

8/28/2013

U.S. Nuclear Regulatory Commission Materials Licensing Section 2442 Warrenville Road, Suite 210 Lisle, IL 60532 – 4352

Dear Sir or Madam:

Hancock Regional Hospital would like to amend its radioactive materials license, number 13-16730-01, to change the radiation safety officer from Mary Lynn Wolfe, M.D. to Justin P. Chang, M.D. Enclosed is a 313a (RSO) with the necessary supporting documentation.

In addition we would like to add Justin P. Chang, M.D. as an authorized user for 35.100, 35.200 and 35.392. Enclosed is a 313a (AUD) and 313a (AUT) with the necessary supporting documentation.

In addition, Hancock Regional Hospital would like to notify the NRC that the administration and Radiation Safety Committee has approved Justin P. Chang, M.D. as a temporary RSO pending this amendment.

Should there be any questions concerning this request, please feel free to contact Mr. Timothy Greist, DABR of Medical Physics Consultants at tgreist@mpcphysics.com or 310-975-4149.

Sincerely,

Bobby Keen, FACHE
President and CEO

Sept. 9, 2013 Date

> 801 North State St. Greenfield, IN 46140

Phone (317) 462.5544



RSO / EXECUTIVE MANAGEMENT LETTER OF UNDERSTANDING

8/28/2013

Justin P. Chang, M.D. Radiation Safety Officer Hancock Regional Hospital 801 N. State St. Greenfield, IN 46140

Dear Dr. Justin P. Chang, RSO:

You have been appointed the Radiation Safety Officer (RSO) of this facility for our United States Nuclear Regulatory Commission Material License. This "Letter of Understanding" is prepared to comply with Title 10 Code of Federal Regulations (CFR) Part 35.24(b). This section of the regulations requires that you agree in writing to the following:

- Assume responsibility for implementing the Radiation Protection Program.
- Ensure that radiation safety activities are being performed in accordance with our own approved procedures and all regulatory requirements.

Furthermore, in compliance with 10 CFR 35.24 (e), (g), the executive management of this facility agrees to provide you as RSO:

- > Specific written notation of your authority, duties and responsibilities, see attached.
- > Sufficient authority, organizational freedom, time, resources and management prerogative to:
 - 1. Identify radiation safety problems;
 - 2. Initiate, recommend, or provide corrective actions;
 - 3. Stop unsafe operations; and,

C. Kan

4. Verify implementation of corrective actions.

Our signatures noted below will attest to the issues noted above. Please make a copy of this document for your files and return the original to my attention.

Sincerely,

Bobby Keen, FACHE

President and CEO

Justin P. Chang, M

Radiation Safety Officer

801 North State Date

Phone (317) 462.5544

RADIATION SAFETY OFFICER AUTHORITY, DUTIES AND RESPONSIBILITIES

The Radiation Safety Officer (RSO) shall:

- 1. Have the authority to implement the Radiation Protection Program as referenced in 10 CFR 20.1101.
- 2. Have the authority, organizational freedom, time, resources, and management prerogative to:
 - a. Identify radiation safety problems;
 - b. Initiate, recommend or provide corrective actions,
 - c. Stop unsafe operations; and,
 - d. Verify implementation of corrective actions.
- 3. Investigate deviations from the radiation safety practices approved by facility management and/or the Radiation Safety Committee, if applicable.
- 4. Collect in a centralized location, executive management approved procedures that can include policy and technical issues which, would makeup the Radiation Protection Program as follows:
 - Authorization for the purchase of radioactive material.
 - b. Receipt and opening of packages containing radioactive material.
 - Storage of radioactive material.
 - d. Inventory control of radioactive material.
 - e. Safe use of radioactive material.
 - f. Emergency procedures in the event of loss, theft, etc.
 - g. Periodic radiation surveys and wipe tests
 - h. Checks of radiation survey and other radiation safety instruments.
 - Disposal of radioactive material.
 - j. Personnel training of those who work in or frequent areas of radioactive material use or storage.
- 5. Oversee a record system of the Radiation Protection Program per 10 CFR 20.2102 to include at least the following:

The provisions of the Radiation Protection Program until the license is terminated by the NRC such as:

- a. All records, reports, written policies and procedures required by regulatory agencies concerning radioactive material.
- b. A copy of the regulations governing the possession, use and disposal of licensed material, such as Title 10 Code of Federal Regulations.

