tolerance. Additionally, a physical examination and fingerstick blood tests will be used for health screening. The purpose of collecting data in the environmental chamber is to compare physiologic and cognitive measurements at different core body temperatures to evaluate factors contributing to individual variability in cognitive and physiologic responses to heat and to evaluate whether core body temperature thresholds exist above which cognitive deficits are observed.

The total estimated burden hours are 103. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (hours)
Miners.				
	Ingestion of temperature pill	30	2	1/60
	Fitting of chest strap	30	2	1/60
	Consent form (field)	30	1	30/60
	Health screening questionnaire (field)	30	1	30/60
	Heat stress app—shift questionnaire (field)	30	4	1/60
	PVT cognitive test (field)	30	5	5/60
	Heat stress app—post—shift questionnaire (field)	30	2	10/60
	Ingestion of temperature pill	15	3	1/60
	Fitting of chest strap	15	3	1/60
	Consent form (chamber)	15	1	30/60
	Physical examination	15	1	10/60
	Health screening questionnaire (chamber)	15	1	30/60
Miners/ firefighters/ construction workers.				
	Fingerstick blood sample for point-of-care testing	8	1	1/60
	Release of Information (HIPPA)	3	1	1/60
	Borg and thermal scale	15	5	1/60
	PANAS KSS fatigue	15	5	2/60
	Cognitive test: PVT (chamber)	15	5	10/60
	Cognitive test: N-back	15	5	1/60
	Pre-test screening questionnaire (chamber)	15	2	5/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2020–18275 Filed 8–19–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Administration for Native Americans (ANA) Ongoing Progress Report (OPR) and Objective Work Plan (OWP)

AGENCY: Administration for Native Americans, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families' (ACF) Administration for Native Americans (ANA) is requesting a revision to the information collection: Ongoing Progress Report (OPR) and the Objective Work Plan (OWP) (OMB #0970–0452). Changes are proposed to reduce the burden on the public by combining ANA's Annual Data Report (OMB #0970–0475) with the OPR.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: Content changes are being made to the currently approved OPR. ANA will continue to use the currently approved OPR with minimal changes to the instructions for the remainder of fiscal year (FY) 2020 and will use the modified OPR beginning FY 2021. The modified OPR combines ANA's Annual Data Report (OMB #0970–0475) with the OPR.

The information in the OPR is collected on a semi-annual basis to monitor the performance of grantees and better gauge grantee progress. The semi-annual data collection replaces the previous quarterly filing requirement of the OPR.

The OPR information collection is conducted in accordance with Sec. 811 [42 U.S.C. 2992] of the Native American Programs Act and will allow ANA to report quantifiable results across all program areas. It also provides grantees with parameters for reporting their progress and helps ANA better monitor and determine the effectiveness of their projects.

There are no changes proposed to the OWP. The OWP information collection is conducted in accordance with 42 U.S.C. of the Native American Programs Act of 1972, as amended. This collection is necessary to evaluate applications for financial assistance and determine the relative merits of the projects for which such assistance is

requested, as set forth in Sec. 806 [42 U.S.C. 2991d–1](a)(1).

Respondents: Federally and state-recognized tribes, Native Pacific Islanders, Tribal Colleges and Universities, native non-profits, and consortia.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Objective Work Plan	300	1	3	900	300
Ongoing Progress Report FY 2020	200	2	1	400	133
Ongoing Progress Report FY 2021—Exp. Date	200	4	2	1600	533

* Burden is annualized over the three year approval period.

Estimated Total Annual Burden Hours: 966.

Authority: Sec. 806 [42 U.S.C. 2991d–1](a)(1) and Sec. 811 [42 U.S.C. 2992].

John M. Sweet Jr.,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–18219 Filed 8–19–20; 8:45 am]

BILLING CODE 4184–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1429]

Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Human Drug Compounding Outsourcing Facilities under Section 503B of the Federal Food, Drug, and Cosmetic Act and Associated Fees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection pertaining to the registration of human drug compounding outsourcing facilities under the Federal Food, Drug, and Cosmetic Act (FD&C Act) and associated fees.

DATES: Submit either electronic or written comments on the collection of information by October 19, 2020.

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 October 19, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of October 19, 2020. 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–2013–N–1429 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Human Drug Compounding Outsourcing Facilities Under Section 503B of the FD&C Act and Associated Fees.'' 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