Event	Date
All Parties Status Conference	April 23, 2019.
Evidentiary Hearing Schedule:	
Position Statement/Prefiled Direct Testimony from NRC Staff	May 17, 2019.
Response Position Statements/Prefiled Direct Testimony Supporting NRC Staff's Prefiled Direct Testimony	May 22, 2019
Response Position Statements/Prefiled Response Testimony Opposing NRC Staff's Prefiled Testimony and any Supporting Prefiled Testimony.	June 28, 2019.
Reply Position Statement/Prefiled Reply Testimony from NRC Staff	July 12, 2019.
Proposed Cross-Examination Questions/Requests for Cross-Examination/In Limine Motions on Direct/Response/Reply Testimony Due.	August 2, 2019.
Responses to Requests for Cross-Examination and In Limine Motions on Direct/Response/Reply Testimony Due	August 9, 2019.
Licensing Board Ruling on Requests for Cross-Examination and In Limine Motions	August 19, 2019.
Evidentiary Hearing	August 28–30, 2019.[24]
Proposed Findings of Fact/Conclusions of Law Due	September 27, 2019.
Reply Findings of Fact/Conclusions of Law Due	October 11, 2019.
Licensing Board Initial Decision	November 29, 2019.

[FR Doc. 2019–09532 Filed 5–8–19; 8:45 am]
BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[NRC–2016–0190]

Program-Specific Guidance About Commercial Radiopharmacy Licenses

AGENCY: Nuclear Regulatory Commission.

ACTION: NUREG; issuance.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) has issued Revision 2 to NUREG–1556, Volume 13, ''Consolidated Guidance About Materials Licenses: Program-Specific Guidance About Commercial Radiopharmacy Licenses.'' Volume 13 of NUREG–1556 has been revised to include information on updated regulatory requirements, safety culture, security of radioactive materials, protection of sensitive information, and changes in regulatory policies and practices consistent with current regulations. This volume is intended for use by applicants, licensees, and the NRC staff.

DATES: NUREG–1556, Volume 13, Revision 2, was published in March 2019.

ADDRESSES: 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 document using any of the following methods:

- *Federal Rulemaking Website:* Go to *http://www.regulations.gov* and search for Docket ID NRC–2016–0190. Address questions about NRC Docket IDs in *regulations.gov* to Jennifer Borges; telephone: 301–287–9127; email: *Jennifer.Borges@nrc.gov.* For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.
- *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.* NUREG–1556, Volumes 13, Revision 2, is located at ADAMS Accession Number ML18180A187. This document is also available on the NRC's public website at *http://www.nrc.gov/reading-rm/doc-collections/nuregs/staff/sr1556/* under ''Consolidated Guidance About Materials Licenses (NUREG–1556).''
- *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.

FOR FURTHER INFORMATION CONTACT: Anthony McMurtray, Office of Nuclear Material Safety and Safeguards; U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–2746; email: *Anthony.McMurtray@nrc.gov.*

SUPPLEMENTARY INFORMATION:

I. Discussion

The NRC issued a revision to NUREG–1556, Volumes 13, to provide guidance to existing materials licensees covered under commercial radiopharmacy licenses and to applicants preparing an application for a commercial radiopharmacy license. This NUREG volume also provides the NRC staff with criteria for evaluating commercial radiopharmacy license applications. The purpose of this notice is to notify the public that the NUREG–1556 volume listed in this document was issued as a final report.

II. Additional Information

The NRC published a notice of the availability of the draft report for comment version of NUREG–1556, Volume 13, Revision 2 in the **Federal Register** on January 24, 2017 (82 FR 8227), with a public comment period of 59 days. The public comment period closed on March 24, 2017. Public comments and the NRC staff responses

[24] A final Board assessment regarding the length of the evidentiary hearing will await the receipt of the parties' direct, response, and reply testimony.

to the public comments for NUREG–1556, Volume 13, Revision 2 are available in ADAMS under Accession No. ML18305B303.

III. Congressional Review Act

This NUREG volume is a rule as defined in the Congressional Review Act (5 U.S.C. 801–808). However, the Office of Management and Budget has not found this NUREG revision to be a major rule as defined in the Congressional Review Act.

Dated at Rockville, Maryland, this 2nd day of May 2019.

For the Nuclear Regulatory Commission.

Andrea L. Kock,

Director, Division of Materials Safety, Security, State and Tribal Programs, Office of Nuclear Material Safety and Safeguards.

[FR Doc. 2019–09485 Filed 5–8–19; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

Advisory Committee on the Medical Uses of Isotopes: Meeting Notice

AGENCY: Nuclear Regulatory Commission.