Audits and other reviews of the Radiation Protection Program content and implementation for a period of three (3) years after the record is made.

- 6. Periodically evaluate "action levels" for continued appropriateness to ensure compliance with 10 CFR 20.1501 and 1502 for the following:
 - a. Personnel exposure investigation levels
 - b. Area surveys dose rate and contamination levels
 - c. Bioassays, if necessary
 - d. Radioactive effluent concentrations, if necessary
- 7. Review the following Radiation Protection Program records, if applicable:
 - a. Sealed source inventories
 - b. Sealed source leak tests
 - c. Dose calibrator linearity tests
 - d. Dose calibrator accuracy tests
 - e. Dose calibrator geometrical variation tests
 - f. Occupational radiation exposure reports
 - g. Medical event documentation
 - h. Spill / incident reports for cause and corrective action
 - i. Dose rate and contamination survey results
 - j. Changes in the radiation safety program
- 8. Ensure the use of reasonable practices and controls to strive to maintain doses to workers and to the public are ALARA, in compliance with 10 CFR 20.1101(b).
- 9. Review with facility management at least annually of the content of the Radiation Protection Program and determine if the written program is being implemented in compliance with 10 CFR 20.1101(c).
- 10. Ensure as a part of the ALARA effort that individual members of the public shall not receive a Total Effective Dose Equivalent (TEDE) of more than 10 mrem (0.1 mSv) per year from airborne radioactive material releases as per 10 CFR 20.1101(d) as necessary.
- 11. Be a member of the Radiation Safety Committee (RSC), if applicable, that will oversee all uses of byproduct material permitted by the license as per 10 CFR 35.24(f).

NRC FORM 313A (AUT)

U.S. NUCLEAR REGULATORY COMMISSION

(05-2012)

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION

APPROVED BY OMB: NO. 3150-0120 EXPIRES: (05/31/2015)

	[10 CFR 35.390, 35.392, 35.394, and 35.396]																
Nai	Name of Proposed Authorized User						State or 1	erritory	Where	License	ed						
Jus	tin C	hang, Ml	D						Indiana								
Re	ques	ted Auth	norizatio	on(s) (c	heck all t	hat ap	ply):										
	<u> </u>	35.300	Use of	unseal	ed bypro	duct m	naterial fo	r whic	ch a writte	en direc	tive is	require	ed				
,	OR																
	✓ 3	35.300			ation of squerels (3		n iodide I- curies)	-131 re	equiring a	a writter	n direc	tive in	quantit	ties les	ss than	or equal	l to
	35.300 Oral administration of sodium iodide I-131 requiring a written directive in quantities greater than 1.22 gigabecquerels (33 millicuries)																
	35.300 Parenteral administration of any beta-emitter, or photon-emitting radionuclide with a photon energy leads than 150 keV for which a written directive is required					energy le	ess										
	3	35.300	Parent	eral adı	ministrati	on of a	any other	radior	nuclide fo	or which	a writ	ten dir	ective i	is requ	uìred		
							ART I 1										
*	* Training and Experience, including board certification, must have been obtained within the 7 years preceding the date of application or the individual must have related continuing education and experience since the required training and experience was completed. Provide dates, duration, and description of continuing education and experience related to the uses checked above. 1. Board Certification a. Provide a copy of the board certification. b. For 35.390, provide documentation on supervised clinical case experience. The table in section 3.c. may be used to document this experience. c. For 35.396, provide documentation on classroom and laboratory training, supervised work experience, and supervised clinical case experience. The tables in sections 3.a., 3.b., and 3.c. may be used to document this experience. d. Skip to and complete Part II Preceptor Attestation.																
	2. 9	Current	35.300	, 35.40	0, or 35.	600 Au	uthorized	l User	r Seeking	Addit	ional <i>i</i>	Author	izatio	ņ			
	a. /	Authoriz	ed Use	r on Ma	iterials Li	cense						under	the re	quiren	nents be	elow or	
		equivale	ent Agre	eement	State red	quirem	ents (che	eck all	that app	ly):							
		35.3	390		35.392		35.394		35.4	490		35.69	0				
	b. If currently authorized for a subset of clinical uses under 35.300, provide documentation on additional required supervised case experience. The table in section 3.c. may be used to document this experience. Also provide completed Part II Preceptor Attestation.																
	doc	cumenta se exper	tion on ience.	classro The tab	om and l	aborat ctions	35.690 a tory traini 3.a., 3.b. Attestatio	ng, su , and :	pervised	work ex	xperie	nce, ar	nd supe	ervise	d clinica	al	

