Pollutant Discharge Elimination System (NPDES) general permits for potable water treatment facility (PWTF) discharges to certain waters of the Commonwealth of Massachusetts (included both Commonwealth and Indian country lands) and the State of New Hampshire. These General Permits replace the previous PWTF GP, which expired on November 15, 2005.

The final PWTF GP establishes Notice of Intent (NOI) requirements, effluent limitations, standards, prohibitions, and management practices for facilities with discharges from potable water treatment facilities. Owners and/or operators of these facilities, including those currently authorized to discharge under the expired General Permit, will be required to submit an NOI to be covered by the PWTF GP to both EPA-New England and the appropriate state agency. After EPA and the State have reviewed the NOI, the facility will receive a written notification from EPA of permit coverage and authorization to discharge under the General Permit.

DATES: The general permits shall be effective on the date of signature and will expire at midnight, five (5) years from the last day of the month preceding the effective date.

ADDRESSES: The required notification information to obtain permit coverage is provided for in the general permits. This information shall be submitted to both EPA and the appropriate State. Notification information may be sent via USPS, e-mail or fax to EPA at EPA–Region 1, Office of Ecosystem Protection (CMU), One Congress Street, Boston, Massachusetts 02114–2023; e-mail address *PWTF.GeneralPermit@EPA.GOV;* or fax number (617) 918–0505. Notification information shall be submitted to the appropriate State agency at the addresses listed in Appendix VI of the general permits.

FOR FURTHER INFORMATION CONTACT: Additional information concerning the final PWTF GP may be obtained between the hours of 9 a.m. and 5 p.m. Monday through Friday, excluding holidays, from Damien Houlihan at *Houlihan.Damien@epa.gov* or (617) 918–1586. The general permits may be viewed over the Internet at the EPA Web site *http://www.epa.gov/region1/npdes/pwtfgp.html.* To obtain a paper copy of the general permits, please contact Mr. Houlihan using the contact information provided above. A reasonable fee may be charged for copying requests.

Dated: September 24, 2009.

Ira Leighton,

Acting Regional Administrator, Region 1.

[FR Doc. E9–23791 Filed 10–1–09; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–ORD–2009–0658; FRL–8965–4]

Human Studies Review Board; Notice of Public Meeting

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The U.S. Environmental Protection Agency's (EPA or Agency) Office of the Science Advisor (OSA) announces a public meeting of the Human Studies Review Board (HSRB) to advise the Agency on EPA's scientific and ethical reviews of research with human subjects.

DATES: The public meeting will be held from October 20–21, 2009, from approximately 10 a.m. to approximately 5:30 p.m., through October 20, 2009 from approximately 8:30 a.m. to approximately 12:30 p.m. Eastern Time.

Location: Environmental Protection Agency, Conference Center—Lobby Level, One Potomac Yard (South Bldg.), 2777 S. Crystal Drive, Arlington, VA 22202.

Meeting Access: Seating at the meeting will be on a first-come basis. To request accommodation of a disability, please contact the person listed under **FOR FURTHER INFORMATION CONTACT** at least 10 business days prior to the meeting, to allow EPA as much time as possible to process your request.

Procedures for Providing Public Input: Interested members of the public may submit relevant written or oral comments for the HSRB to consider during the advisory process. Additional information concerning submission of relevant written or oral comments is provided in Unit I.D. of this notice.

FOR FURTHER INFORMATION CONTACT: Any member of the public who wishes further information should contact Jim Downing, EPA, Office of the Science Advisor, (8105R), Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460; telephone number: (202) 564–2468; fax: (202) 564–2070; e-mail addresses: *downing.jim@epa.gov.* General information concerning the EPA HSRB can be found on the EPA Web site at *http://www.epa.gov/osa/hsrb/.*

ADDRESSES: Submit your written comments, identified by Docket ID No. EPA–HQ–ORD–2009–0658, by one of the following methods:

Internet: http://www.regulations.gov: Follow the on-line instructions for submitting comments.

E-mail: ord.docket@epa.gov.

Mail: Environmental Protection Agency, EPA Docket Center (EPA/DC), ORD Docket, Mailcode: 28221T, 1200 Pennsylvania Ave., NW., Washington, DC 20460.

Hand Delivery: The EPA/DC Public Reading Room is located in the EPA Headquarters Library, Room Number 3334 in the EPA West Building, located at 1301 Constitution Ave., NW., Washington, DC 20460. The hours of operation are 8:30 a.m. to 4:30 p.m. Eastern Time, Monday through Friday, excluding Federal holidays. Please call (202) 566–1744 or e-mail the ORD Docket at *ord.docket@epa.gov* for instructions. Updates to Public Reading Room access are available on the Web site (*http://www.epa.gov/epahome/dockets.htm*).

Instructions: Direct your comments to Docket ID No. EPA–HQ–ORD–2009–0658. EPA's policy is that all comments received will be included in the public docket without change and may be made available online at *http://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 *http://www.regulations.gov* or e-mail. The *http://www.regulations.gov* website 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 e-mail comment directly to EPA, without going through *http://www.regulations.gov,* your e-mail 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.

