**Preparation, Transfer for Commercial Distribution, and Use of Byproduct Material for Medical Use**

 **(59 FR 61767; 59 FR 65243; 60 FR 322) RATS ID 1995‑1 Effective 1/1/95**

| **Change to** **NRC****Section** | **Title** |  **State** **Section** | **Compatibility****Category** | **Summary of Change** | **Difference****Yes/No** | **Significant****Yes/No** | **If Difference, Why or Why Not Was a Comment Generated** |
| --- | --- | --- | --- | --- | --- | --- | --- |
| 30.4 | Definitions |  | C | **Added Definition:**Medical Use |  |  |  |
| 32.72 |  |  | B |  |  |  |  |
| 35.32 | Definitions |  | D | **Added Definition:**Authorized Nuclear Pharmacist | N/A |  |  |
| 35.32 | Definitions |  | C | **Added Definition:**Authorized User |  |  |  |
| 35.32 | Definitions |  | C | **Added Definition:**Medical Use |  |  |  |
| 35.32 | Definitions |  | C | **Added Definition:**Misadministration |  |  |  |
| 35.32 | Definitions |  | D | **Added Definition:**Pharmacist | N/A |  |  |
| 35.32 | Definitions |  | D | **Added Definition:**Recordable Event |  |  |  |
| 35.32 | Definitions |  | C | **Added Definition:**Written Directive | N/A |  |  |
| 35.11 | License Required |  | C |  |  |  |  |
| 35.32(a)(b)(c) | Quality management program |  | H&S |  |  |  |  |
| 35.32(d)(e)(f) | Quality management program |  | D |  | N/A |  |  |
| 35.33 | Notifications |  | C |  |  |  |  |
| 35.52 | Possession, use, calibration, and check of instruments to measure dosages of alpha- or beta-emitting radionuclides |  | H&S |  |  |  |  |
| 35.53(a)(b) | Measurements of dosages of unsealed byproduct material for medical use |  | H&S |  |  |  |  |
| 35.53(c) | Measurements of dosages of unsealed byproduct material for medical use |  | D | N/A | N/A |  |  |
| 35.75(a) | Release of individuals containing radio pharmaceuticals or permanent implants |  | C |  |  |  |  |
| 35.75(b) | Release of individuals containing radio pharmaceuticals or permanent implants |  | H&S |  |  |  |  |
| 35.75(c)(d) | Release of individuals containing radio pharmaceuticals or permanent implants |  | D | N/A | N/A |  |  |
| 35.100 | Use of unsealed byproduct material for uptake, dilution, and excretion studies |  | H&S |  |  |  |  |
| 35.200 | Use of unsealed byproduct material for imaging and localization studies |  | H&S |  |  |  |  |
| 35.300 | Use of unsealed byproduct material for therapeutic administration |  | H&S |  |  |  |  |
| 35.404 | Release of patients or human research subjects treated with temporary implants |  | C |  |  |  |  |
| 35.406(a)(c) | Brachytherapy sources inventory |  |  H&S |  |  |  |  |
| 35.406 | Brachytherapy sources inventory |  | D | N/A | N/A |  |  |
| 35.610 | Safety instructions |  | H&S |  |  |  |  |
| 35.615 | Safety precautions |  | D/H&S |  |  |  |  |