

3701:1-58-07

Application for license, amendment or renewal.

- (A) An application must be signed by the applicant's or licensee's management as defined in rule 3701:1-38-01 of the Administrative Code.
- (B) An application for a license for medical use of radioactive material as described in rules 3701:1-58-32, 3701:1-58-34, 3701:1-58-37, 3701:1-58-43, 3701:1-58-53, 3701:1-58-55, and 3701:1-58-72 of the Administrative Code must be made by:
 - (1) Submitting documentation in accordance with rule 3701:1-40-14 of the Administrative Code, and including the facility diagram, equipment, and training and experience qualifications of the radiation safety officer, authorized user(s), authorized medical physicist(s), and authorized nuclear pharmacist(s);
 - (2) Submitting the appropriate license fees listed in rule 3701:1-38-02 of the Administrative Code after receiving an invoice from the department; and
 - (3) Submitting procedures required by rules 3701:1-58-58 and 3701:1-58-64 to 3701:1-58-66 of the Administrative Code, as applicable.
- (C) A request for a license amendment or renewal must be made by:
 - (1) Submitting documentation in accordance with rule 3701:1-40-14 of the Administrative Code;
 - (2) Submitting the appropriate license or amendment fees listed in rule 3701:1-38-02 of the Administrative Code after receiving an invoice from the department; and
 - (3) Submitting procedures required by rules 3701:1-58-58 and 3701:1-58-64 to 3701:1-58-66 of the Administrative Code, as applicable.
- (D) In addition to the requirements in paragraphs (B) and (C) of this rule, an application for a license or amendment for medical use of radioactive material as described in rule 3701:1-58-72 of the Administrative Code must also include information regarding any radiation safety aspects of the medical use of the material that is not addressed in rules 3701:1-58-01 to 3701:1-58-31 of the Administrative Code.
 - (1) The applicant shall also provide specific information on:
 - (a) Radiation safety precautions and instructions;
 - (b) Methodology for measurement of dosages or doses to be administered to patients or human research subjects; and
 - (c) Calibration, maintenance, and repair of instruments and equipment necessary for radiation safety.

- (2) The applicant or licensee shall also provide any other information requested by the director in review of the application.

- (E) An applicant that satisfies the requirements specified in rule 3701:1-40-23 of the Administrative Code may apply for a type A specific license of broad scope.

Five Year Review (FYR) Dates: 06/12/2015 and 06/01/2020

CERTIFIED ELECTRONICALLY

Certification

06/12/2015

Date

Promulgated Under: 119.03
Statutory Authority: 3748.02, 3748.04
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