

**NUCLEAR REGULATORY COMMISSION**

[NRC–2016–0190]

**Program-Specific Guidance About Commercial Radiopharmacy Licenses****AGENCY:** Nuclear Regulatory Commission.**ACTION:** Draft NUREG; request for comments.

**SUMMARY:** The U.S. Nuclear Regulatory Commission (NRC) is revising its licensing guidance for licenses authorizing commercial nuclear pharmacy use of byproduct material. The NRC is requesting public comment on draft NUREG–1556, Volume 13, Revision 2, “Consolidated Guidance About Materials Licenses: Program-Specific Guidance About Commercial Radiopharmacy Licenses.” The document has been updated from the previous revision to include information on safety culture, security of radioactive materials, protection of sensitive information, and changes in regulatory policies and practices. This document is intended for use by applicants, licensees, and the NRC staff.

**DATES:** Submit comments by March 24, 2017. Comments received after this date will be considered if it is practical to do so, but the NRC is only able to assure consideration of comments received on or before this date.

**ADDRESSES:** You may submit comments by any of the following methods (unless this document describes a different method for submitting comments on a specific subject):

- *Federal Rulemaking Web site:* Go to <http://www.regulations.gov> and search for Docket ID NRC–2016–0190. Address questions about NRC dockets to Carol Gallagher; telephone: 301–415–3463; email: [Carol.Gallagher@nrc.gov](mailto:Carol.Gallagher@nrc.gov). For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.

- *Mail comments to:* Cindy Bladey, Office of Administration, Mail Stop: OWFN–12–H8, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

For additional direction on accessing information and submitting comments, see “Obtaining Information and Submitting Comments” in the **SUPPLEMENTARY INFORMATION** section of this document.

**FOR FURTHER INFORMATION CONTACT:** Said Daibes, Office of Nuclear Material Safety and Safeguards; U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–6863; email: [Said.Daibes@nrc.gov](mailto:Said.Daibes@nrc.gov).

**SUPPLEMENTARY INFORMATION:****I. Obtaining Information and Submitting Comments***A. Obtaining Information*

Please refer to Docket ID NRC–2016–0190 when contacting the NRC about the availability of information regarding this document. You may obtain publicly-available information related to this action by the following methods:

- *Federal Rulemaking Web site:* Go to <http://www.regulations.gov> and search for Docket ID NRC–2016–0190.

- *NRC’s Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly-available documents online in the ADAMS Public Documents collection at <http://www.nrc.gov/reading-rm/adams.html>. To begin the search, select “ADAMS Public Documents” and then select “Begin Web-based ADAMS Search.” For problems with ADAMS, please contact the NRC’s Public Document Room (PDR) reference staff at 1–800–397–4209, 301–415–4737, or by email to [pdr.resource@nrc.gov](mailto:pdr.resource@nrc.gov). The draft NUREG–1556, Volume 13, Revision 2, is available in ADAMS under Accession No. ML16356A040.

- *NRC’s PDR:* You may examine and purchase copies of public documents at the NRC’s PDR, Room O1–F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852.

The draft NUREG–1556, Volume 13, Revision 2, is also available on the NRC’s public Web site on the: (1) “Consolidated Guidance About Materials Licenses (NUREG–1556)” page at <https://www.nrc.gov/reading-rm/doc-collections/nuregs/staff/sr1556>; and the (2) “Draft NUREG–Series Publications for Comment” page at <http://www.nrc.gov/public-involve/doc-comment.html#nuregs>.

*B. Submitting Comments*

Please include Docket ID NRC–2016–0190 in the subject line of your comment submission, in order to ensure that the NRC is able to make your comment submission available to the public in this docket.

The NRC cautions you not to include identifying or contact information that you do not want publicly disclosed in your comment submission. The NRC will post all comment submissions at <http://www.regulations.gov> as well as enter the comment submissions into ADAMS, and the NRC does not routinely edit comment submissions to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for

submission to the NRC, then you should inform those persons not to include identifying or contact information that they do not want to be publicly disclosed in their comment submission. Your request should state that the NRC does not routinely edit comment submissions to remove such information before making the comment submissions available to the public or entering the comment submissions into ADAMS.

**II. Further Information**

NUREG–1556, Volume 13, Revision 2 provides program-specific guidance to assist applicants and licensees in preparing applications for materials licenses for commercial radiopharmacies. In particular, it describes the types of information needed to complete NRC Form 313, “Application for Materials License.” It also provides the NRC with criteria for evaluating a license application. The purpose of this notice is to provide the public with an opportunity to review and provide comments on draft NUREG–1556, Volume 13, Revision 2, “Consolidated Guidance About Materials Licenses: Program-Specific Guidance About Commercial Radiopharmacy Licenses.” These comments will be considered in the final version or subsequent revisions.

This draft NUREG–1556, Volume 13, Revision 2 does not include any revisions associated with the proposed rule “Medical Use of Byproduct Material-Medical Event Definitions, Training and Experience, and Clarifying Amendments.” That proposed rule would amend the following requirements in parts 30, 32, and 35 of title 10 of the *Code of Federal Regulations* related to commercial nuclear pharmacies:

- Removal of the requirement for the board certified nuclear pharmacist to have an attestation statement in addition to the board certificate;
- measuring molybdenum contamination and reporting of failed technetium generators;
- labeling requirements for radioactive drugs; and
- clarifying other revisions to the regulations.

This draft NUREG–1556, Volume 13, Revision 2 does not include any guidance for the proposed rule revisions because that rule is not final at this time.

