FORM DHHS/RHS-1M Supplement B-Therapy



STATE OF NEW HAMPSHIRE DEPARTMENT OF HEALTH AND HUMAN SERVICES RADIOLOGICAL HEALTH SECTION

AUTHORIZED USER TRAINING AND EXPERIENCE AND PRECEPTOR ATTESTATION

(For uses defined under New Hampshire Rules for the Control of Radiation He-P 4035.35)

Name of Proposed Authorized User:	State or Territory Where Licensed:					
Requested Authorization(s) – Check all that apply:						
4035.35 Use of Unsealed Byproduct Material for which a Written Directive is Require	red					
OR						
4035.35 Oral administration of sodium iodide I-131 requiring a written directive in quantities less than or equal to 1.22 gigabecquerels (33 millicuries)						
4035.35 Oral administration of sodium iodide I-131 requiring a written directive in quantities greater than 1.22 gigabecquerels (33 millicuries)						
4035.35 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						
4035.35 Parenteral administration of any other radionuclide for which a written direc	tive is required					
PART I – TRAINING AND EXPERIENC (He-P 4035.65 & 4035.66)	CE					
* Provide dates, duration, and description of training, continuing education, and experience related to the uses checked above and in accordance with He-P 4035.73.						
Board Certification						
a. Provide a copy of the board certification.						
b. For 4035.65, provide documentation on supervised clinical case experience. document this experience.	The table in section 3.c. may be used to					
c. For 4035.66, provide documentation on classroom and laboratory training, so clinical case experience. The tables in sections 3.a., 3.b., and 3.c. may be use						
d. Skip to and complete Part II Preceptor Attestation.						
OR						
2. Current 4035.35, 4035.41, or 4035.47 Authorized User Seeking Additional A	<u>Authorization</u>					
a. Authorized user on Materials License under the requ Nuclear Regulatory Commission or Agreement State requirements (check all	nirements below or equivalent U.S. l that apply):					
☐ 4035.59 ☐ 4035.65 ☐ 4035.66 (<33 mCi I-131) ☐ 4035.66	5 (>33 mCi I-131) 4035.69					
b. If currently authorized for a subset of clinical uses under 4035.35, provide do supervised case experience. The table in section 3.c. may be used to docume completed Part II Preceptor Attestation.						
c. If currently authorized under 4035.59 or 4035.69 and requesting authorization classroom and laboratory training, supervised work experience, and supervise sections 3.a., 3.b., and 3.c. may be used to document this experience. Also pattestation.	ed clinical case experience. The tables in					

OR

3.	Training and Experie	nce for Proposed Autl	horized User				
a. Classroom and Laboratory Training							
	4035.65	☐ 4035.66 (<33 mC	i I-131)] 4035.66 (>33 mCi	I-131)	4035	.66 (Parenteral)
	Description of	Training	Location of Training		Clock Hours		Dates of Training*
	Radiation physics and	instrumentation					
	Radiation protection						
	Mathematics pertaining measurement of radios						
	Chemistry of byprodu material for medical u						
	Radiation biology						
			Total I	Hours of Training	·		
	b. Supervised Work E 4035.65 (If more than one sup Total Hours of Ex	4035.66 (<33 mC		4035.66 (>33 mCi supervised work ex			.66 (Parenteral) ple copies of this section.)
	_	of Experience Include		tion of Experience/ Permit Number of Fa	acility	Confirm	Dates of Experience*
	Ordering, receiving, and unpacking radioactive materials safely and performing the related radiation surveys					☐ Yes ☐ No	
Performing quality control procedures on instruments used to determine the activity of dosages and performing checks for proper operation of survey meters Calculating, measuring, and safely preparing patient or human research subject dosages Using administrative controls to prevent a medical event involving the use of unsealed byproduct material					☐ Yes ☐ No		
					☐ Yes ☐ No		
					☐ Yes ☐ No		
	Using procedures to contain spilled byproduct material safely and using proper decontamination procedures					☐ Yes ☐ No	
	Supervising Individual License/Permit number listing supervising individual as a authorized user					ising individual as an	
	Supervising individual requirements (check al	meets the requirements (I that apply)**:	below, or equiva	alent U.S. Nuclear Re	gulatory (Commission	or Agreement State
	☐ 4035.65 With experience, that includes at least 3 cases of, administering dosages of:						
	4035.66(a)	Oral administration of sodium iodide I-131 requiring a written directive in quantities less than or					
	4035.66(d)	equal to 1.22 gigabecquerels (33 millicuries) Oral administration of sodium iodide I-131 requiring a written directive in quantities greater than					
	4035.66(i)	or a definition of southin found 1-131 requiring a written directive in quantities greater than					
Parenteral administration of any beta-emitter or photon-emitting radionuclide with a photo less than 150 keV, and/or parenteral administration of any other radionuclide, for which a directive is required							
		ized user must have exp		istering dosages in th	e same do	sage categor	y or categories as the