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued)

3. Training and Experience for Proposed Authorized User (continued)

b. Supervised Work Experience (continued)

Supervising In	dividual	License/Permit Number listing supervising individual as an authorized user				
Darlene Fink-	Bennett, MD	21-01333-01				
Supervising apply)**:	individual meets the requirements below, o	r equivalent Agreement State requirements (check all that				
35.390	With experience administering dosages of:					
✓ 35.392✓ 35.394	✓ Oral Nal-131 requiring a written direct gigabecquerels (33 millicuries)	Nal-131 requiring a written directive in quantities less than or equal to 1.22 ecquerels (33 millicuries)				
	✓ Oral Nal-131 in quantities greater that	n 1.22 gigabecquerels (33 millicuries)				
✓ 35.396	Parenteral administration of beta-emi energy less than 150 keV requiring a	itter, or photon-emitting radionuclide with a photon written directive is required				
	Parenteral administration of any other	r radionuclide requiring a written directive				
** Supervising Authorized User must have experience in administering dosages in the same dosage category or categories as the individu requesting authorized user status.						

c. Supervised Clinical Case Experience

If more than one supervising individual is necessary to document supervised work experience, provide multiple copies of this page.

Description of Experience	Number of Cases Involving Personal Participation	Location of Experience/License or Permit Number of Facility	Dates of Experience*
Oral administration of sodium iodide I-131 requiring a written directive in quantities less than or equal to 1.22 gigabecquerels (33 millicuries)	4	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	3/6/08; 3/12/08; 4/1/08; 4/25/08
Oral administration of sodium iodide I-131 requiring a written directive in quantities greater than 1.22 gigabecquerels (33 millicuries)	3	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	4/23/10; 4/23/10; 4/27/10
Parenteral administration of any beta-emitter, or photon-emitting radionuclide with a photon energy less than 150 keV for which a written directive is required			
Parenteral administration of any other radionuclide for which a written directive is required (List radionuclides)			

