Act (PRA). These requirements have been approved by OMB (OMB No.: 0625–0245; Expiration Date: 12/31/2014). Public reporting for this collection of information is estimated to be less than 10 minutes per response, including the time for reviewing instructions, and completing and reviewing the collection of information.

Paperwork Reduction Act Data:
OMB Number: 0625–0245.
ITA Number: ITA–4141P.
Type of Review: Regular Submission.

AFFECTED PUBLIC: Business or other for-profit.

Estimated Number of Registered Users: 3,500.
Estimated Time per Response: Less than 10 minutes.
Estimated Total Annual Burden Hours: 92,878 hours.
Estimated Total Annual Costs: $0.00.

Notwithstanding any other provision of law, no person is required to respond to nor shall a person be subject to a penalty for failure to comply with a collection of information subject to the requirements of the Paperwork Reduction Act unless that collection of information displays a current valid OMB Control Number.

Executive Order 12866

This rule has been determined to be not significant for purposes of Executive Order 12866.

Executive Order 13132

This rule does not contain policies with federalism implications as that term is defined in EO 13132.

List of Subjects in 19 CFR Part 360

Administrative practice and procedure, Business and industry, Imports, Reporting and recordkeeping requirements, Steel.

For reasons discussed in the preamble, we propose amending 19 CFR 360 as follows:

PART 360—STEEL IMPORT MONITORING AND ANALYSIS SYSTEM

1. The authority citation for part 360 continues to read as follows:

Authority: 13 U.S.C. 301(a) and 302.

2. Section 360.105 is revised to read as follows.

§ 360.105 Duration of the steel import licensing requirement.

The licensing program will be in effect through March 21, 2017, but may be extended upon review and notification in the "Federal Register" prior to this expiration date. Licenses will be required for all subject imports entered during this period, even if the entry summary documents are not filed until after the expiration of this program. The licenses will be valid for 10 business days after the expiration of this program to allow for the final filing of required Customs documentation.

Dated: November 2, 2012.

Francisco J. Sanchez,
Under Secretary for International Trade.

[FR Doc. 2012–27539 Filed 11–9–12; 8:45 am]

BILLING CODE 3510–05–P

ARCHITECTURAL AND TRANSPORTATION BARRIERS COMPLIANCE BOARD

36 CFR Part 1195

[Docket No. ATBCB–2012–0003]

RIN 3014–AA40

Medical Diagnostic Equipment Accessibility Standards Advisory Committee

AGENCY: Architectural and Transportation Barriers Compliance Board.

ACTION: Notice of advisory committee meeting.

SUMMARY: The Medical Diagnostic Equipment Accessibility Standards Advisory Committee (Committee) will hold its second meeting. The second Committee meeting was originally planned for October 29 and 30, 2012 but cancelled on these dates due to the imminent approach of Hurricane Sandy. On July 5, 2012, the Architectural and Transportation Barriers Compliance Board (Access Board) established an advisory committee to make recommendations to the Board on matters associated with comments received and responses to questions included in a previously published NPRM on Medical Diagnostic Equipment Accessibility Standards. See 77 FR 6916 (February 9, 2012). The NPRM and information related to the proposed standards are available on the Access Board’s Web site at: http://www.access-board.gov/medical-equipment.htm.

The advisory committee will hold its second meeting on December 3 and 4, 2012. The agenda for the meeting is based on the one originally planned for the October 29 and 30, 2012 meeting dates that were cancelled because of Hurricane Sandy. The agenda includes the following:

• Review of previous committee work;
• Formation of subcommittees based on medical diagnostic equipment type;
• Presentation on the proposed transfer surface size and anthropometric data of people who use wheeled mobility devices by Edward Steinfeld, Arch. D., AIA, Director of the Center for Inclusive Design and Environmental Access;
• Continued discussion on transfer surface height and size;
• Review and discussion on permitted obstructions to the transfer surface;
• Consideration of and possible discussion on issues proposed by committee members; and
• Discussion of administrative issues.

The preliminary meeting agenda, along with information about the committee, is available at the Access Board’s Web site (http://www.access-board.gov/medical-equipment.htm).

Committee meetings are open to the public and interested persons can attend the meetings and communicate their views. Members of the public will have opportunities to address the committee on issues of interest to them during public comment periods scheduled on each day of the meeting.

The meetings will be accessible to persons with disabilities. An assistive listening system, computer assisted real-time transcription (CART), and sign language interpreters will be provided. Persons attending the meetings are requested to refrain from using perfume, cologne, and other fragrances for the
comprehension of other participants (see www.access-board.gov/about/policies/fragrance.htm for more information). Also, persons wishing to provide handouts or written information to the committee are requested to provide electronic formats to Rex Pace via email prior to the meetings so that alternate formats can be distributed to committee members.

David M. Capozzi, Executive Director.

