index lesion = 1.1 mm

# RESULTS

## Prostate MRI for assessment of EPE according to subjective analysis

	Reader-1	Reader-2
Sensitivity	60.0%	53.3%
Specificity	86.8%	92.1%
PPV	64.3%	72.7%
NPV	84.6%	83.3%



# RESULTS (MRI ANALYSIS)

- Cohen's Kappa statistics showed good inter-rater reliability with respect to the Likert scoring (ICC: 0.727)

	Reader-1	Reader-2
EPE absent or probably not present	%53.8	%55.6
EPE equivocal	%28.3	%24.5
EPE probably present	%11.3	%13.2
EPE absolutely present	%6.6	%6.6

# RESULTS

	Reader-1	Reader-2
The mean MRI-determined RD	1.9 mm (ranged between 0.5 mm to 5.2 mm)	2.0 mm (ranged between 0.9 mm to 4.8 mm)
The median MRI-determined RD	1.9 mm	1.7 mm

- With respect to measurement of MRI determined RD ICC analysis showed good agreement between the readers for patients with true positive results (ICC=0.898)

# RESULTS

- Between pathological RD & MRI determined RD
  - Likert score-5 = good reliability( single measure ICC = 0.820; reader-1,single measure ICC = 0.880; reader-2)
  - Patients with true positive results=good reliability (single measure ICC = 0.77; reader-1,single measure ICC = 0.73; reader-2).
  - Likert score-4=poor reliability( single measure ICC = 0.248; reader-1,single measure ICC = 0.404; reader-2)

# CONCLUSION

- To the best of our knowledge we firstly examined the ability of prostate MRI in determining pathological RD.
- We suggested that prostate MRI shows promising results on determining pathological RD in patients with gross measurable tumor on EPE assesment.
- In conclusion, prostate MRI may have a potential for assessing the degree of EPE as a prognostic marker.