DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Advance Planning Document (APD) Process.

OMB No.: 0970–0417.

Description: The Advance Planning Document (APD) process, established in the rules at 45 CFR part 95, subpart F, is the procedure by which states request and obtain approval for Federal Financial Participation (FFP) in their cost of acquiring Automated Data Processing (ADP) equipment and services.

State child support agencies are required to establish and operate a federally approved statewide ADP and information retrieval system to assist in child support enforcement. States are required to submit an initial APD, containing information to assist the Secretary of the Department of Health and Human Services (HHS) in determining if the state computerized support enforcement project planning and implementation meets federal certification requirements needed for the approval of FFP. States are then required to submit annual APD updates to provide project status updates to HHS, as well as, to request ongoing FFP for systems development, enhancements, operations and maintenance. As-Needed APDs are also submitted to acquire FFP when major milestone are missed or significant changed to project schedules occur. Based on an assessment of the information provided in APD, states that do not meet the federal requirements necessary for approval are required to conduct periodic independent verification and validation (IV&V) services for high risk project oversight.

In addition to the APDs providing HHS with the information necessary to determine the allowable level of federal funding for state systems projects, states also submit associated procurement and data security documents, such as the request for proposals (RFPs), contracts, contract amendments, and the biennial security review reports.

Respondents: State Child Support Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
RFP and Contract	54	1.5	4	324
Emergency Funding Request	5	.1	2	1
Biennial Reports	54	1	1.50	81
Advance Planning Document	34	1.2	120	4,896
Operational Advance Planning Document	20	1	30	600
Independent Verification and Validation (ongoing)	3	4	10	120
Independent Verification and Validation (semiannually)	1	2	16	32
Independent Verification and Validation (quarterly)	1	4	30	120
System Certification	1	1	240	240

Estimated Total Annual Burden Hours: 6,414.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–17196 Filed 8–14–17; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Reallotment of FY 2017 Funds

AGENCY: Administration on Intellectual and Developmental Disabilities (AIDD), Administration on Disabilities (AoD), Administration for Community Living (ACL), HHS.

ACTION: Notice of reallotment of FY 2017 funds.

SUMMARY: AIDD intends to reallot funds under authority of the Development Disabilities Assistance and Bill of Rights Act of 2000 which states: ''If the Secretary determines that an amount of an allotment to a State for a period (of a fiscal year or longer) will not be required by the State during the period for the purpose for which the allotment was made, the Secretary may reallot the amount.''

AIDD will be reallotting FY 2017 funds awarded to the State Council on Developmental Disabilities (SCDD) and the Protection & Advocacy (P&A) agency located within the Commonwealth of Puerto Rico. This determination is based on the limited reported expenditures and requests for reimbursement over the last several years from the SCDD and P&A in the Commonwealth of Puerto Rico.

The Puerto Rico SCDD will have up to $1.9 million rescinded and proportionately redistributed to the remaining SCDDs. SCDDs that receive FY 2017 realloted funds will have through the end of FY 2018 to obligate the funds and until the end of FY 2019 to liquidate the funds.

The Puerto Rico P&A will have up to $550,000 rescinded and proportionately redistributed to the remaining P&As. P&As that receive the FY 2017 funds will have through the end of FY 2019 to spend the funds.

Realloted funds for both the SCDDs and the P&As must be used according to

the terms as outlined in the FY 2017 Notice of Award for each program.

DATES: Funds will be realloted after September 1, 2017 and before September 30, 2017.

ADDRESSES: The reallotment amounts to SCDDs and P&As can be found at *https://www.acl.gov/node/110.*

FOR FURTHER INFORMATION CONTACT: Andrew Morris, Office of Policy and Development, Center on Policy & Evaluation, Administration for Community Living, 330 C St. SW., Washington, DC 20201. Telephone (202) 795–7408. Email *andrew.morris@acl.hhs.gov.* Please note the telephone number is not toll free. This document will be made available in alternative formats upon request. Written correspondence can be sent to Administration for Community Living, U.S. Department of Health and Human Services, 330 C St. SW., Washington, DC 20201.

Dated: August 4, 2017.

Melissa Ortiz,

Commissioner, Administration on Disabilities.

[FR Doc. 2017–17195 Filed 8–14–17; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–1197]

The Food and Drug Administration's Proposed Method for Adjusting Data on Antimicrobials Sold or Distributed for Use in Food-Producing Animals Using a Biomass Denominator; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability for public comment of a proposed method for applying a food animal biomass denominator to annual data on antimicrobials sold and distributed for use in food animals in the United States. This method will allow us to obtain a corrected estimate of antimicrobial drug sales relative to the animal population potentially being treated with those drugs, thereby lending further context to the antimicrobial sales data we are collecting and analyzing.

DATES: Submit either electronic or written comments on the proposed method by November 13, 2017.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before November 13, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of November 13, 2017. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2017–N–1197 for ''FDA's Proposed Method for Adjusting Data on Antimicrobials Sold or Distributed for Use in Food-Producing Animals Using a Biomass Denominator.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Submit written requests for single copies of the proposed method to the Policy and Regulations Staff (HFV–6), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your requests. Persons with access to the Internet may obtain the proposed method at either *https://www.fda.gov/ForIndustry/UserFees/AnimalDrugUserFeeAct*