Ohio Administrative Code

Rule 3701:1-58-37 Use of unsealed radioactive material for which a written directive is required.

Effective: August 15, 2021

A licensee may use any unsealed radioactivematerial identified in paragraph (B)(1)(b)(vi) of rule 3701:1-58-40 theAdministrative Code prepared for medical use and for which a written directiveis required that is:

(A) Obtained from:

(1) A manufacturer or preparer licensed under rule 3701:1-46-43 of the Administrative Code or equivalent United States nuclear regulatory commission or agreement state requirements; or

(2) A PET radioactive drug producer licensed in accordance with paragraph (I) of rule 3701:1-40-14 of the Administrative Code or equivalent United States nuclear regulatory commission or agreement state requirement; or

(B) Prepared by, excluding production of PET radionuclides:

(1) An authorized nuclear pharmacist;

(2) A physician who is an authorized user and who meets the requirements specified in rule 3701:1-58-36 or 3701:1-58-40 of the Administrative Code; or

(3) An individual under the supervision, as specified in rule 3701:1-58-14 of the Administrative Code, of the authorized nuclear pharmacist in paragraph (B)(1) of this rule or the physician who is an authorized user in paragraph (B)(2) of this rule; or

(C) Obtained from and prepared by an United States nuclear regulatory commission or agreement state licensee for use in research in accordance with an investigational new drug protocol accepted by United States food and drug administration; or

(D) Prepared by the licensee for use in research in accordance with an investigational new drug protocol accepted by United States food and drug administration.