NRC FORM 313A (AUT) (05-2012)		U.S. NUCLEAR REGULATORY COMMISSION
' '	ED USER TRAINING AND EXPERIE	ENCE AND PRECEPTOR ATTESTATION (continued)
3. <u>Training and E</u>	xperience for Proposed Authorized	<u>d User</u> (continued)
c. Supervised (Clinical Case Experience (continued)	
Supervising Indivi	dual	License/Permit Number listing supervising individual as an authorized user
Darlene Fink-Ber	inett, MD	21-01333-01 William Beaumont Hospital
Supervising indiapply)**:	vidual meets the requirements below	, or equivalent Agreement State requirements (check all that
✓ 35.390 V	Vith experience administering dosage	es of:
✓ 35.392 ✓ 35.394	 Oral Nal-131 requiring a written dia gigabecquerels (33 millicuries) 	rective in quantities less than or equal to 1.22
▼ 35.394 ▼ 35.396	☑ Oral Nal-131 in quantities greater	than 1.22 gigabecquerels (33 millicuries)
35.390	Parenteral administration of beta-e energy less than 150 keV requiring	emitter, or photon-emitting radionuclide with a photon g a written directive is required
	Parenteral administration of any of	her radionuclide requiring a written directive
	horized User must have experience in adminis orized user status.	stering dosages in the same dosage category or categories as the individual
d. Provide com	pleted Part II Preceptor Attestation.	
	PART II – PRECI	EPTOR ATTESTATION
individual as	long as the preceptor provides, direc	eceptor. The preceptor does not have to be the supervising ts, or verifies training and experience required. If more than ce, obtain a separate preceptor statement from each.
	the boxes below, the preceptor is atte ought and not attesting to the individe	esting that the individual has knowledge to fulfill the duties of ual's "general clinical competency."
First Section Check one of the fo	llowing for each requested authori	zation:
For 35.390:		
Board Certif	ication	
I attest th	Name of Proposed Authorized User	has satisfactorily completed the training and experience
requireme	ents in 35.390(a)(1).	
		OR
Training and	1 Experience	
✓ I attest th		has satisfactorily completed the 700 hours of training
	Name of Proposed Authorized User	
	rience, including a minimum of 200 h	ours of classroom and laboratory training, as required by

NRC FORM 313A (AUT) (05-2012) PAGE 4

NRC FORM 313A (AUT) (05-2012) PAGE 5

Parenteral administration of any other radionuclide requiring a written directive

Darlene Fink-Bennett, MD

License/Permit Number/Facility Name

21-01333-01 William Beaumont Hospital, Royal Oak, MI

(248) 898-4126

08/29/2013

NRC FORM 313A (AUD)

U.S. NUCLEAR REGULATORY COMMISSION

(05-2012)

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION

(for uses defined under 35,100, 35,200, and 35,500)

APPROVED BY OMB: NO. 3150-0120 EXPIRES: (05/31/2015)

[10 CFR 35.190	•						
Name of Proposed Authorized User		State or Territory Where License	ed				
Justin Chang, MD		Indiana					
Requested Authorization(s) (check all that	t apply)						
√ 35.100 Uptake, dilution, and excretion	studies						
√ 35.200 Imaging and localization studie	es						
35.500 Sealed sources for diagnosis (specify device)						
		G AND EXPERIENCE three methods below)					
* Training and Experience, including boa the date of application or the individual the required training and experience was education and experience related to the	must have obtaine as completed. Pro	ed related continuing education ovide dates, duration, and described	n and experie	ence since			
1. Board Certification							
a. Provide a copy of the board certific	cation.						
 b. If using only 35.500 materials, stop Preceptor Attestation. 	here. If using 35	.100 and 35.200 materials, sk	ip to and com	iplete Part II			
2. <u>Current 35.390 Authorized User</u>	Seeking Addition	nal 35.290 Authorization					
 Authorized user on Materials Lice State requirements seeking autho 		meeting 10 CFR 35.3	90 or equival	ent Agreement			
 b. Supervised Work Experience. (If more than one supervising indiscopies of this section.) 	vidual is necessar	y to document supervised wor	k experience,	provide multiple			
Description of Experience		f Experience/License or : Number of Facility	Clock Hours	Dates of Experience*			
Eluting generator systems appropriate for the preparation of radioactive drugs for imaging and localization studies, measuring and testing the eluate for radionuclidic purity, and processing the eluate with reagent kits to prepare labeled radioactive drugs							
	Total Hours of Experience:						
Supervising Individual License/Permit Number listing supervising individual as an authorized user							
Supervisor meets the requirements be 35.290 35.390 + ger	·	t Agreement State requiremer	its (check all i	that apply).			

(05-2012)

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued)

√ 3. Training and Experience for Proposed Authorized User

a. Classroom and Laboratory Training.

