Advanced Technology Consortium (ATC)

CREDENTIALING PROCEDURES FOR PROSTATE IMPLANT PROTOCOLS

FACILITY QUESTIONNAIRE

Institutions wishing to enter patients onto RTOG, NCCTG or ACOSOG protocols that include permanent prostate brachytherapy implants must be credentialed prior to participation in the study. Because these procedures require a team effort, the institution, the radiation oncologist and radiation physicist are credentialed as a team.

RTOG and NCCTG:

If there is more than one radiation oncologist who wishes to be credentialed, please refer to the letter from Drs. Kuban and Ibbott (RTOG) or Drs. Pisansky and Ibbott (NCCTG). The treatment team is required to submit three items.

- 1) Facility Questionnaire and Reference Cases
- 2) Knowledge Assessment Form
- 3) Data for a recent prostate implant performed by the radiation oncologist and physicist using the technique proposed for the RTOG or NCCTG protocol.

RTOG/NCCTG requires that the prostate team must have performed at least 10 TRUS guided prostate implants. At the bottom of the ATC Knowledge Assessment Form you will be asked to attest to this by your signature.

RTOG and NCCTG packages are to be submitted to:

Attention: Irene Harris Radiological Physics Center 1515 Holcombe Blvd. – Box 547 Houston, TX 77030

ACOSOG: The treatment team is required to submit the following three items:

- 1) TIPPB Facility Questionnaire and Reference Cases
- 2) A case list of 15 permanent prostate seed implants which must have been performed over the last 12 months. Use Radiation Therapy Skill Verification Form (www.qarc.org)
- 3) Data for the 3 most recent cases performed by the brachytherapy team (See Radiation Therapy Skill Verification Form for submission requirements)

ACOSOG packages are to be submitted to:

Physics Division Quality Assurance Review Center 825 Chalkstone Ave. Providence, RI 02908-4735

An institution that has been credentialed for one radiation source model does not need to submit the complete package to become credentialed for a second source model. It is only necessary to submit the two reference cases, performed with the new source model. Similarly, changing to a different treatment planning system requires re-credentialing, and only the two reference cases need to be submitted.

Institutions that have the capability to submit treatment plans electronically will be expected to transmit data in digital form to the Image-Guided Therapy Center (ITC) in St. Louis (http://itc.wustl.edu/). All RTOG participants are expected to submit data electronically.

For any questions, please contact the RPC (<u>http://rpc.mdanderson.org</u>) at (713) 745-8989 or QARC (<u>www.qarc.org</u>) at (401) 456-6500.

Allow at least 4 weeks for review and approval. Incomplete submissions will take longer.

ATC CREDENTIALING PROCEDURES FOR PROSTATE IMPLANT PROTOCOLS FACILITY QUESTIONNAIRE

I. Radiation Oncology Facility: Study Group:	RTF #: RTOG / NCCTG / ACOSOG #:	
Is this Facility also known by other name(s)? If so, please provide:	
•	ered:	
Address:		
PERSONNEL CONTACT INFORMATION		
A. Radiation Oncologist Responsible for Implant Pat	ents	
Name:	Phone:	
Address:	Fax:	
	E-mail:	
B. Chair/Chief of Radiation Oncology		
Name:	Phone:	
Address:	Fax:	
	E-mail:	
C. Physicist Responsible for Implant Patients		
Name:	Phone:	
Address:	Fax:	
	E-mail [.]	

Prostate Brachytherapy QA

D. Dosimetrist Responsible for Implant Patients

N	ame: Phone:
Add	ress: Fax:
	E-mail:
F Da	ta Manager Responsible for Implant Patients
N	ame: Phone:
Add	ress: Fax:
	E-mail:
	perience of personnel: For the Radiation Oncologist named above
A.	-
	How many ultrasound guided prostate implants have been performed in the past
	6 months? 12 months? career total?
-	Has this person's skill been verified? by RTOG? by NCCTG? by ACOSOG? date:
В.	For the Physicist named above
	How many ultrasound guided prostate implants have been preplanned using ultrasound in the past 6 months? 12 months? career total?
	How many ultrasound guided prostate implants have been evaluated with post implant CT in the past 6 months? 12 months? career total?
	Has this person's skill been verified? by RTOG? by NCCTG? by ACOSOG? date:
III. Ec	juipment:
	Ultrasound unit (vendor and model):
	CT scanner (vendor and model):
	Treatment planning system
	eplan:
	Vendor and version:
	How are ultrasound images entered for pre planning? videotape digitized Other (explain):

