PDR

DESIGNATED ORIGINAL

Standard Form 83
(Rev. September 1983)

Request for OMB Review

Paulette Smith

Important

Read instructions before completing form. Do not use the same SF 83 to request both an Executive Order 12291 review and approval under the Paperwork Reduction Act.

Answer all questions in Part I. If this request is for review under E.O. 12291, complete Part II and sign the regulatory certification. If this request is for approval under the Paperwork Reduction Act and 5 CFR 1320, skip Part II, complete Part III and sign the paperwork certification.

Send three copies of this form, the material to be reviewed, and for paperwork—three copies of the supporting statement, to:

Office of Information and Regulatory Affairs
Office of Management and Budget
Attention: Docket Library, Room 3201
Washington, DC 20503

PART I.—Complete This Part for All Requests.

1. Department/agency and Bureau/office originating request

U. S. Nuclear Regulatory Commission

2. Agency code

3 1 5 0

3. Name of person who can best answer questions regarding this request

Michael Lamastra

Telephone number

(301) 504-3416

4. Title of information collection or rulemaking

10 CFR Part 33 - Specific Domestic Licenses of Broad Scope for Byproduct Material

5. Legal authority for information collection or rule *(cite United States Code, Public Law, or Executive Order)*

42 USC 2201(o) or

6. Affected public *(check all that apply)*

1 ☐ Individuals or households
2 ☒ State or local governments
3 ☐ Farms
4 ☒ Businesses or other for-profit
5 ☒ Federal agencies or employees
6 ☒ Non-profit institutions
7 ☒ Small businesses or organizations

PART II.—Complete This Part Only If the Request is for OMB Review Under Executive Order 12291

7. Regulation Identifier Number (RIN)

___ ___ ___ ___ ___ ___ ___ ___ ___ ___ or ☐ None assigned

8. Type of submission *(check one in each category)*

Classification
1 ☐ Major
2 ☐ Nonmajor

Stage of development
1 ☐ Proposed or draft
2 ☐ Final or interim final, with prior proposal
3 ☐ Final or interim final, without prior proposal

Type of review requested
1 ☐ Standard
2 ☐ Pending
3 ☐ Emergency
4 ☐ Statutory or judicial deadline

9. CFR section affected

_____ CFR _____

10. Does this regulation contain reporting or recordkeeping requirements that require OMB approval under the Paperwork Reduction Act and 5 CFR 1320? ☐ Yes ☐ No

11. If a major rule, is there a regulatory impact analysis attached? 1 ☐ Yes 2 ☐ No

If "No," did OMB waive the analysis? 3 ☐ Yes 4 ☐ No

Certification for Regulatory Submissions

In submitting this request for OMB review, the authorized regulatory contact and the program official certify that the requirements of E.O. 12291 and any applicable policy directives have been complied with.

Signature of program official — Date

Signature of authorized regulatory contact 050001 — Date

12. *(OMB use only)*

D502 1

ART III.—Complete This Part Only if the Request is for Approval of a Collection of Information Under the Paperwork Reduction Act and 5 CFR 1320.

2. Abstract—Describe needs, uses and affected public in 50 words or less

"Radioactive Materials, Radiation Safety"
10 CFR Part 33 specifies requirements for applying for and being granted licenses authorizing broad scope use of radioactive byproduct material.

4. Type of information collection *(check only one)*

Information collections not contained in rules

1 ☐ Regular submission
2 ☐ Emergency submission *(certification attached)*

Information collections contained in rules

3 ☒ Existing regulation *(no change proposed)*
4 ☐ Notice of proposed rulemaking (NPRM)
5 ☐ Final, NPRM was previously published
6 Final or interim final without prior NPRM
A ☐ Regular submission
B ☐ Emergency submission *(certification attached)*

7. Enter date of expected or actual Federal Register publication at this stage of rulemaking *(month, day, year)*: _____

8. Type of review requested *(check only one)*

1 ☐ New collection
2 ☐ Revision of a currently approved collection
3 ☒ Extension of the expiration date of a currently approved collection without any change in the substance or in the method of collection
4 ☐ Reinstatement of a previously approved collection for which approval has expired
5 ☐ Existing collection in use without an OMB control number

9. Agency report form number(s) *(include standard/optional form number(s))*

Not applicable

10. Annual reporting or disclosure burden

1 Number of respondents	1
2 Number of responses per respondent	1
3 Total annual responses *(line 1 times line 2)*	1
4 Hours per response	1
5 Total hours *(line 3 times line 4)*	1