SUPPLEMENTARY INFORMATION:

I. Public Meeting

A. Does This Action Apply to Me?

This action is directed to the public in general. This action may, however, be of interest to persons who conduct or assess human studies, especially studies on substances regulated by EPA or to persons who are or may be required to conduct testing of chemical substances under the Federal Food, Drug, and Cosmetic Act (FFDCA) or the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). Since other entities may also be interested, the Agency has not attempted to describe all the specific entities that may be affected by this action. If you have any questions regarding the applicability of this action to a particular entity, consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

B. How Can I Access Electronic Copies of This Document and Other Related Information?

In addition to using regulations.gov, you may access this **Federal Register** document electronically through the EPA Internet under the ''Federal Register'' listings at *http://www.epa.gov/fedrgstr/*.

Docket: All documents in the docket are listed in the *http://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 *http://www.regulations.gov* or in hard copy at the ORD Docket, EPA/DC, Public Reading Room. The EPA/DC Public Reading Room is located in the EPA Headquarters Library, Room Number 3334 in the EPA West Building, located at 1301 Constitution Ave., NW., Washington, DC 20460. The hours of operation are 8:30 a.m. to 4:30 p.m. EST, Monday through Friday, excluding Federal holidays. Please call (202) 566–1744 or e-mail the ORD Docket at *ord.docket@epa.gov* for instructions. Updates to Public Reading Room access are available on the Web site (*http://www.epa.gov/epahome/dockets.htm*).

EPA's position paper(s), charge/questions to the HSRB, and the meeting agenda will be available by early October 2009. In addition, the Agency may provide additional background documents as the materials become available. You may obtain electronic copies of these documents, and certain other related documents that might be available electronically, from the regulations.gov website and the EPA HSRB Web site at *http://www.epa.gov/osa/hsrb/*. For questions on document availability or if you do not have access to the Internet, consult the person listed under **FOR FURTHER INFORMATION**.

C. What Should I Consider as I Prepare My Comments for EPA?

You may find the following suggestions helpful for preparing your comments:

a. Explain your views as clearly as possible.

b. Describe any assumptions that you used.

c. Provide copies of any technical information and/or data that you used to support your views.

d. Provide specific examples to illustrate your concerns and suggest alternatives.

e. To ensure proper receipt by EPA, be sure to identify the docket ID number

assigned to this action in the subject line on the first page of your response. You may also provide the name, date, and **Federal Register** citation.

D. How May I Participate in This Meeting?

You may participate in this meeting by following the instructions in this section. To ensure proper receipt by EPA, it is imperative that you identify docket ID number EPA–HQ–ORD–2009–0658 in the subject line on the first page of your request.

a. Oral comments. Requests to present oral comments will be accepted up to October 13, 2009. To the extent that time permits, interested persons who have not pre-registered may be permitted by the Chair of the HSRB to present oral comments at the meeting. Each individual or group wishing to make brief oral comments to the HSRB is strongly advised to submit their request (preferably via e-mail) to the person listed under **FOR FURTHER INFORMATION CONTACT** no later than noon, Eastern time, October 13, 2009, in order to be included on the meeting agenda and to provide sufficient time for the HSRB Chair and HSRB Designated Federal Officer (DFO) to review the agenda to provide an appropriate public comment period. The request should identify the name of the individual making the presentation, the organization (if any) the individual will represent, and any requirements for audiovisual equipment (e.g., overhead projector, LCD projector, chalkboard). Oral comments before the HSRB are limited to five minutes per individual or organization. Please note that this limit applies to the cumulative time used by all individuals appearing either as part of, or on behalf of an organization. While it is our intent to hear a full range of oral comments on the science and ethics issues under discussion, it is not our intent to permit organizations to expand these time limitations by having numerous individuals sign up separately to speak on their behalf. If additional time is available, there may be flexibility in time for public comments. Each speaker should bring 25 copies of his or her comments and presentation slides for distribution to the HSRB at the meeting.

b. Written comments. Although you may submit written comments at any time, for the HSRB to have the best opportunity to review and consider your comments as it deliberates on its report, you should submit your comments at least five business days prior to the beginning of the meeting. If you submit comments after this date, those comments will be provided to the Board members, but you should recognize that the Board members may not have adequate time to consider those comments prior to making a decision. Thus, if you plan to submit written comments, the Agency strongly encourages you to submit such comments no later than noon, Eastern Time, October 13, 2009. You should submit your comments using the instructions in Unit I.C. of this notice. In addition, the Agency also requests that person(s) submitting comments directly to the docket also provide a copy of their comments to the person listed under **FOR FURTHER INFORMATION CONTACT.** There is no limit on the length of written comments for consideration by the HSRB.

E. Background

a. Topics for discussion. The HSRB is a Federal advisory committee operating in accordance with the Federal Advisory Committee Act (FACA) 5 U.S.C. App. 2 § 9. The HSRB provides advice, information, and recommendations to EPA on issues related to scientific and ethical aspects of human subjects research. The major objectives of the HSRB are to provide advice and recommendations on: (1) Research proposals and protocols; (2) reports of completed research with human subjects; and (3) how to strengthen EPA's programs for protection of human subjects of research. The HSRB reports to the EPA Administrator through EPA's Science Advisor.

