Focal prostate brachytherapy – our experience so far

Bing Wei & Thang Nguyen Charge Radiation Therapists – Icon Cancer Care





- Rationale for focal brachytherapy
- Our focal experience
- Future direction





Side effects from active prostate cancer treatment

- Radical Prostatectomy invasive, risk of impotence, urinary incontinence (1)
- Radiation Therapy rectal injury, impotence, bladder dysfunction post treatment QoL similar or worse than RP (1)
- Hormone Therapy anaemia, osteoporosis, impotence, cognitive impairment and increases risk of sudden death (1)
- Brachytherapy incontinence, impotence, retention, irritation (2)





Rationale for focal treatment

- Early prostate cancer should have a treatment that is *curative, minimally or non invasive, single session* without debilitating side effects
- "Lumpectomy" for favourable breast cancer
- Challenging in prostate cancer due to the multi-focal nature of the disease, *although* shown to be less of a concern than previously thought (1)
- Localised destruction of the largest focus area could potentially reduce local progression and metastasis (3)





Rationale for focal treatment cont.

- Recent advances in medical imaging *may* have overcome these obstacles (2):
- mpMRI used as non-invasion tool in intra-prostatic location of prostate cancer
- T2 MRI is able to provide clinicians with a 3D map of the target tissue







What is focal brachytherapy

- An approach that maximises tumour control, minimises side effects
- Treating the only the portion of gland which contains significant cancer
- Has the potential to reduce injury to adjacent organs whilst maintaining excellent oncological outcome







Patient selection

- Patients with low or moderate risk
- Index nodule identified on MRI matches template biopsy
- Patients with dominate Gleason 4 or clinical stage T2b greater are *excluded*

79 years old	68 years old	63 years old	75 years old
PSA 6.8	PSA 7.5	PSA 5.1	PSA 8.5
3 + 4 = 7	3 + 4 =7	3 + 4 = 7	3 + 4 = 7
Left anterior zone	Right anterior TZ	Right anterolateral mid gland	Right anterior TZ











Methodology

- 145Gy encompassing PTV
- Range of activities from 0.311mCi 0.500mCi

Priority	Target	Dose-volume objectives (145Gy
1	PTV	V100% ≥ 98%
		V 150 ≤ 52-70%
		D90 > prescription dose





Normal tissue tolerance/ dose constraints

Priority	Normal tissue	Dose-volume constraints (145Gy)
1	Urethra	D10% < 150% of prescription dose
		D30% < 130% of prescription dose
2	Rectum	D2cc < 145Gy
		D0.1cc < 200Gy





	Patient 1	Patient 2	Patient 3	Patient 4
V100	99.25%	100%	99.17%	99.46%
V150	76.25%	79.26%	91.3%	80.27%
D90	184Gy	199Gy	225Gy	195Gy







	Patient 1	Patient 2	Patient 3	Patient 4
V100	99.25%	100%	99.17%	99.46%
V150	76.25%	79.26%	91.3%	80.27%
D90	184Gy	199Gy	225Gy	195Gy
D90	184Gy	199Gy	225Gy	195







	Patient 1	Patient 2	Patient 3	Patient 4
V100	99.25%	100%	99.17%	99.46%
V150	76.25%	79.26%	91.3%	80.27%
D90	184Gy	199Gy	225Gy	195Gy







	Patient 1	Patient 2	Patient 3	Patient 4
V100	99.25%	100%	99.17%	99.46%
V150	76.25%	79.26%	91.3%	80.27%
D90	184Gy	199Gy	225Gy	195Gy
		+ + + + + + + +	++++	







Preplan - OAR

	Patient 1	Patient 2	Patient 3	Patient 4
Urethra D10%	113.94%	148.72%	61.72%	117.62%
Urethra D30%	107.49%	140.44%	39.56%	93.59%
Rectum D2cc	21.13%	18.67%	5.37%	9.59%
Rectum D0.1cc	39.41%	32.53%	10.88%	13.22%





Preplan - OAR

	Patient 1	Patient 2
Urethra D10%	113.94%	148.72%
Urethra D30%	107.49%	140.44%
Rectum D2cc	21.13%	18.67%
Rectum D0.1cc	39.41%	32.53%







Proportion of prostate treated



Post plan

Recommended evaluated postoperative dosimetric parameters:

- V100
- V150
- V200
- D90
- Urethral doses should include UV125, UV150, UD50, UD30, UD5 and/or maximum and minimum dose
- Rectal doses cubic centimeters of rectum which received ≥ prescription dose (RV100)





Post plan results - PTV

	Patient 1	Patient 2	Patient 3	Patient 4
V100	89.59%	89.03%	91.70%	72.25%
V150	42.70%	47.70%	68.76%	48.49%
V200	19.97%	24.04%	47.23%	20.80%
D90	127Gy	143.14Gy	150.58Gy	96.99Gy





Post plan results - OARs

Urethra	Patient 1	Patient 2	Patient 3	Patient 4
UV150	26.34%	29.29%	0%	10.59%

Rectum				
V100 <1.3cm ³	0cm³	0.04cm³	0cm³	0cm³







Patient		Baseline PSA	3	6	9	12	15	18	21	24	FU in 27month
	1	6.800		0.560	0.440	0.300	0.210	0.090	0.050	0.070	0.060
	2	7.500		2.190		1.400					
	3	8.100		1.490		0.920				1.000	
	4	5.100		1.900		1.400		1.500			





Conclusion

- Valid option
- Our experience has been positive smooth and straight forward process with the aid of technology that incorporates MRI and ultrasound fusion
- Routine follow up with PSA, MRI and repeat template biopsy
- Challenges remain in how to interpret PSA result and surveillance of the untreated gland





Future direction

- The success of focal therapy relies upon the accuracy of preintervention diagnostics
- Prostate specific membrane antigen (PSMA) scan has the ability to highlight the presence of active local disease (4)
- Formalise a phase II trial employing focal LDR brachytherapy targeting the index lesion incorporating the use of PSMA scans





Acknowledgments

- All at Icon Cancer Centre
- All work conducted at Epworth Radiation Oncology





References

- 1. Potters L, Morgenstern C, Calugaru E, et al. 12-Year Outcomes Following Permanent Prostate Brachytherapy in Patients With Clinically Localized Prostate Cancer. J. Urology, 2005; 173:1562-1566
- 2. (104) Onik G, Miessau M et al. Three dimensional prostate mapping biopsy has a potentially significant impact on prostate cancer management. Journal Clinical Oncol. 2009;27:4321-4326
- 3. Al-Qaisieh B, Mason J, Bownes P et.al. *Dosimetry modelling for focal low-dose-rate prostate brachytherapy*. IJROBP 2015 Jul 12;92(4):787-93.
- 4. Afshar-Oromieh A, et al. *The diagnostic value of PET/CT imaging with the ⁶⁸Galabelled PSMA ligand HBED-CC in the diagnosis of recurrent prostate cancer*. *European Journal of Nuclear Medicine and Molecular Imaging*. 2015.



