ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act (44 U.S.C. 3501 *et seq.*), this document announces that the following Information Collection Request (ICR) has been forwarded to the Office of Management and Budget (OMB) for review and approval: NESHAP, Subpart GGG, Pharmaceutical Production, OMB Control Number 2060–0358, expiration date 7/31/00. The ICR describes the nature of the information collection and its expected burden and cost; where appropriate, it includes the actual data collection instrument.

DATES: Comments must be submitted on or before July 27, 2000.

FOR FURTHER INFORMATION CONTACT: For a copy of the ICR contact Sandy Farmer at EPA by phone at (202) 260–2740, by E–Mail at Farmer.Sandy@epamail.epa.gov or download off the Internet at *http://www.epa.gov/icr* and refer to EPA ICR No. 1781.02. For technical questions about the ICR contact Marcia Mia at 202–564–7042.

SUPPLEMENTARY INFORMATION:

Title: NESHAP, subpart GGG, Pharmaceuticals Production (OMB Control No. 2060–0358; EPA ICR No. 1781.02) expiring 07/31/00. This is a request for extension of a currently approved collection.

Abstract: In general all NESHAP require initial notifications, performance tests, and periodic reports. Owners or operators are also required to maintain records of the occurrence and duration of any startup, shutdown, or malfunction in the operation of an affected facility, or any period during which the monitoring system is inoperative.

Any owner or operator subject to the provisions of this part shall maintain a file of these measurements, and retain the file for at least 5 years following the date of such measurements, maintenance reports, and records. All reports are sent to the delegated State or Local authority and are entered into the AIRS database.

The National Emission Standards for Hazardous Air Pollutants (NESHAP) for Pharmaceuticals Production were proposed on April 2, 1997 and promulgated on September 21, 1998. These standards apply to the facilities in Pharmaceuticals Production that are major sources of hazardous air pollutants (HAP). The affected facility is all pharmaceutical manufacturing operations including process vents, storage tanks, equipment components, and wastewater systems commencing construction or reconstruction after the date of proposal. This information is being collected to assure compliance with 40 CFR part 63, subpart GGG.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control numbers for EPA's regulations are listed in 40 CFR part 9 and 48 CFR Chapter 15. The **Federal Register** document required under 5 CFR 1320.8(d), soliciting comments on this collection of information was published on 03/31/00 (65 FR 17258); no comments were received.

Burden Statement: The annual public reporting and recordkeeping burden for this collection of information is estimated to average 409 hours per response. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, or disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; develop, acquire, install, and utilize technology and systems for the purposes of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; adjust the existing ways to comply with any previously applicable instructions and requirements; train personnel to be able to respond to a collection of information; search data sources; complete and review the collection of information; and transmit or otherwise disclose the information.

Respondents/Affected Entities: Pharmaceutical Production Plants.

Estimated Number of Respondents: 103.

Frequency of Response: Initial, quarterly, semiannually and on occasion.

Estimated Total Annual Hour Burden: 84,275 hours.

Estimated Total Annualized Capital, O&M Cost Burden: $0.

Send comments on the Agency's need for this information, the accuracy of the provided burden estimates, and any suggested methods for minimizing respondent burden, including through the use of automated collection techniques to the following addresses. Please refer to EPA ICR No. 1781.02 and OMB Control No. 2060–0358 in any correspondence.

Ms. Sandy Farmer,
U.S. Environmental Protection Agency,
Office of Environmental Information,
Collection Strategies Division (2822),
1200 Pennsylvania Ave., NW,
Washington, DC 20460; and

Office of Information and Regulatory Affairs,
Office of Management and Budget,
Attention: Desk Officer for EPA,
725 17th Street, NW,
Washington, DC 20503.

Dated: June 19, 2000.

Oscar Morales,

Director, Collection Strategies Division.

[FR Doc. 00–16178 Filed 6–26–00; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–6725–5]

National Drinking Water Advisory Council; Contaminant Candidate List and 6-Year Review of Existing Regulations Working Group; Notice of Open Meeting

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: Under section 10(a)(2) of Public Law 92–423, ''The Federal Advisory Committee Act,'' notice is hereby given that a meeting of the Contaminant Candidate List (CCL) Regulatory Determination and 6-Year Review of Existing Regulations Working Group of the National Drinking Water Advisory Council established under the Safe Drinking Water Act, as amended (42 U.S.C. S300f *et seq.*), will be held July 10, 2000, from 1:00 pm–5:00 pm ET (approximately), at the U.S. EPA, 401 M Street, S.W., Suite 925B, Washington, D.C. 20460. The meeting is open to the public to observe and statements will be taken from the public as time allows. Seating is limited.

This is the second of three scheduled meetings to address the 6-Year Review of Existing Regulations. The Working Group will recommend a protocol for selecting existing NPDWRs for possible revision and develop specific recommendations for analyzing and presenting the available scientific data (The Working Group does not plan to discuss specific contaminants as a part of this exercise.) Final recommendations will be forwarded to the full NDWAC for further consideration.

At the last meeting, the Working Group formed three sub-groups to revise specific portions of the strawman protocol. The sub-groups will forward their final products to EPA for consolidation. EPA will consolidate comments and distribute a revised draft to Working Group members for discussion on July 10, 2000.