At its meeting on October 20–21, 2009, EPA's Human Studies Review Board will consider scientific and ethical issues surrounding three topics:

1. Two published reports of completed, pre-rule research on the effects of exposure to pesticides containing pyrethrins/pyrethroids. EPA requests the advice of the HSRB on the scientific merit, relevancy, and limitations of these studies, and on their ethical acceptability. EPA intends to incorporate into a future revision of the EPA White Paper, ''A Review of the Relationship between Pyrethrins, Pyrethroid Exposure and Asthma and Allergies,'' a discussion of either or both of these studies if they are deemed to be scientifically sound, relevant and ethically acceptable.

2. A proposal for new research to be conducted by Carroll-Loye Biological Research to evaluate in the laboratory the repellent efficacy to ticks of two registered products containing 20% picaridin. EPA requests the advice of the HSRB concerning whether, if it is revised as suggested in EPA's review and if it is performed as described, this research is likely to generate scientifically reliable data, useful for assessing the efficacy of the tested materials in repelling ticks, and to meet the applicable requirements of 40 CFR part 26, subparts K and L.

3. A new scenario design and associated protocol from the Antimicrobials Exposure Assessment Task Force II (AEATF–II), describing proposed research to monitor at three sites the dermal and inhalation exposure of professional janitorial workers who apply an antimicrobial pesticide formulated as an aerosol spray. EPA requests the advice of the HSRB concerning whether, if it is revised as suggested in EPA's review and if it is performed as described, this research is likely to generate scientifically reliable data, useful for assessing the exposure of those who apply antimicrobial pesticides as aerosols, and to meet the applicable requirements of 40 CFR part 26, subparts K and L.

In addition, the Board will be reviewing its draft June 24–25, 2009, meeting report for subsequent Board approval. Finally, the HSRB may also discuss planning for future HSRB meetings.

b. Meeting minutes and reports. Minutes of the meeting, summarizing the matters discussed and recommendations, if any, made by the advisory committee regarding such matters, will be released within 90 calendar days of the meeting. Such minutes will be available at *http://www.epa.gov/osa/hsrb/* and *http://www.regulations.gov*. In addition, information concerning a Board meeting report, if applicable, can be found at *http://www.epa.gov/osa/hsrb/* or from the person listed under **FOR FURTHER INFORMATION CONTACT.**

Dated: September 24, 2009.

Kevin Teichman,

EPA Acting Science Advisor.

[FR Doc. E9–23795 Filed 10–1–09; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection Being Submitted to the Office of Management and Budget for Review and Approval, Comments Requested

September 28, 2009.

SUMMARY: The Federal Communications Commission, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s), as required by the Paperwork Reduction Act of 1995, 44 U.S.C. 3501–3520. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number. Comments are requested concerning (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology.

DATES: Persons wishing to comment on this information collection should submit comments on November 2, 2009. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Nicholas A. Fraser, Office of Management and Budget (OMB), via fax at (202) 395–5167, or via the Internet at Nicholas_A._Fraser@omb.eop.gov and to Judith B. Herman, Federal Communications Commission (FCC). To submit your comments by e–mail send then to: PRA@fcc.gov. To view a copy of this information collection request (ICR) submitted to OMB: (1) Go to web page: *http://www.reginfo.gov/public/do/PRAMain*, (2) look for the section of the web page called ''Currently Under Review'', (3) click on the downward–pointing arrow in the ''Select Agency'' box below the ''Currently Under Review'' heading, (4) select ''Federal Communications Commission'' from the list of agencies presented in the ''Select Agency'' box, (5) click the ''Submit'' button to the right of the ''Select Agency'' box, and (6) when the FCC list appears, look for the title of this ICR (or its OMB Control Number, if there is one) and then click on the ICR.

FOR FURTHER INFORMATION CONTACT: Judith B. Herman, 202–418–0214. For additional information about the information collection(s) send an e–mail to *PRA@fcc.gov* or contact Judith B. Herman, 202–418–0214.

SUPPLEMENTARY INFORMATION:

OMB Control No: 3060–0233.

Title: Part 36, Separations.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for–profit.

Number of Respondents: 1,997 respondents; 7,562 responses.

Estimated Time Per Response: 5 – 22 hours.

Frequency of Response: On occasion, annual and quarterly reporting requirements and third party disclosure requirements.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for information collection is contained in 47 U.S.C. sections 151, 154(i), and (j), 221(c) and 410(c).

Total Annual Burden: 71,283 hours.

Privacy Act Impact Assessment: N/A

Nature and Extent of Confidentiality: No assurance of confidentiality has been given regarding the information.

Need and Uses: The Commission will submit this information collection to the Office of Management and Budget (OMB) during this 30 day comment period in order to obtain the full three year clearance from them. The Commission is requesting an extension (no change in the reporting and/or third party disclosure requirements) of this information collection. There is a change in the estimated respondents/responses and the annual burden hours. The Commission is reporting a 1,774 adjusted increase in the number of responses and a 12,865 hour increase in the total annual burden hours.

In order to determine which carriers are entitled to universal service support,