

Improving Head Start for School Readiness Act of 2007 and the Consolidated Appropriations Act of 2017.

Mary B. Jones,
ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Native Employment Works (NEW) Program Plan Guidance and Report Requirements, (OMB No.: 0970-0174)

AGENCY: Division of Tribal TANF Management, Office of Family Assistance, Administration for Children and Families, HHS.

ACTION: Request for Public Comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a three-year extension of the form OFA-0086, NEW Plan Guidance and NEW Program Report (OMB #0970-0174, expiration 7/31/2019). There are changes requested to these forms, including the deletion of guidance for NEW programs included in Public Law 102-477 programs.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing *infocollection@acf.hhs.gov*. Alternatively, copies can

also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The NEW program plan guidance documents specify the information needed to complete a NEW program plan and explains the process for plan submission every third year and to complete the annual program report. The program plan is the application for NEW program funding and documents how the grantee will carry out its NEW program. The program report provides HHS, Congress, and grantees information to document and assess the activities and accomplishments of the NEW program.

Respondents: Indian tribes and tribal coalitions that run NEW programs.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
NEW program plan guidance for non-477 Tribes	1 15	15	1	29	435
NEW program report	2 44	44	1	15	660

¹ We estimate that 44 of the 78 NEW grantees will not include their NEW programs in Public Law 102-477 projects. 44 grantees divided by 3 (because grantees submit the NEW plan once every 3 years) = 15.

² We estimate that 44 of the 78 NEW grantees will not include their NEW programs in Public Law 102-477 projects and therefore will submit the NEW program report to HHS.

Estimated Total Annual Burden Hours: 1095.

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. 612.

Mary B. Jones,
ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Continued Information Collection Activity; Evaluation of the Child Welfare Capacity Building Collaborative (OMB Number: 0970-0484)

AGENCY: Children's Bureau, Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a three-year extension of the previously approved forms that include satisfaction surveys; a leadership interview protocol; a web-based collaboration survey; assessment tools; and service-specific feedback forms (OMB #0970-0484, expiration 8/31/2019). There are no changes to the forms.

DATES: *Comments due within 60 days of publication.* In compliance with the

requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing *infocollection@acf.hhs.gov*. Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The Evaluation of the Child Welfare Capacity Building Collaborative is sponsored by the Children's Bureau, Administration for Children and Families of the U.S. Department of Health and Human Services. The Capacity Building Collaborative includes three centers (Center for States, Center for Tribes,

Center for Courts) funded by the Children’s Bureau to provide national child welfare expertise and evidence-informed training and technical assistance services to state, tribal and territorial public child welfare agencies and Court Improvement Programs (CIPs). The Centers offer a wide array of services including, but not limited to: Web-based content and resources, product development and dissemination, self-directed and group-based training, virtual learning and peer networking events, and tailored consultation and coaching. During the project period, Center services are evaluated by both Center-specific evaluations and a Cross-Center Evaluation. The Center-specific evaluations are designed to collect data on Center-specific processes and outcomes, which are used to support service delivery and continuous quality improvement. The Cross-Center Evaluation is designed to respond to a set of cross-cutting evaluation questions posed by the Children’s Bureau. The Cross-Center Evaluation examines: How

and to what extent key partners across and within Centers collaborate; whether Center capacity building service interventions are evaluable; the degree to which Centers follow common protocols; what service interventions are delivered and in what services do jurisdictions participate; how satisfied recipients are with services; what outcomes are achieved in jurisdictions receiving Center services and under what conditions are services effective; and what are the costs of services.

The Cross-Center Evaluation uses a longitudinal, mixed methods approach to evaluate Center services as they develop and mature over the course of the study. Multiple data collection strategies are used to efficiently capture quantitative and qualitative data to enable analyses that address each evaluation question. Cross-Center Evaluation data sources for this effort include (1) satisfaction surveys to assess recipient satisfaction with services, such as the Learning Experiences Satisfaction Survey; (2) a leadership interview used to assess perceptions of state child

welfare directors, tribal child welfare directors, and CIP directors; and (3) a web-based collaboration survey used to assess perceptions of collaboration within and between the capacity building centers. Center-specific data sources for this effort include (1) assessment tools such as the Center for Tribes Needs and Fit Exploration Tools; and (2) service-specific feedback forms, such as the Center for States Intensive Projects instrument and the Center for Courts CQI Workshops instrument.