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

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

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) Oral administration of sodium iodide I-131 requiring a written directive in quantities greater than 1.22 gigabecquerels (33 millicuries) 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) Supervising Individual		License/Permit number listing supervisin	g individual as an
Supervising individual meets the requirements requirements (check all that apply)**:	ents below, or equiva	lent U.S. Nuclear Regulatory Commission or	Agreement State
□ 4035.65 □ 4035.66(a) □ 4035.66(d) □ 4035.66(d) □ 4035.66(i) □ 4035.66(i) □ Parenteral administration of any beta-emitter or photon-emitting radionuclide, for which a written directive is required			
** Supervising authorized user must have individual requesting authorized user statements.		istering dosages in the same dosage category of	or categories as the

d. Provide completed Part II Preceptor Attestation.

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 4035.68.)

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."

<u>FIRST SECTION</u> – Check one of the following for each requested authorization:				
For 4035.65: 1. Board Certification				
I attest that	_ has satisfactorily completed the			
requirements in 4035.65(b)(1).				
OR				
2. <u>Training and Experience</u>				
I attest that	has satisfactorily completed the training			
and experience as required by 4035.65(c).				
For 4035.66 (Identical Attestation Statement Regardless of Training and E	<u>experience Pathway)</u> :			
I attest that has satisfact Name of Proposed Authorized User	orily completed the training as required			
in He-P 4035.66(b)(1), and the experience required in 4035.66(b)(2).				
AND				
<u>SECOND SECTION</u> – Complete for all submittals.				
I attest that has satisfactor	rily completed the required clinical case experience			
Name of Proposed Authorized User				
required in 4035.65(c)(2)b. listed below:				
Oral administration of less than or equal to 33 millicuries of sodium iodide I-131 for which a written directive is required				
☐ Oral administration of greater than 33 millicuries of sodium iodide I-131				
Parenteral administration of any beta-emitter, or a photon-emitting rad and/or parenteral administration of any other radionuclide, for which				
AND				
THIRD SECTION – Complete for all submittals.				
I attest that has satisfactor Name of Proposed Authorized User	rily achieved a level of competency to function			
independently as an authorized user for: Oral administration of less than or equal to 33 millicuries of sodium in	odida I 131 for which a written directive is required			
Oral administration of fess than or equal to 33 minicules of sodium to 37 minicules of sodium to	•			
Parenteral administration of any beta-emitter, or a photon-emitting rad				
and/or parenteral administration of any other radionuclide, for which				

FOURTH SECTION

Complete for 4035.66 (Current 4035.59 or 4035.69 Authorized User):						
I attest that	attest that is an authorized user under 4035.59 or 4035.69 or equivalent Name of Proposed Authorized User					
required by 4035.66(g)(4), and	U.S. Nuclear Regulatory Commission or Agreement State requirements, has satisfactorily completed the training, as required by 4035.66(g)(4), and the experience required by 4035.66(g)(5), and has achieved a level of competency sufficient to function independently as an authorized user for:					
	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					
Parenteral administration of	f any other radionuclide for which a writ	ten directive is required				
	AND					
Board Certification:						
I attest that	I attest that has satisfactorily completed the board certification Name of Proposed Authorized User					
requirements of $4035.66(g)(1)$, has satisfactorily completed the training required by $4035.66(g)(4)$ and the experience required by $4035.66(g)(5)$, and has achieved a level of competency sufficient to function independently as an authorized user for:						
	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					
Parenteral administration of	f any other radionuclide for which a writ	ten directive is required				
FIFTH SECTION – Complete for a	ll submittals.					
	equivalent U.S. Nuclear Regulatory Cor	nmission or Agreement Sta	ate requirements, as an			
☐ 4035.65 ☐ 4035		4035.66(i)				
I have experience administering dosages in the following categories for which the proposed authorized user is requesting authorization:						
	☐ Oral administration of less than or equal to 33 millicuries of sodium iodide I-131 for which a written directive is required ☐ Oral administration of greater than 33 millicuries of sodium iodide I-131					
Parenteral administration of any beta-emitter, or a photon-emitting radionuclide with a photon energy less than 150 keV, and/or parenteral administration of any other radionuclide, for which a written directive is required						
Name of Preceptor:	Signature:	Telephone Number:	Date:			
License/Permit Number/Facility Name:						