11. Annual recordkeeping burden

1 Number of recordkeepers	
2 Annual hours per recordkeeper	
3 Total recordkeeping hours *(line 1 times line 2)*	
4 Recordkeeping retention period	years

12. Total annual burden

1 Requested *(line 17-5 plus line 18-3)*	1
2 In current OMB inventory	1
3 Difference *(line 1 less line 2)*	0
Explanation of difference	
4 Program change	
5 Adjustment	

13. Current *(most recent)* OMB control number or comment number

3150-0015

14. Requested expiration date

3 years from approval date

22. Purpose of information collection *(check as many as apply)*

1 ☐ Application for benefits
2 ☐ Program evaluation
3 ☐ General purpose statistics
4 ☒ Regulatory or compliance
5 ☐ Program planning or management
6 ☐ Research
7 ☐ Audit

23. Frequency of recordkeeping or reporting *(check all that apply)*

1 ☐ Recordkeeping

Reporting

2 ☐ On occasion
3 ☐ Weekly
4 ☐ Monthly
5 ☐ Quarterly
6 ☐ Semi-annually
7 ☐ Annually
8 ☐ Biennially
9 ☒ Other *(describe)*: Initial application and five year renewal

24. Respondents' obligation to comply *(check the strongest obligation that applies)*

1 ☐ Voluntary
2 ☐ Required to obtain or retain a benefit
3 ☒ Mandatory

15. Are the respondents primarily educational agencies or institutions or is the primary purpose of the collection related to Federal education programs? ☐ Yes ☒ No

16. Does the agency use sampling to select respondents or does the agency recommend or prescribe the use of sampling or statistical analysis by respondents? ☐ Yes ☒ No

17. Regulatory authority for the information collection

10 CFR Part 33 ; or _____ FR _____ ; or, Other *(specify)*: _____

Paperwork Certification

By submitting this request for OMB approval, the agency head, the senior official or an authorized representative, certifies that the requirements of 5 CFR 1320, the Privacy Act, statistical standards or directives, and any other applicable information policy directives have been complied with.

Signature of program official	Date

Signature of agency head, the senior official or an authorized representative	Date
Gerald F. Cranford, Designated Senior Official for Information Resources Management	12-30-92

☆ GPO : 1984 O - 453-776

Supporting Statement
for
10 CFR Part 33
Specific Domestic Licenses of
Broad Scope for Byproduct Material
(3150-0015)

Description of the Information Collection

10 CFR Part 33 specifies requirements for applying for and being granted licenses authorizing broad scope use of byproduct material. Three types of licenses may be issued: Type A, Type B, or Type C, with Type A being the largest program. The types of licenses are defined by Section 33.11. Applicants and licensees are primarily medical institutions, colleges, universities, government agencies, and large private companies engaged in broad educational, research, and development activities.

A. Justification

Need for the Collection of Information

Section 33.12 specifies that an applicant for a broad license must complete NRC Form 313, "Application for Materials License." Form 313 has previously been cleared under OMB Clearance No. 3150-0120, which should be referred to for further supporting information and burden and cost data.

Sections 33.13, 33.14, and 33.15 specify the information which must be included in an application in order for NRC to issue a license based on a finding that the public health and safety will be adequately protected.

Section 33.13 specifies the requirements for a Type A broad license. It requires that an applicant have equipment and facilities adequate to protect health and minimize danger to life or property, have personnel with adequate training and experience, and have adequate administrative controls and procedures, including a radiation safety committee and radiation safety officer. The applicant must describe his program for meeting these requirements when he submits a license application, in order to establish that he can safely use radioactive material.

Section 33.14 specifies the requirements for a Type B broad license. The requirements are similar to those specified in Section 33.13, except that the applicant need not demonstrate as extensive experience in use of radioactive materials as is required for a Type A broad license, and need not have a radiation safety committee. The application submitted by the applicant must demonstrate that he can meet these requirements.

Section 33.15 specifies the requirements for a Type C broad license. It requires that an applicant have equipment and facilities adequate to protect health and minimize danger to life or property, have personnel meeting certain training and experience requirements, supervise use of radioactive material, and have adequate administrative controls and procedures to assure safe use of radioactive material. The application submitted by the applicant must demonstrate that he can meet these requirements.

Agency Use of Information

NRC reviews the information submitted in order to determine whether the applicant's training, experience, equipment, facilities, and procedures for the use of byproduct material are adequate to protect the public health and safety as required by the Atomic Energy Act, so that the Commission may determine whether to issue, amend, or renew a broad license.