Description of Training	Location of Training	Clock Hours	Dates of Training*				
Radiation physics and instrumentation	William Beaumont Hospital, Royal Oak, Michigan	35	7-1-2007 - 6-30-2012				
Radiation protection	William Beaumont Hospital, Royal Oak, Michigan	20	7-1-2007 - 6-30-2012				
Mathematics pertaining to the use and measurement of radioactivity	William Beaumont Hospital, Royal Oak, Michigan	6	7-1-2007 - 6-30-2012				
Chemistry of byproduct material for medical use (not required for 35.590)	William Beaumont Hospital, Royal Oak, Michigan	24	7-1-2007 - 6-30-2012				
Radiation biology	William Beaumont Hospital, Royal Oak, Michigan	10	7-1-2007 - 6-30-2012				
Total Hours of Training: 95							

Supervised Work Experience (completion of this table is not required for 35.590). (If more than one supervising individual is necessary to document supervised work experience, provide multiple copies of this section.)

Supervised Work Experience	Total Hours of Experience:				
Description of Experience Must Include:	Location of Experience/License or Permit Number of Facility	Confirm	Dates of Experience*		
Ordering, receiving, and unpacking radioactive materials safely and performing the related radiation surveys	William Beaumont Hospital, Royal Oak, Michigan	✓ Yes	7-1-2007 - 6-30-2012		
Performing quality control procedures on instruments used to determine the activity of dosages and performing checks for proper operation of survey meters	William Beaumont Hospital, Royal Oak, Michigan	✓ Yes	7-1-2007 - 6-30-2012		

(05-2012)

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued)

		•
3.	Training and Experience for Proposed Authorized User (continued)	

h	Suponicod	Mork	Evnorionco	(continued)

Description of Experience Must Include:	Location of Experience/License or Permit Number of Facility	Confirm	Dates of Experience*	
Calculating, measuring, and safely preparing patient or human research subject dosages	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	✓ Yes	7-1-2007 - 6-30-2012	
Using administrative controls to prevent a medical event involving the use of unsealed byproduct material	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	✓ Yes	7-1-2007 - 6-30-2012	
Using procedures to contain spilled byproduct material safely and using proper decontamination procedures	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	✓ Yes	7-1-2007 - 6-30-2012	
Administering dosages of radioactive drugs to patients or human research subjects	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	✓ Yes	7-1-2007 - 6-30-2012	
Eluting generator systems appropriate for the preparation of radioactive drugs for imaging and localization studies, measuring and testing the eluate for radionuclidic purity, and processing the eluate with reagent kits to prepare labeled radioactive drugs	William Beaumont Hospital, Royal Oak, Michigan 21-01333-01	✓ Yes ☐ No	7-1-2007 - 6-30-2012	
Supervising Individual	License/Permit Number listing	supervising indi-	vidual as an	
Darlene Fink-Bennett, MD 21-01333-01 William Beaumont Hospital				
Supervisor meets the requirements bel	low, or equivalent Agreement State requireme	ents (check one	∍).	
35.190 35.290	35.390 3 5.390 + generator experien	ice in 35.290(c	(1)(ii)(G)	

c. For 35.590 only, provide documentation of training on use of the device.

Device	Type of Training	Location and Dates

d. For 35.500 uses only, stop here. For 35.100 and 35.200 uses, skip to and complete Part II Preceptor Attestation.

