Without pulse surveys, it will be difficult to measure changes employee sentiment on critical PMA issues in a short time frame. Having three pulses allows us to trend questions as conditions change.

Technical experts will review and approve the survey content.

Some questions may be asked that are of a personal or sensitive nature (*e.g.,* questions on Diversity, Equity, Inclusion, and Accessibility).

B. Annual Reporting Burden

Respondents: 367,000.
Responses Per Respondent: 1.
Total Annual Responses: 3.
Hours per Response: 0.0106 (38 seconds).
Total Burden Hours: 11,621.67.

C. Public Comments

GSA invites comments on: whether this collection of information is necessary, whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the Regulatory Secretariat Division by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. ''3090–XXXX Generic Clearance for the Collection of the Government-wide Pulse Survey'' in all correspondence.

Beth Anne Killoran,

Deputy Chief Information Officer.

[FR Doc. 2022–13988 Filed 6–29–22; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Sexual Risk Avoidance Education Performance Analysis Study—Extension (Office of Management and Budget (OMB) #0970–0536)

AGENCY: Office of Planning, Research, and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS).

ACTION: Request for public comment.

SUMMARY: OPRE and the Family and Youth Services Bureau (FYSB) request an extension without changes to a currently approved information collection activity as part of Sexual Risk Avoidance Education Performance Analysis Study (SRAE PAS)(OMB Control No. 0970–0536; expiration date October 31, 2022). The goal of the study is to collect, analyze, and report on performance measures data for the SRAE program.

DATES: *Comments are due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act (PRA) of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing *opreinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The purpose of the SRAE program is to educate youth on how to voluntarily refrain from nonmarital sexual activity and prevent other youth risk behaviors. The requested extension will allow ACF to continue to collect the performance measures from SRAE grantees. Data will continue to be used to determine if the SRAE grantees are meeting performance benchmarks related to their program's mission and priorities. The program office will continue to use the data to provide technical assistance to grantees and for its own reporting purposes.

Respondents: Departmental Sexual Risk Avoidance Education (DSRAE), State Sexual Risk Avoidance Education (SSRAE), and Competitive Sexual Risk Avoidance Education (CSRAE) grantees, their sub recipients, and program participants.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
(1) Participant Entry Survey					
DSRAE participants	378,390	1	0.1333	50,439	16,813
SSRAE participants	952,899	1	0.1333	127,021	42,340
CSRAE participants	60,408	1	0.1333	8,052	2,684
(2) Participant Exit Survey					
DSRAE participants	302,712	1	0.1667	50,462	16,821
SSRAE participants	762,319	1	0.1667	127,079	42,360
CSRAE participants	48,326	1	0.1667	8,056	2,685
(3) Performance reporting data entry form: grantees					
DSRAE grantees	119	6	16	11,424	3,808
SSRAE grantees	39	6	16	3,744	1,248
CSRAE grantees	34	6	16	3,264	1,088
(4) Performance reporting data entry form: subrecipients					
DSRAE subrecipients	252	6	13	19,656	6,552

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
SSRAE subrecipients	426	6	13	33,228	11,076
CSRAE subrecipients	63	6	13	4,914	1,638

Estimated Total Annual Burden Hours: 149,113.

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 42 U.S.C. 1310.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–14026 Filed 6–29–22; 8:45 am]

BILLING CODE 4184–83–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–E–2251]

Determination of Regulatory Review Period for Purposes of Patent Extension; TEPEZZA

AGENCY: Food and Drug Administration, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or the Agency) has determined the regulatory review period for TEPEZZA and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of a patent which claims that human biological product.

DATES: Anyone with knowledge that any of the dates as published (see **SUPPLEMENTARY INFORMATION**) are incorrect may submit either electronic or written comments and ask for a redetermination by August 29, 2022. Furthermore, any interested person may petition FDA for a determination regarding whether the applicant for extension acted with due diligence during the regulatory review period by December 27, 2022. See ''Petitions'' in the **SUPPLEMENTARY INFORMATION** section for more information.

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 August 29, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of August 29, 2022. 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–2020–E–2251 for ''Determination of Regulatory Review Period for Purposes of Patent Extension; TEPEZZA.'' 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, 240–402–7500.

- 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