**Medical Use of Byproduct Material—Authorized User Clarification, Part 35**

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| **Change to NRC**  **Section** | **Title** | **State**  **Section** | **Compatibility**  **Category** | **Summary of Change to CFR** | **Difference**  **Yes/No** | **Significant**  **Yes/No** | **If Difference, Why or Why Not Was a Comment Generated** |
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| § 35.50 | Training for Radiation Safety Officer |  | B | **In § 35.50, paragraph (a)(2)(ii)(B) is**  **revised to read as follows:**  \* \* \* \* \*  (a) \* \* \*  (2) \* \* \*  (ii) \* \* \*  (B) In clinical nuclear medicine  facilities providing diagnostic and/or therapeutic services under the direction of physicians who meet the requirements for authorized users in §§ 35.57, 35.290, or 35.390; |  |  |  |
| § 35.51 | Training for an authorized medical  physicist. |  | B | **In § 35.51, paragraphs (a)(2)(ii) and (b)(2) are revised to read as follows:**  (a) \* \* \*  (2) \* \* \*  (ii) In clinical radiation facilities  providing high-energy, external beam therapy (photons and electrons with energies greater than or equal to 1 million electron volts) and brachytherapy services under the direction of physicians who meet the requirements in § 35.57, 35.490, or 35.690; and \* \* \* \* \*  (b) \* \* \*  (2) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraphs (c) and (a)(1) and (a)(2), or (b)(1) and (c) of this section, and has achieved a level of competency sufficient to function independently as an authorized medical physicist for each type of therapeutic medical unit for which the individual is requesting authorized medical physicist status. The written attestation must be signed by a preceptor authorized medical physicist who meets the requirements in §§ 35.51, 35.57, or equivalent Agreement State requirements for an authorized medical physicist for each type of therapeutic medical unit for which the individual is requesting authorized medical physicist status; and  \* \* \* \* \* |  |  |  |
| § 35.57 | Training for experienced Radiation Safety Officer, teletherapy or medical  physicist, authorized medical physicist,  authorized user, nuclear pharmacist, and  authorized nuclear pharmacist**.** |  | B | **In § 35.57, a new paragraph (c) is added to read as follows:**  (c) Individuals who need not comply with training requirements as described in this section may serve as preceptors for, and supervisors of, applicants seeking authorization on NRC licenses for the same uses for which these individuals are authorized. |  |  |  |
| § 35.190 | Training for uptake, dilution, and excretion studies. |  | B | **In § 35.190, the introductory text of paragraph (c)(1)(ii) and paragraph (c)(2) are revised to read as follows:**  (c)(1) \* \* \*  (ii) Work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57, 35.190, 35.290, 35.390, or equivalent Agreement State requirements, involving—  \* \* \* \* \*  (2) Has obtained written attestation, signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.190, 35.290, or 35.390, or equivalent Agreement State requirements, that the individual has satisfactorily completed the requirements in paragraph (a)(1) or (c)(1) of this section and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under  § 35.100. |  |  |  |
| § 35.290 | Training for imaging and  localization studies. |  | B | **In § 35.290, the introductory text of paragraph (c)(1)(ii) and paragraph (c)(2)**  **are revised to read as follows:**  (c)(1) \* \* \*  (ii) Work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57, 35.290, or 35.390 and 35.290(c)(1)(ii)(G), or equivalent Agreement State requirements, involving—  \* \* \* \* \*  (2) Has obtained written attestation, signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.290, or 35.390 and 35.290(c)(1)(ii)(G), or equivalent Agreement State requirements, that the individual has satisfactorily completed the requirements in paragraph (a)(1) or (c)(1) of this section and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under §§ 35.100 and 35.200. |  |  |  |
| § 35.390 | Training for use of unsealed  byproduct material for which a written  directive is required. |  | B | **In § 35.390, the introductory text of paragraph (b)(1)(ii) and paragraph (b)(2) are revised to read as follows:**  (b)(1) \* \* \*  (ii) Work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57, 35.390, or equivalent Agreement State requirements. A supervising authorized user, who meets the requirements in  § 35.390(b), must also have experience in administering dosages in the same dosage category or categories (i.e.,  § 35.390(b)(1)(ii)(G)) as the individual requesting authorized user status. The work experience must involve—  \* \* \* \* \*  (2) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraphs (a)(1) and (b)(1)(ii)(G) or (b)(1) of this section, and has achieved a level of competency sufficient to function independently as an authorized user for the medical uses authorized under § 35.300. The written attestation must be signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.390, or equivalent Agreement State requirements. The preceptor authorized user, who meets the requirements in  § 35.390(b) must have experience in administering dosages in the same dosage category or categories (i.e.,  § 35.390(b)(1)(ii)(G)) as the individual requesting authorized user status. |  |  |  |
| § 35.392 | Training for the oral  administration of sodium iodide I-131  requiring a written directive in quantities  less than or equal to 1.22 gigabecquerels  (33 millicuries). |  | B | **In § 35.392, the introductory text of paragraph (c)(2) and paragraph (c)(3) are revised to read as follows:**  (c) \* \* \*  (2) Has work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57,  35.390, 35.392, 35.394, or equivalent Agreement State requirements. A supervising authorized user who meets the requirements in § 35.390(b) must also have experience in administering dosages as specified in  §§ 35.390(b)(1)(ii)(G)(1) or  35.390(b)(1)(ii)(G)(2). The work experience must involve—  \* \* \* \* \*  (3) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraphs (c)(1) and (c)(2) of this section, and has achieved a level of competency sufficient to function independently as an authorized user for medical uses authorized under  § 35.300. The written attestation must be signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.390, 35.392, 35.394, or equivalent Agreement State requirements. A preceptor authorized user, who meets the requirement in § 35.390(b), must also have experience in administering dosages as specified in  §§ 35.390(b)(1)(ii)(G)(1) or  35.390(b)(1)(ii)(G)(2). |  |  |  |