Reduction of Burden Through Information Technology

There are no legal obstacles to reducing the burden associated with this information collection. Applicants and licensees may use electronic information processing systems to prepare and submit required information.

Effort to Identify Duplication

The Information Requirements Control Automated System (IRCAS) was searched to determine duplication. None was found.

Effort to Use Similar Information

There is no similar information available to the NRC.

Effort to Reduce Small Business Burden

The majority of licensees who use byproduct material are small businesses. Since the health and safety consequences of improper handling or use of radioactive byproduct material are the same for large and small entities, it is not possible to reduce the burden on small businesses by less frequent or less complete accounting or control procedures.

Consequences of Less Frequent Collection

Applications for new licenses and amendments are submitted only once. Applications for renewal of licenses are submitted every five years. This is the minimum frequency necessary to assure that licensees will continue to conduct programs in a manner that will assure adequate protection of the public health and safety.

Circumstances Which Justify Variation from OMB Guidelines

There are no variations from OMB Guidelines.

Consultations Outside the Agency

There have been no outside consultations since the previous clearance.

Confidentiality of Information

None, except for proprietary information.

Sensitive Questions

None.

Estimate of Compliance Burden

Reporting Requirements (§§33.12, 33.13, 33.14, 33.15 Combined)

Annual Responses	Staff Hours per Submittal	Annual Licensee Burden
[120]	[20]	[2,400]

This burden data is provided for information only. The burden for this information collection is attributable to and is reported for inventory purposes under NRC Form 313, OMB Clearance No. 3150-0120.

Estimated annual cost to the industry to respond to the collection requirements is $295,200 (2,400 hrs @ $123/hr).

Source of burden data and Method of Estimating Burden

Burden and cost estimates are based on the number of applications received by NRC annually in previous years and on staff experience and discussions with a small number of licensees.

Estimated Annual Cost to the Federal Government

Professional: 120 responses x 5 hrs/response x $123/hr = $73,800 annually

Clerical and other: See OMB Clearance No. 3150-0120

Reasons for Change in Burden

There is no change in burden.

Publication for Statistical Use

None.

B. Collection of Information Employing Statistical Methods

Statistical methods are not used in this collection of information.

[7590-01]

NUCLEAR REGULATORY COMMISSION

Documents Containing Reporting or Recordkeeping Requirements: Office of Management and Budget Review

AGENCY: Nuclear Regulatory Commission

ACTION: Notice of the Office of Management and Budget review of information collection.

SUMMARY: The Nuclear Regulatory Commission (NRC) has recently submitted to the Office of Management and Budget (OMB) for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35).

1. Type of submission, new, revision, or extension: Extension.
2. The title of the information collection:
   10 CFR Part 33 - Specific Domestic Licenses of Broad Scope for Byproduct Material
3. The form number if applicable: Not applicable.
4. How often the collection is required: New applications may be submitted at any time. Renewal applications are submitted every five years.
5. Who will be required or asked to report: Persons applying for or holding an NRC license for broad scope use of radioactive byproduct material. Applicants and licensees are primarily medical institutions, colleges, universities, government agencies, and large private companies engaged in broad educational, research, and development activities.

6. An estimate of the number of responses: 120

7. An estimate of the total number of hours needed to complete the requirement or request: Twenty hours per submittal. The total industry burden is 2,400 hours.

8. An indication of whether Section 3504(h), Pub. L. 96-511 applies:

   Not applicable.

9. Abstract: 10 CFR Part 33 specifies requirements for applying for and being granted licenses authorizing broad scope use of radioactive byproduct material. The information submitted is reviewed by NRC to determine whether the applicant has adequate equipment, facilities, procedures, training, and experience to safely use radioactive material.

Copies of the submittal may be inspected or obtained for a fee from the NRC Public Document Room, 2120 L Street, N.W. (Lower Level), Washington, DC.

Comments and questions may be directed by mail to the OMB reviewer:

Ronald Minsk
Office of Information and Regulatory Affairs (3150-0015)
NEOB-3019
Office of Management and Budget
Washington, DC 20503

Comments may also be communicated by telephone at (202) 395-3084.

The NRC Clearance officer is Brenda Jo. Shelton, (301) 492-8132.

Dated at Bethesda, Maryland, this 30th day of December 1992

For the Nuclear Regulatory Commission

Gerald F. Cranford

Designated Senior Official

for Information Resources Management