NRC FORM 313A (AUD) U.S. NUCLEAR REGULATORY COMMISSION (05-2012) AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued) PART II - PRECEPTOR ATTESTATION Note: This part must be completed by the individual's preceptor. The preceptor does not have to be the supervising individual as long as the preceptor provides, directs, or verifies training and experience required. If more than one preceptor is necessary to document experience, obtain a separate preceptor statement from each. (Not required to meet training requirements in 35.590) By checking the boxes below, the preceptor is attesting that the individual has knowledge to fulfill the duties of the position sought and not attesting to the individual's "general clinical competency." First Section Check one of the following for each use requested: For 35, 190 **Board Certification** has satisfactorily completed the requirements in I attest that Name of Proposed Authorized User 10 CFR 35.190(a)(1) and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under 10 CFR 35.100. OR Training and Experience ✓ I attest that Justin Chang, MD has satisfactorily completed the 60 hours of training and Name of Proposed Authorized User experience, including a minimum of 8 hours of classroom and laboratory training, required by 10 CFR 35.190(c)(1), and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under 10 CFR 35.100. For 35.290 **Board Certification** has satisfactorily completed the requirements in I attest that Name of Proposed Authorized User 10 CFR 35.290(a)(1) and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under 10 CFR 35.100 and 35.200. Training and Experience ✓ I attest that Justin Chang, MD has satisfactorily completed the 700 hours of training Name of Proposed Authorized User and experience, including a minimum of 80 hours of classroom and laboratory training, required by 10 CFR 35.290(c)(1), and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under 10 CFR 35.100 and 35.200. Second Section Complete the following for preceptor attestation and signature:

I meet the requirements below, or equivalent Agreement State requirements, as an authorized user for:

√ 35.390 ✓ 35.190 ✓ 35.290

√ 35.390 + generator experience

Name of Preceptor Signature 2 Telephone Number Date 08/29/2013 (248) 898-4126

License/Permit Number/Facility Name

21-01333-01 William Beaumont Hospital

Darlene Fink-Bennett, MD

NRC FORM 313A (RSO) (05-2012)	U.S. NUCLEAR REGULATORY COMMISSION	4.					
AND PRE	FFICER TRAINING AND EXPERIENCE CEPTOR ATTESTATION [10 CFR 35.50]		D BY OMB: NO. 3150-0120 (05/31/2015)				
Name of Proposed Radiation Safety Offic							
Justin Chang	M.D.						
	se authorizes the following medical uses (check all	that apply):					
	35.300 35.400 35.500 3	5.600 (remo	ote afterloader)				
35.600 (teletherapy)	35.600 (gamma stereotactic radiosurgery) 3	5.1000 ()				
PART I TRAINING AND EXPERIENCE (Select one of the four methods below) *Training and Experience, including board certification, must have been obtained within the 7 years preceding the date of application or the individual must have obtained related continuing education and experience since the required training							
	vide dates, duration, and description of continuing e						
1. Board Certification							
a. Provide a copy of the board ce	ertification.						
 b. Use Table 3.c. to describe train all types of medical use on the lice 	ning in radiation safety, regulatory issues, and eme ense.	rgency proc	edures for				
c. Skip to and complete Part II Pr	eceptor Attestation.						
	OR		İ				
2. Current Radiation Safety Office Officer for the Additional Media	er Seeking Authorization to Be Recognized as a lical Uses Checked Above	Radiation	Safety				
 a. Use the table in section 3.c. to procedures for the additional type 	describe training in radiation safety, regulatory issues of medical use for which recognition as RSO is s	ies, and eme ought.	ergency				
b. Skip to and complete Part II Pro	eceptor Attestation.						
	OR .						
3. Structured Educational Programme	am for Proposed Radiation Safety Officer						
a. Classroom and Laboratory Tra	aining						
Description of Training	Location of Training	Clock Hours	Dates of Training*				
Radiation physics and instrumentation							
Radiation protection							
Mathematics pertaining to the use and measurement of radioactivity							
Radiation biology							
Radiation dosimetry							

Total Hours of Training:

RADIATION SAFETY OFFICER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued)

3. Structured Educational Program for Proposed Radiation Safety Officer (continued)

Supervised Radiation Safety Experience
 (If more than one supervising individual is necessary to document supervised work experience, provide multiple copies of this section.)