| § 35.394 | Training for the oral  administration of sodium iodide I-131  requiring a written directive in quantities  greater than 1.22 gigabecquerels (33  millicuries). |  | B | **In § 35.394, the introductory text of paragraph (c)(2) and paragraph (c)(3) are**  **revised to read as follows:**  (c) \* \* \*  (2) Has work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57,  35.390, 35.394, or equivalent Agreement State requirements. A supervising authorized user, who meets the requirements in  § 35.390(b), must also have experience in administering dosages as specified in  § 35.390(b)(1)(ii)(G)(2). The work experience must involve—  \* \* \* \* \*  (3) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraphs (c)(1) and (c)(2) of this section, and has achieved a level of competency sufficient to function independently as an authorized user for medical uses authorized under  § 35.300. The written attestation must be signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.390, 35.394, or equivalent Agreement State requirements. A preceptor authorized user, who meets the requirements in § 35.390(b), must also have experience in administering osages as specified in  § 35.390(b)(1)(ii)(G)(2). |  |  |  |
| § 35.396 | Training for the parenteral  administration of unsealed byproduct  material requiring a written directive. |  | B | **In § 35.396, the introductory text of paragraph (d)(2) and paragraph (d)(3) are revised to read as follows:**  (d) \* \* \*  (2) Has work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57,  35.390, 35.396, or equivalent Agreement State requirements, in the parenteral administration, for which a written directive is required, of any beta emitter, or any 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. A supervising authorized user who meets the requirements in § 35.390 must have experience in administering dosages as specified in §§ 35.390(b)(1)(ii)(G)(3) and/or 35.390(b)(1)(ii)(G)(4). The work experience must involve—  \* \* \* \* \*  (3) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraph (b) or (c) of this section, and has achieved a level of competency sufficient to function independently as an authorized user for the parenteral administration of unsealed byproduct material requiring a written directive. The written attestation must be signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.390, 35.396, or equivalent Agreement State requirements. A preceptor authorized user, who meets the requirements in § 35.390, must have experience in administering dosages as specified in  §§ 35.390(b)(1)(ii)(G)(3)  and/or 35.390(b)(1)(ii)(G)(4). |  |  |  |
| § 35.490 | Training for use of manual  brachytherapy sources. |  | B | **In § 35.490, the introductory text of paragraph (b)(1)(ii) and paragraphs (b)(2) and (b)(3) are revised to read as follows**:  (b)(1) \* \* \*  (ii) 500 hours of work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57, 35.490, or equivalent Agreement State requirements at a medical institution, involving—  \* \* \* \* \*  (2) Has completed 3 years of supervised clinical experience in radiation oncology, under an authorized user who meets the requirements in §§ 35.57, 35.490, or equivalent Agreement State requirements, as part of a formal training program approved by the Residency Review Committee for Radiation Oncology of the Accreditation Council for Graduate Medical Education  or the Royal College of Physicians and Surgeons of Canada or the Committee on Postdoctoral Training of the American Osteopathic Association. This experience may be obtained concurrently with the supervised work experience required by paragraph (b)(1)(ii) of this section; and  (3) Has obtained written attestation, signed by a preceptor authorized user who meets the requirements in  §§ 35.57, 35.490, or equivalent Agreement State requirements, that the individual has satisfactorily completed the requirements in paragraph (a)(1), or paragraphs (b)(1) and (b)(2), of this section and has achieved a level of competency sufficient to function independently as an authorized user of manual brachytherapy sources for the medical uses authorized under § 35.400. |  |  |  |
| § 35.491 | Training for ophthalmic use of  strontium-90. |  | B | **In § 35.491, paragraph (b)(3) is revised to read as follows:**  (b) \* \* \*  (3) Has obtained written attestation, signed by a preceptor authorized user who meets the requirements in  §§ 35.57, 35.490, 35.491, or equivalent Agreement State requirements, that the individual has satisfactorily completed the requirements in paragraph (b) of this section and has achieved a level of competency sufficient to function independently as an authorized user of strontium-90 for ophthalmic use. |  |  |  |
| § 35.690 | Training for use of remote  afterloader units, teletherapy units, and  gamma stereotactic radiosurgery units. |  | B | **In § 35.690, the introductory text of paragraph (b)(1)(ii) and paragraphs (b)(2) and (b)(3) are revised to read as follows:**  (b)(1) \* \* \*  (ii) 500 hours of work experience, under the supervision of an authorized user who meets the requirements in §§ 35.57, 35.690, or equivalent Agreement State requirements at a medical institution, involving—  \* \* \* \* \*  (2) Has completed 3 years of supervised clinical experience in radiation therapy, under an authorized user who meets the requirements in §§ 35.57, 35.690, or equivalent Agreement State requirements, as part of a formal training program approved by the Residency Review Committee for Radiation Oncology of the Accreditation Council for Graduate Medical Education or the Royal College of Physicians and Surgeons of Canada or the Committee on Postdoctoral Training of the American Osteopathic Association. This experience may be obtained concurrently with the supervised work experience required by paragraph (b)(1)(ii) of this section; and  (3) Has obtained written attestation that the individual has satisfactorily completed the requirements in paragraph (a)(1) or paragraphs (b)(1) and  (b)(2), and paragraph (c), of this section, and has achieved a level of competency sufficient to function independently as an authorized user of each type of therapeutic medical unit for which the individual is requesting authorized user status. The written attestation must be signed by a preceptor authorized user who meets the requirements in §§ 35.57, 35.690, or equivalent Agreement State requirements for an authorized user for each type of therapeutic medical unit for which the individual is requesting authorized user status; and |  |  |  |