For more information, contact April McLaughlin, Designated Federal Officer,

Contaminant Candidate List and Regulatory Determination and 6-Year Review of Existing Regulations Working Group, U.S. EPA (4607), Office of Ground Water and Drinking Water, 401 M Street SW, Washington, DC 20460. The email address is: mclaughlin.april@epa.gov. or call 202–260–5524.

Dated: June 20, 2000.

Janet Pawlukiewicz,

Acting Deputy Director, Office of Ground Water and Drinking Water.

[FR Doc. 00–16179 Filed 6–26–00; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–6725–3]

Science Advisory Board; Notification of Public Advisory Committee Meeting

Meeting Notice—Executive Committee—July 12–13, 2000

Pursuant to the Federal Advisory Committee Act, Public Law 92–463, notice is hereby given that the Science Advisory Board's (SAB's) Executive Committee will conduct a public meeting on Wednesday and Thursday, July 12–13, 2000. The meeting will convene each day at 8:30 am at the EPA Office of Administration Auditorium located at 79 T.W. Alexander Drive in Research Triangle Park, NC and adjourn no later than 5:30 pm. All times noted are Eastern Daylight Time. The meeting is open to the public, however, seating is limited and available on a first come basis.

Purpose of the Meeting—At this meeting, the Executive Committee will receive updates from its committees and subcommittees concerning their recent and planned activities. As part of these updates, some committees will present draft reports for Executive Committee review and approval. Tentatively anticipated drafts include, but are not limited to the *Executive Committee Scientific and Technological Achievement Awards Subcommittee:* Review of the Report on ''Scientific and Technological Achievement Awards.''

As part of this two day meeting, the Executive Committee will also: (a) meet with various Agency officials to discuss matters of mutual interest such as the scope and breadth of R&D activities performed at RTP, including a poster presentation the afternoon of July 12 to be held in Classroom One of the Environmental Research Center, Highway 54 and T. W. Alexander Drive, Research Triangle Park, NC; (b) receive briefings from Agency staff on various topics, including an update of the Integrated Risk Information System (IRIS) project; (c) conduct the third in a series of Workshops on the role of science in some of the Agency's innovative approaches to environmental decisionmaking focusing on new approaches to stakeholder involvement; and, (d) discuss options for activities the Board might undertake to improve the use of science at the science policy interface.

Availability of Materials—The timing of these events will be included in an agenda for the meeting that should be available one week prior to the meeting. Drafts of the reports that will be reviewed at the meeting should be available to the public at the SAB website (http://www.epa.gov/sab) by close-of-business on July 5.

For Further Information—Any member of the public wishing further information concerning this meeting or wishing to submit *brief* oral comments should contact Dr. John R. Fowle III, Designated Federal Officer (DFO) for the Executive Committee, *in writing,* no later than close of business July 7, 2000 at USEPA Science Advisory Board (1400A), 1200 Pennsylvania Avenue, NW, Washington, DC 20460; fax (202) 501–0323; or via e-mail at <fowle.john@epa.gov>. Those wishing further information concerning the meeting should contact Dr. Fowle at (202) 564–4533.

Providing Oral or Written Comments at SAB Meetings

It is the policy of the Science Advisory Board to accept written public comments of any length, and to accommodate oral public comments whenever possible. The Science Advisory Board expects that public statements presented at its meetings will not be repetitive of previously submitted oral or written statements. *Oral Comments:* In general, each individual or group requesting an oral presentation at a face-to-face meeting will be limited to a total time of ten minutes. For teleconference meetings, opportunities for oral comment will usually be limited to no more than three minutes per speaker and no more than fifteen minutes total. Deadlines for getting on the public speaker list for a meeting are given above. Speakers should bring at least 35 copies of their comments and presentation slides for distribution to the reviewers and public at the meeting. *Written Comments:* Although the SAB accepts written comments until the date of the meeting (unless otherwise stated), written comments should be received in the SAB Staff Office at least one week prior to the meeting date so that the comments may be made available to the committee for their consideration. Comments should be supplied to the appropriate DFO at the address/contact information noted above in the following formats: one hard copy with original signature, and one electronic copy via e-mail (acceptable file format: WordPerfect, Word, or Rich Text files (in IBM–PC/Windows 95/98 format). Those providing written comments and who attend the meeting are also asked to bring 25 copies of their comments for public distribution.

General Information—Additional information concerning the Science Advisory Board, its structure, function, and composition, may be found on the SAB Website (http://www.epa.gov/sab) and in *The FY1999 Annual Report of the Staff Director* which is available from the SAB Publications Staff at (202) 564–4533 or via fax at (202) 501–0256. Committee rosters, draft Agendas and meeting calendars are also located on our website.

Meeting Access—Individuals requiring special accommodation at this meeting, including wheelchair access to the conference room, should contact the DFO at least five business days prior to the meeting so that appropriate arrangements can be made.

Dated: June 19, 2000.

Donald G. Barnes,

Staff Director, Science Advisory Board.

[FR Doc. 00–16177 Filed 6–26–00; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–6725–7]

Regulatory Reinvention (XL) Pilot Projects; Project XL Proposed Final Project Agreement: Progressive Insurance Company

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of availability of the Project XL Proposed Final Project Agreement: Progressive Insurance Project—Pay-as-you-Drive Auto Insurance.

SUMMARY: EPA is requesting comments on a proposed Project XL Final Project Agreement (FPA) for the Progressive Auto Insurance Company (hereafter ''Progressive''). The FPA is a voluntary agreement developed collaboratively by Progressive and the EPA.

DATES: Comments are due on or before July 11, 2000.