Description of Experience	Location of Training/ License or Permit Number of Facility	Dates of Training*
Shipping, receiving, and performing related radiation surveys		
Using and performing checks for proper operation of instruments used to determine the activity of dosages, survey meters, and instruments used to measure radionuclides		
Securing and controlling byproduct material		
Using administrative controls to avoid mistakes in administration of byproduct material		
Using procedures to prevent or minimize radioactive contamination and using proper decontamination procedures		
Using emergency procedures to control byproduct material		
Disposing of byproduct material		
Licensed Material Used (e.g., 35.100, 35.200, etc.)+		1

Choose all applicable sections of 10 CFR Part 35 to describe radioisotopes and quantities used: 35.100, 35.200, 35.300, 35.400, 35.500, 35.600 remote afterloader units, 35.600 teletherapy units, 35.600 gamma stereotactic radiosurgery units, emerging technologies (provide list of devices).

RC FORM 313A (RSO) -2012)	U.S. NUCLEAR REGU	LATORY COMMISSION
RADIATION SAFETY OFFICER TRAINING A	AND EXPERIENCE AND PRECEPTOR ATTESTAT	ION (continued)
3. Structured Educational Program for Prop	osed Radiation Safety Officer (continued)	
b. Supervised Radiation Safety Experience	(continued)	
(If more than one supervising individual is copies of this section.)	s necessary to document supervised work experienc	e, provide multiple
Supervising Individual	License/Permit Number listing supervising in Radiation Safety Officer	ndividual as a
This license authorizes the following medical	uses;	
35.100 35.200 35.300	35.400	
35.500 35.600 (remote afterload	er) 35.600 (teletherapy)	
35.600 (gamma stereotactic radiosurgery	35.1000 ()	
c. Describe training in radiation safety, regulause on the license.	atory issues, and emergency procedures for all type	
Description of Training	Training Provided By	Dates of Training*
Radiation safety, regulatory issues, and emergency procedures for 35.100, 35.200, and 35.500 uses	Mary Lynn Wolfe, M.D.	to 8-30-13
		0-30-13
Radiation safety, regulatory issues, and emergency procedures for 35,300 uses	Maybeyn Welfe 4. D	+0
	0 0	8-30-13
Radiation safety, regulatory issues, and emergency procedures for 35.400 uses	maybegan Wolfe to D	10
	. 00	8-30-13
Radiation safety, regulatory issues, and emergency procedures for 35.600 - teletherapy uses		
Radiation safety, regulatory issues, and emergency procedures for 35.600 - remote afterloader uses		
Radiation safety, regulatory issues, and emergency procedures for 35.600 - gamma stereotactic radiosurgery uses		
Radiation safety, regulatory issues, and emergency procedures for 35.1000, specify use(s):		

NRC FORM 313A (RSO) (05-2012)	U.S. NUCLEAR REGULATORY COMMISSION
RADIATION SAFETY OFFICER TRAINING AND EXPERI	ENCE AND PRECEPTOR ATTESTATION (continued)
3. Structured Educational Program for Proposed Radiati	on Safety Officer (continued)
 Training in radiation safety, regulatory issues, and eme license (continued) 	ergency procedures for all types of medical use on the
Supervising Individual If training was provided by supervising RSO, AU, AMP, or ANP. (If more than one supervising individual is necessary to document supervised training, provide multiple copies of this page.) Mary Lynn Wolfe, MD	icense/Permit Number listing supervising individual
License/Permit lists supervising individual as:	Authorized Nuclear Pharmacist
Authorized as RSO, AU, ANP, or AMP for the following	medical uses:
☑ 35.100 ☑ 35.200 ☑ 35.300	35.400
35.500 35.600 (remote afterloader)	35.600 (teletherapy)
35.600 (gamma stereotactic radiosurgery)	35.1000 ()
d. Skip to and complete Part II Preceptor Attestation.	
OR	
 Authorized User, Authorized Medical Physicist, or A the licensee's license 	Authorized Nuclear Pharmacist identified on
a. Provide license number.	
b. Use the table in section 3.c. to describe training in raprocedures for all types of medical use on the license	
c. Skip to and complete Part II Preceptor Attestation.	
PART II - PRECEPTOR Note: This part must be completed by the individual's preceptor individual as long as the preceptor provides, directs, or wone preceptor is necessary to document experience, obsiriest Section	or. The preceptor does not have to be the supervising verifies training and experience required. If more than
Check one of the following:	
1. Board Certification	
Name of Proposed Radiation Safety Officer has	satisfactorily completed the requirements in
10 CFR 35.50(a)(1)(i) and (a)(1)(ii); or 35.50 (a)(2)(i) and	d (a)(2)(ii); or 35.50(c)(1).
OR	
2. Structured Educational Program for Proposed Radiati	on Safety Officers
I attest that has	satisfactorily completed a structural educational
Name of Proposed Radiation Safety Officer	
program consisting of both 200 hours of classroom and I radiation safety experience as required by 10 CFR 35.50	
OR	