ACTION: Notice of meeting.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) will convene a teleconference meeting of the Advisory Committee on the Medical Uses of Isotopes (ACMUI) on June 10, 2019, to discuss: (1) The revisions to the ACMUI bylaws and (2) the draft report of the ACMUI Regulatory Guide 8.39 Subcommittee. This report will include the subcommittee's comments and recommendations on Phase 1 of the revisions to Regulatory Guide 8.39, ''Release of Patients Administered Radioactive Material.'' A phased approach is being conducted by the NRC staff to comprehensively update Regulatory Guide 8.39. Phase 1 of the revision provides licensees with more detailed instructions to patients before and after they have been administered radioactive material than what is currently provided in Regulatory Guide 8.39. Meeting information, including a copy of the agenda and handouts, will be available at *http://www.nrc.gov/reading-rm/doc-collections/acmui/meetings/2019.html.* The agenda and handouts may also be obtained by contacting Ms. Kellee Jamerson using the information below.

DATES: The teleconference meeting will be held on Monday, June 10, 2019, 2:00 p.m. to 4:00 p.m. Eastern Time.

Public Participation: Any member of the public who wishes to participate in the teleconference should contact Ms. Jamerson using the contact information below or may register for the GoToWebinar for the June 10, 2019, meeting at *https://attendee.gotowebinar.com/register/8147066000509193473.*

Contact Information: Kellee Jamerson, email: *Kellee.Jamerson@nrc.gov,* telephone: (301) 415–7408.

Conduct of the Meeting

Dr. Christopher Palestro, ACMUI Chairman, will preside over the meeting. Dr. Palestro will conduct the meeting in a manner that will facilitate the orderly conduct of business. The following procedures apply to public participation in the meeting:

1. Persons who wish to provide a written statement should submit an electronic copy to Ms. Jamerson at the contact information listed above. All submittals must be received by June 4, 2019, three business days prior to the June 10, 2019, meeting, and must pertain to the topic on the agenda for the meeting.

2. Questions and comments from members of the public will be permitted during the meeting at the discretion of the Chairman.

3. The draft transcript and meeting summary will be available on ACMUI's website *http://www.nrc.gov/reading-rm/doc-collections/acmui/meetings/2019.html* on or about July 23, 2019.

This meeting will be held in accordance with the Atomic Energy Act of 1954, as amended (primarily Section 161a); the Federal Advisory Committee Act (5 U.S.C. App); and the Commission's regulations in 10 CFR part 7.

Dated: May 6, 2019.

Russell E. Chazell,

Federal Advisory Committee Management Officer.

[FR Doc. 2019–09525 Filed 5–8–19; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[NRC–2019–0103]

Information Collection: Criteria and Procedures for Determining Eligibility for Access to or Control Over Special Nuclear Material

AGENCY: Nuclear Regulatory Commission.

ACTION: Renewal of existing information collection; request for comment.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) invites public comment on the renewal of Office of Management and Budget (OMB) approval for an existing collection of information. The information collection is entitled, ''Criteria and Procedures for Determining Eligibility for Access to or Control Over Special Nuclear Material.''

DATES: Submit comments by July 8, 2019. Comments received after this date will be considered if it is practical to do so, but the Commission is able to ensure consideration only for comments received on or before this date.

ADDRESSES: You may submit comments by any of the following methods:

- *Federal Rulemaking website:* Go to *http://www.regulations.gov* and search for Docket ID NRC–2019–0103. Address questions about NRC dockets to Jennifer Borges; telephone: 301–287–9127; email: *Jennifer.Borges@nrc.gov.* For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.
- *Mail comments to:* David Cullison, Office of the Chief Information Officer, Mail Stop: T6–A10M, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

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

FOR FURTHER INFORMATION CONTACT: David Cullison, Office of the Chief Information Officer, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–2084; email: *Infocollects.Resource@nrc.gov.*

SUPPLEMENTARY INFORMATION:

I. Obtaining Information and Submitting Comments

A. Obtaining Information

Please refer to Docket ID NRC–2019–0103 when contacting the NRC about the availability of information for this action. You may obtain publicly-available information related to this action by any of the following methods:

- *Federal Rulemaking website:* Go to *http://www.regulations.gov* and search for Docket ID NRC–2019–0103.
- *NRC's Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly available documents online in the ADAMS Public Document collection at *https://www.nrc.gov/reading-rm/adams.html.* To begin the search, select ''Begin Web-based ADAMS Search.'' For problems with ADAMS, contact the NRC's Public Document Room (PDR)