How are prostate and normal tissue contours entered?	
Defined on planning system 🗌 defined on ultrasound unit and i	nput as above 🗌
Other (explain):	
Is a point source approximation used? Yes No	
If yes, do you use an: anisotropy constant 🗌 anisotropy factors	
If not, explain your procedures for determining and accounting for seed or	ientation.
st Implant Plan:	
Vendor and version:	
How are the CT images entered for post planning? CD tape op	tical disc 🗌
The are the CT intrages entered for post plaining : CD tape of	
	Dhicaliv via netwo
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	•
digitized from hardcopyelectronically from the scannerelectronically from	•
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digitized from hardcopy electronically from the scanner electronically from the scanner Other (explain):	/e 🗌
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digitized from hardcopy electronically from the scanner electronically from the scanner Other (explain):	/e 🗌
digitized from hardcopy electronically from the scanner Other (explain): How are prostate and normal tissue contours entered? Defined on planning system Defined on planning system Other (explain): Dose calculation matrix resolution is mm x mm.	/e 🗌
digitized from hardcopy □ electronically from the scanner □ electronically from the scanner □ Other (explain):	/e 🗌
digitized from hardcopy □ electronically from the scanner □ electronically from the scanner □ electronically from the scanner □ Other (explain):	/e 🗌

Ε.

- D. Radiation Sources:
 - ¹²⁵I: Vendor/Model: ______Typical source strength/seed: ______Typical source strength/seed: ______Typical source strength/seed: ______
 ¹⁰³Pd: Vendor/Model: ______Typical source strength/seed: ______Typical source strength/seed: _______Typical source strength/seed: _______
 Special ordered for each patient ______Prepared from in-house inventory
 Implant Technique: ______Loose seeds (e.g., Mick applicator) ______Disposable preloaded needles (seeds with spacers interspersed) _______Stranded products or dissolvable sutures
- F. Do you immobilize the prostate during the implant procedure, for example, with anchor needles?

IV. Quality Assurance Procedures: (attach additional sheets if necessary)

- A. Source strength verification:
 - 1. Dosimetry system used for in-house verification of seed activity:

Vendor: _____ Model: _____

- 2. How is the calibration of this system directly traceable to NIST? (Attach copies of ADCL certificates)
- 3. What are the QA procedures to verify that the calibration of this system has not changed?

- 4. For each seed model, what is the NIST calibration date to which your chamber calibration is traceable?
- 5. How frequently are these QA procedures performed?
- 6. Describe your measurement technique for verifying seed strengths of individual patients.

	7.	Number of seeds assayed per patient:% orseeds					
	8.	What is your criterion for agreement with the vendor? +/-5% \Box , +/-7% \Box , +/-10% \Box ,					
		Other (explain)					
	9.	What seed strength is used for treatment planning? your own measurements 🗌 vendor 🗌					
		Other (explain)					
В.	So	urce accounting:					
	1. Are radiographs taken at the completion of the implant? Yes 🗌 No						
		If yes: AP lateral oblique stereo other:					

2. Describe procedures used to account for all seeds at the time of implant:

3. Describe procedures used to account for all seeds at the time of post implant planning:

4. Describe techniques used to identify seeds and avoid identifying the same seed on multiple CT slices:

5. What is the discrepancy limit for unaccounted seeds and what action do you take if the discrepancy exceeds the limit?

- C. Other dosimetry and QA procedures:
 - 1. Describe any calculations done at the time of commissioning to verify the accuracy of the computer generated treatment plan:

2. Describe your method for ensuring that the dosimetric parameters you use are consistent with the NIST calibration of the source and your calculation method (point source approximation vs. line source):

3. Describe any other procedures followed to assure that the dose calculations are in accordance with the requirements of the protocol:

3. Describe any other quality assurance procedures pertinent to these brachytherapy procedures:

Reference Cases

Please calculate and attach isodose distributions for the single seed and geometric implant described below. Sources should be the model ¹²⁵I or ¹⁰³Pd seeds (from the list of models complying with the AAPM prerequisites – see <u>http://rpc.mdanderson.org</u>) that you choose to treat your patients, with source strength specified exact at the beginning of the implant. Do the calculations as you would do them clinically using the TG-43 dosimetry, detailing any assumptions necessary.

Case 1: A single seed, strength 0.5U (μGy m² h⁻¹) (use 2.5 U for ¹⁰³Pd): If your software allows and you use a line source approximation, calculate both in the longitudinal and mid-transverse planes of the seed. Please submit isodose lines from 0.2 to 100 Gy. (Lines 0.2, 0.5, 1, 5, 10, 50, & 100 Gy are preferred.)

