Shoreline II, Outfitter and Guide Management Plan), Comment Period Ends:, 03/14/2016, Contact: Carey Case 907–772–3871.

EIS No. 20160018, Draft, NPS, MD, Assateague Island National Seashore General Management Plan, Comment Period Ends: 05/01/2016, Contact: Deborah Darden 410–629–6080.

EIS No. 20160019, Final, USFS, OR, Kahler Dry Forest Restoration, Review Period Ends: 03/14/2016, Contact: John Evans 541–278–3869.

EIS No. 20160020, Draft, USFS, CA, Lassen National Forest Over-Snow Vehicle (OSV) Use Designation, Comment Period Ends: 03/15/2016, Contact: Chris Obrien 530–252–6698.

EIS No. 20160021, Draft, USACE, NY, Mamaroneck and Sheldrake Rivers Flood Risk Management Village of Mamaroneck General Reevaluation, Comment Period Ends: 03/14/2016, Contact: Matthew Voisine 917–790–8718.

Amended Notices

EIS No. 20150312, Draft, FRA, NY, NEC FUTURE Tier 1, Comment Period Ends: 02/16/2016, Contact: Rebecca Reyes-Alicea 212–668–2282, Revision to FR Notice Published 11/13/2015; Extending Comment Period from 01/30/2016 to 02/16/2016.

EIS No. 20150353, Draft, FRA, MD, Baltimore and Potomac Tunnel Project, Comment Period Ends: 02/19/2016, Contact: Michelle W. Fishburne 202–293–0398, Revision to FR Notice Published 12/18/20105; Extending Comment Period from 02/05/2016 to 02/19/2016.

EIS No. 20150358, Draft, USACE, FL, Herbert Hoover Dike Dam Safety Modification, Comment Period Ends: 02/23/2016, Contact: Stacie Auvenshine 904–232–3694, Revision to FR Notice Published 12/24/2015; Correction to Comment Period from 02/08/2016 to 02/23/2016.

EIS No. 20160001, Final, FHWA, CO, I–70 East, Review Period Ends: 03/02/2016, Contact: Chris Horn 720–963–3017, Revision to FR Notice Published 01/15/2016; Extending Comment Period from 02/16/2016 to 03/02/2016.

EIS No. 20160008, Draft, USFS, WY, Withdrawn—Bear Lodge Project, Contact: Jeanette Timm 307–283–1361, Revision to FR Notice Published 01/15/2016; The U.S. Department of Agriculture's Forest Service has Officially Withdrawn this EIS.

Dated: January 26, 2016.

Dawn Roberts,

Management Analyst, NEPA Compliance Division, Office of Federal Activities.

[FR Doc. 2016–01706 Filed 1–28–16; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–153]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by March 29, 2016.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–R–153 Medicaid Drug Use Review (DUR) Program

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Medicaid Drug Use Review (DUR) Program; *Use:* States must provide for a review of drug therapy before each prescription is filled

or delivered to a Medicaid patient. This review includes screening for potential drug therapy problems due to therapeutic duplication, drug-disease contraindications, drug-drug interactions, incorrect drug dosage or duration of drug treatment, drug-allergy interactions, and clinical abuse/misuse. Pharmacists must make a reasonable effort to obtain, record, and maintain Medicaid patient profiles. These profiles must reflect at least the patient's name, address, telephone number, date of birth/age, gender, history, *e.g.,* allergies, drug reactions, list of medications, and pharmacist's comments relevant to the individual's drug therapy.

The State must conduct RetroDUR which provides for the ongoing periodic examination of claims data and other records in order to identify patterns of fraud, abuse, inappropriate or medically unnecessary care. Patterns or trends of drug therapy problems are identified and reviewed to determine the need for intervention activity with pharmacists and/or physicians. States may conduct interventions via telephone, correspondence, or face-to-face contact.

Annual reports are submitted to CMS for the purposes of monitoring compliance and evaluating the progress of States' DUR programs. The information submitted by States is reviewed and results are compiled by CMS in a format intended to provide information, comparisons and trends related to States' experiences with DUR. The States benefit from the information and may enhance their programs each year based on State reported innovative practices that are compiled by CMS from the DUR annual reports. *Form Number:* CMS–R–153 (OMB Control Number 0938–0659); *Frequency:* Yearly, quarterly, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 510; *Total Annual Hours:* 20,808. (For policy questions regarding this collection contact Renee Hilliard at 410–786–2991).

Dated: January 25, 2016.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2016–01688 Filed 1–28–16; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10519]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by February 29, 2016.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR,* Email: *OIRA_submission@omb.eop.gov.*

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Physician Quality Reporting System (PQRS) and the Electronic Prescribing Incentive (eRx) Program Data Assessment, Accuracy and Improper Payments Identification Support; *Use:* The incentive and reporting programs have data integrity issues, such as rejected and improper payments. This four year project will evaluate incentive payment information for accuracy and identify improper payments, with the goal of recovering these payments. Additionally, based on the project's results, recommendations will be made so that we can avoid future data integrity issues.

Data submission, processing, and reporting will be analyzed for potential errors, inconsistencies, and gaps that are related to data handling, program requirements, and clinical quality measure specifications of PQRS and eRx program. Surveys of Group Practices, Registries, and Data Submission Vendors (DSVs) will be conducted in order to evaluate the PQRS and eRx Incentive Program. Follow-up interviews will occur with a small number of respondents. *Form Number:* CMS–10519 (OMB control number: 0938–1255); *Frequency:* Annually; *Affected Public:* Business or other for-profits; *Number of Respondents:* 115;