[FR Doc. 2012–27516 Filed 11–9–12; 8:45 am]

BILLING CODE 8150–01–P

ENVIRONMENTAL PROTECTION AGENCY
40 CFR Part 52

Approval and Promulgation of Air Quality Implementation Plans; Montana; Revisions to the Administrative Rules of Montana—Air Quality, Subchapter 7, Subchapter 16 and Subchapter 17

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: EPA is proposing to approve new rules and revisions as submitted by the State of Montana on September 23, 2011, as revisions to Montana’s State Implementation Plan. Montana adopted these rules on December 2, 2005, and March 23, 2006. The new rules adopted on December 2, 2005, became state-effective on January 1, 2006; the new rules and revisions adopted on March 23, 2006, became state-effective on April 7, 2006. These new rules and revisions meet the requirements of the Clean Air Act and EPA’s minor new source review regulations. The intended effect of this action is to propose to approve these rules as they are consistent with the Clean Air Act. This action is being taken under section 110 of the Clean Air Act.

DATES: Comments must be received on or before December 13, 2012.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R08–OAR–2012–0846, by one of the following methods:

• www.regulations.gov. Follow the on-line instructions for submitting comments.

• Email: daly.carl@epa.gov and leone.kevin@epa.gov.

• Fax: (303) 312–6064 (please alert the individual listed in the FOR FURTHER INFORMATION CONTACT if you are faxing comments).

• Mail: Carl Daly, Director, Air Program, Environmental Protection Agency (EPA), Region 8, Mailcode 8P–AR, 1595 Wynkoop Street, Denver, Colorado 80202–1129.

• Hand Delivery: Carl Daly, Director, Air Program, Environmental Protection Agency (EPA), Region 8, Mailcode 8P–AR, 1595 Wynkoop Street, Denver, Colorado 80202–1129.Such deliveries are only accepted Monday through Friday, 8:00 a.m. to 4:30 p.m., excluding Federal holidays. Special arrangements should be made for deliveries of boxed information.

Instructions: Direct your comments to Docket ID No. EPA–R08–OAR–2012–0846. EPA’s policy is that all comments received will be included in the public docket without change and may be made available online at www.regulations.gov, including any personal information provided, unless the comment includes information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you consider to be CBI or otherwise protected through www.regulations.gov or email. The www.regulations.gov web site is an “anonymous access” system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an email comment directly to EPA, without going through www.regulations.gov, your email address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the Internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD–ROM you submit. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses. For additional information about EPA’s public docket visit the EPA Docket Center homepage at http://www.epa.gov/epahome/dockets.htm. For additional instructions on submitting comments, go to Section I. General Information of the SUPPLEMENTARY INFORMATION section of this document.

Docket: All documents in the docket are listed in the www.regulations.gov index. Although listed in the index, some information is not publicly available, e.g., CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, will be publicly available only in hard copy. Publicly-available docket materials are available either electronically in www.regulations.gov or in hard copy at the Air Program, Environmental Protection Agency (EPA), Region 8, 1595 Wynkoop Street, Denver, Colorado 80202–1129. EPA requests that if at all possible, you contact the individual listed in the FOR FURTHER INFORMATION CONTACT section to view the hard copy of the docket. You may view the hard copy of the docket Monday through Friday, 8:00 a.m. to 4:00 p.m., excluding Federal holidays.

FOR FURTHER INFORMATION CONTACT:
K. Leone, Air Program, Mailcode 8P–AR, Environmental Protection Agency, Region 8, 1595 Wynkoop Street, Denver, Colorado 80202–1129, (303) 312–6227, or leone.kevin@epa.gov.

SUPPLEMENTARY INFORMATION:

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I. General Information
II. What is being addressed in this proposed action?
III. What Authorities Apply to EPA’s Proposed Action
IV. EPA’s Review and Proposed Action on SIP Revisions
V. Summary of EPA’s Proposed Action
VI. Statutory and Executive Order Reviews

Definitions

For the purpose of this document, we are giving meaning to certain words or initials as follows:

(i) The words or initials Act or CAA mean or refer to the Clean Air Act, unless the context indicates otherwise.

(ii) The initials ARM mean or refer to the Administrative Rule of Montana.

(iii) The words EPA, we, us or our mean or refer to the United States Environmental Protection Agency.

(iv) The initials MACT mean Maximum Achievable Control Technology.

(v) The initials MAQP mean Montana Air Quality Permit.

(vi) The initials MHRP mean Monitoring, Reporting and Recordkeeping.

(vii) The initials NAAQS mean National Ambient Air Quality Standards.


(ix) The initials NSR mean or refer to new source review, a phrase intended to encompass the stationary source regulatory programs that regulate the construction and modification of stationary sources as provided under CAA section 110(a)(2)(C), CAA Title I,