The proposed rule, “Medical Use of Byproduct Material-Medical Event Definitions, Training and Experience, and Clarifying Amendments,” and proposed changes to NUREG–1556 commercial radiopharmacy licenses

associated with the proposed rule were published for public comment in the **Federal Register** (79 FR 42409 and 79 FR 42224) on July 21, 2014. Comments received on those changes in the proposed rule and guidance are being considered by the NRC staff separately. If the proposed rule becomes final, the proposed revisions to NUREG–1556, Volume 13 addressing the implementation of the proposed rule will be incorporated into NUREG–1556, Volume 13, Revision 2 before its final publication.

Dated at Rockville, Maryland, this 13th day of January, 2017.

For the U.S. Nuclear Regulatory Commission.

**Pamela J. Henderson,**

*Deputy Director, Division of Material Safety, State, Tribal and Rulemaking Programs, Office of Nuclear Material Safety and Safeguards.*

[FR Doc. 2017–01546 Filed 1–23–17; 8:45 am]

**BILLING CODE 7590–01–P**

**U.S. OFFICE OF PERSONNEL MANAGEMENT**

**Submission for Review: Revision of an Existing Information Collection, USAJOBS®, OMB Control No. 3206–0219**

**AGENCY:** U.S. Office of Personnel Management.

**ACTION:** 60-Day Notice and request for comments.

**SUMMARY:** The Office of Personnel Management (OPM) offers the general public and other Federal agencies the opportunity to comment on a revised information collection request (ICR), OMB Control No. 3206–0219, USAJOBS.

**DATES:** Comments are encouraged and will be accepted until March 27, 2017. This process is conducted in accordance with 5 CFR 1320.1.

**ADDRESS:** Interested persons are invited to submit written comments on the proposed information collection to the U.S. Office of Personnel Management, Chief Information Officer, Employee Services IT PMO, USAJOBS, 1900 E. Street NW., Washington, DC 20415, Attention: John Still or send them via electronic mail to [john.still@opm.gov](mailto:john.still@opm.gov).

**FOR FURTHER INFORMATION CONTACT:** A copy of this ICR, with applicable

supporting documentation, may be obtained by contacting the U.S. Office of Personnel Management, Chief Information Officer, Employee Services IT PMO, USAJOBS, 1900 E. Street NW., Washington, DC 20415, Attention: John Still, or by sending a request via electronic mail to [john.still@opm.gov](mailto:john.still@opm.gov)

**SUPPLEMENTARY INFORMATION:** As required by the Paperwork Reduction Act of 1995, (Pub. L. 104–13, 44 U.S.C. chapter 35) as amended by the Clinger-Cohen Act (Pub. L. 104–106), OPM is soliciting comments for this collection. USAJOBS is the Federal Government’s centralized source for most Federal jobs and employment information, including both positions that are required by law to be posted at that location and positions that can be posted there at an agency’s discretion. The Applicant Profile and Resume Builder are two components of the USAJOBS application system.

USAJOBS reflects the minimal critical elements collected across the Federal Government to begin an application for Federal jobs under the authority of sections 1104, 1302, 3301, 3304, 3320, 3361, 3393, and 3394 of title 5, United States Code. This revision proposes to renew a currently approved collection. Therefore, we invite comments that:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected; and
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

**Analysis:**  
**Agency:** Office of Personnel Management.

**Title:** USAJOBS.  
**OMB Number:** 3206–0219.  
**Frequency:** Annually.

**Affected Public:** Individuals.  
**Number of Respondents:** 4,196,336.  
**Estimated Time per Respondent:** 43 minutes.

**Total Burden Hours:** 3,007,374.

U.S. Office of Personnel Management.

**Beth F. Cobert,**  
*Acting Director.*

[FR Doc. 2017–01475 Filed 1–23–17; 8:45 am]

**BILLING CODE 6325–38–P**

**OFFICE OF PERSONNEL MANAGEMENT**

**Excepted Service**

**AGENCY:** U.S. Office of Personnel Management (OPM).

**ACTION:** Notice.

**SUMMARY:** This notice identifies Schedule A, B, and C appointing authorities applicable to a single agency that were established or revoked from September 1, 2016 to September 30, 2016.

**FOR FURTHER INFORMATION CONTACT:** Senior Executive Resources Services, Senior Executive Service and Performance Management, Employee Services, 202–606–2246.

**SUPPLEMENTARY INFORMATION:** In accordance with 5 CFR 213.103, Schedule A, B, and C appointing authorities available for use by all agencies are codified in the Code of Federal Regulations (CFR). Schedule A, B, and C appointing authorities applicable to a single agency are not codified in the CFR, but the Office of Personnel Management (OPM) publishes a notice of agency-specific authorities established or revoked each month in the **Federal Register** at [www.gpo.gov/fdsys/](http://www.gpo.gov/fdsys/). OPM also publishes an annual notice of the consolidated listing of all Schedule A, B, and C appointing authorities, current as of June 30, in the **Federal Register**.

**Schedule A**

No schedule A authorities to report during September 2016.

**Schedule B**

No schedule B authorities to report during September 2016.

The following Schedule C appointing authorities were approved during September 2016.

Agency name	Organization name	Position title	Request No.	Effective date
DEPARTMENT OF AGRICULTURE.	Office of Communications .....	Press Assistant .....	DA160166 .....	09/06/2016
		Deputy Press Secretary .....	DA160176 .....	09/26/2016
	Foreign Agricultural Service .....	Speechwriter & Communications Advisor.	DA160169 .....	09/06/2016