Dosimetry Calculations:

Write below the equation that will be used for hand calculating the instantaneous dose-rate to an arbitrary point from a single seed in the TG-43 formalism. (If possible give notations used by your treatment planning computer). The intent is for you to be able to verify that the values of various parameters in your treatment planning system are the same as in TG-43.

Define the variables in the equation:

For each seed model used to treat patients on this protocol, submit the data used by your treatment planning computer for the following parameters:

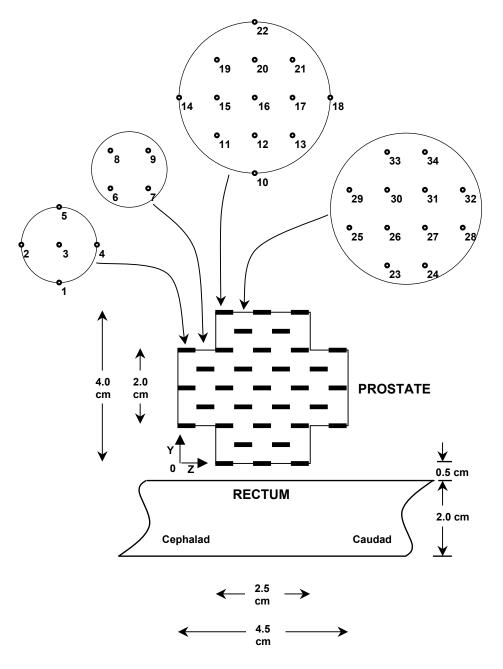
- Dose rate constant (Λ)
- Anisotropy constant (ϕ) and/or factors
- Radial dose function
- The units of S_K are :
- Do your ¹²⁵I dose calculations agree with TG-43 to within <u>+</u>5% from 5-70 mm? Yes No
- Do your ¹⁰³Pd dose calculations agree with TG-43 to within <u>+</u>5% from 5-50 mm? Yes □ No □

Case 2: The diagram on the following page represents a cylindrical prostate and rectum. Assume the dimensions of the prostate and rectum in the drawing to be both the PTV and ETV.

The purpose of this reference case is principally to verify the consistency of your calculations with TG-43. Perform calculations for the implant shown in the diagram using seeds of source strength 0.5 U (μ Gy m² h⁻¹) (use 2.5 U for ¹⁰³Pd).

- Submit isodose distributions in all axial planes. (0.5 cm spacing)
- Submit the following isodose lines: 290, 217, 145, 130, 73 and 36 Gy
- Submit the volumes of prostate and rectum as calculated by the treatment planning system.
- Submit integral dose-volume histogram (DVH) tables in 10 Gy increments for:
 - (i) Planning target volume (PTV).
 - (ii) Rectum in prostate region.

<u>Note</u>: If you wish to use more than one of the approved seed models, please submit cases 1 and 2 for each seed model.



REFERENCE CASES – PAGE 2

Prostate Reference Case Case #2 3/98 (x,y,z) Coordinates of center of seeds

Seed #	х	У	z1	z2	z3	z4
1	2	1	0.25			4.25
2	1	2	0.25			4.25
3	2	2	0.25			4.25
4	3	2	0.25			4.25
5	2	3	0.25			4.25
6	1.5	1.5	0.75			3.75
7	2.5	1.5	0.75			3.75
8	1.5	2.5	0.75			3.75
9	2.5	2.5	0.75			3.75
10	2	0	1.25	2.25	3.25	
11	1	1	1.25	2.25	3.25	
12	2	1	1.25	2.25	3.25	
13	3	1	1.25	2.25	3.25	
14	0	2	1.25	2.25	3.25	
15	1	2	1.25	2.25	3.25	
16	2	2	1.25	2.25	3.25	
17	3	2	1.25	2.25	3.25	
18	4	2	1.25	2.25	3.25	
19	1	3	1.25	2.25	3.25	
20	2	3	1.25	2.25	3.25	
21	3	3	1.25	2.25	3.25	
22	2	4	1.25	2.25	3.25	
23	1.5	0.5	1.75	2.75		
24	2.5	0.5	1.75	2.75		
25	0.5	1.5	1.75	2.75		
26	1.5	1.5	1.75	2.75		
27	2.5	1.5	1.75	2.75		
28	3.5	1.5	1.75	2.75		
29	0.5	2.5	1.75	2.75		
30	1.5	2.5	1.75	2.75		
31	2.5	2.5	1.75	2.75		
32	3.5	2.5	1.75	2.75		
33	1.5	3.5	1.75	2.75		
34	2.5	3.5	1.75	2.75		

Total # of seeds: (5x2 + 4x2 + 13x3 + 12x2) = 81