NRC FORM 313A (RS (05-2012)	U.S. NUCLEAR REGULATORY COMMISSIO
	AFETY OFFICER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION (continued)
Preceptor Attest	ation (continued)
First Section (co Check one of the	
3. Addition	nal Authorization as Radiation Safety Officer
attest th	at Justin Chung is an Name of Proposed Radiation Safety Officer
Au	thorized User Authorized Nuclear Pharmacist
Au	thorized Medical Physicist
aspec	ied on the Licensees license and has experience with the radiation safety ts of similar type of use of byproduct material for which the individual has ion Safety Officer responsibilities
	AND
Second Section Complete for all (check all that apply):
1 attest that	Name of Proposed Radiation Safety Officer has training in the radiation safety, regulatory issues, and
	ocedures for the following types of use:
√ 35.100	
√ 35.200	
35.300	oral administration of less than or equal to 33 millicuries of sodium iodide I-131, for which a written directive is required
 ✓ 35.300	oral administration of greater than 33 millicuries of sodium iodide I-131
35.300	parenteral administration of any beta-emitter, or a photon-emitting radionuclide with a photon energy less than 150 keV for which a written directive is required
√ 35.300	parenteral administration of any other radionuclide for which a written directive is required
35.400	
35.500	
35.600	remote afterloader units
35.600	teletherapy units
35.600	gamma stereotactic radiosurgery units
35.1000	emerging technologies, including:

•				AND PRECEPT		(==::::::::::::::::
Third or or or			AND			
Third Section Complete for ALL						
I attest that	Justin Cha	I N Q tion Safety Officer		ed a level of radi	ation safety kno	wiedge
sufficient to functio				er for a medical u	se licensee.	
	ем жана жана жана жана жа			*********		
Fourth Section Complete the following	g for Preceptor A	Attestation a	nd signature			
I am the Radiation Saf	,	Hancock Regio				
. an mo radiadon dan				Name of Facility		
License/Permit Number:	13-16730-01					
						•
			,			
				Teleph	one Number	Date ,
me of Preceptor	1Sic	mature				
me of Preceptor	Welle 4.	nature	y Lypn U		8-4606	8/28/20

Organized through the cooperation of the American College of Radiology, the American Roentgen Ray Society, the American Radium Society, the Radiological Society of North America, the Section on Radiology of the American Medical Association,

Justin Philip Chang, MD

the American Society for Radiation Oncology, the Association of University Radiologists, and the American Association of Physicists in Medicine, Hereby certifies that

Has pursued an accepted course of graduate study and clinical work; has met certain standards and qualifications, including passing the examinations conducted under the authority of the American Board of Radiclogy, demonstrating to the satisfaction of the Board qualification to practice; and is therefore awarded the Board's certification in

Diagnostic Radiology

AM Eliqible

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COLUMBIA
1934

Certificate No. 61405

Ongoing validity of this certificate is contingent upon meeting the requirements of Maintenance of Certification.

This diplomate of the American Board of Radiology is permitted to use the DABS mark to signify this certification.

fine Shappy

Richard I Monic Secretary-Treasurer Hauf Schulds
Exercising Director

ATTE



Effective: July 01, 2012

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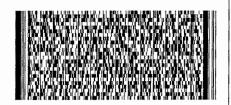
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