Respondents: Respondents of data collection instruments include (1) child welfare and judicial professionals who use the Collaborative’s products and online courses, that participate in webinars, virtual or in-person trainings, or peer events, and that receive brief or intensive tailored services from the Centers; (2) all State child welfare directors, and Tribal child welfare directors, and CIP coordinators that receive services from the Centers; and (3) directors and staff of the three Capacity Building Centers.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Web Pages and Products Satisfaction Survey	6,240	2,080	1	.08	166
Learning Experiences Satisfaction Survey (single) ¹	2,000	666	1	.33	220
Learning Experiences Satisfaction Survey (intensive) ²	3,600	1200	1	.08	96
Webinars, Events, and In-Person Meetings Satisfaction Survey	22,008	7,336	1	.08	587
Center for States Information and Referral Survey	48	16	1	.05	1
Center for States Intensive Projects Survey	1,320	440	2	.33	290
Center for States Constituency Groups Surveys	1,600	533	2	.33	352
Center for States Brief Tailored Services Survey	500	166	1	.33	55
Center for Tribes Contact Form	200	22	1	.05	1
Center for Tribes Demographic Survey	80	26	1	1.75	46
Center for Tribes Needs and Fit Exploration Tool Phase 1	120	40	1	1.5	60
Center for Tribes Needs and Fit Exploration Tool Phase 2	100	33	1	3.0	99
CIP Annual Meeting Survey	800	266	1	.13	35
Center for Courts CQI Workshops Survey	192	63	1	.17	11
Assessment and Capacity Building Work Plan Satisfaction Survey	1,800	600	1	.066	40
Leadership Interview—States and Territories	52	17	2	1	34
Leadership Interview—CIPs	52	17	2	1	34
Leadership Interview—Tribes	32	10	2	1.25	25
Leadership Interview Part II—Tribes	32	10	2	.67	13
Annual Collaboration Survey	920	306	1	.36	110
Total					2,275

¹ For Learning Experiences that consist of a single event (e.g., on-line session or in-person training).

² For more intensive Learning Experiences that require administration of multiple surveys over a series of events, modules, or units.

Estimated Total Annual Burden Hours: 2,275.

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. 5106.

Mary B. Jones,
ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2009-N-0232]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Interstate Shellfish Dealer’s Certificate

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by May 24, 2019.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, Fax: 202-395-7285, or emailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910-0021. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-3794, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Interstate Shellfish Dealer’s Certificate OMB Control Number 0910-0021—Revision

Under section 243 of the Public Health Service Act (42 U.S.C. 243) FDA is required to cooperate with and aid State and local authorities in the enforcement of their health regulations, and is authorized to assist States in the prevention and suppression of communicable diseases. Under this authority, we participate with State regulatory agencies, some foreign nations, and the molluscan shellfish industry in the National Shellfish Sanitation Program (NSSP). NSSP is a voluntary, cooperative program to promote the safety of molluscan shellfish by providing for the classification and patrol of shellfish growing waters and for the inspection and certification of shellfish processors.

Each NSSP-participating State and foreign nation monitors its molluscan shellfish processors and for purposes of interstate or international commerce issues certificates for those that meet the State or foreign shellfish control authority’s criteria. Each participating State and nation provides a certificate of its certified shellfish processors to FDA on Form FDA 3038, “Interstate Shellfish Dealer’s Certificate.” We use this information to publish the “Interstate Certified Shellfish Shippers List,” a monthly comprehensive listing of all molluscan shellfish processors certified under the cooperative program. If we did not collect the information necessary to compile this list, participating States would not be able to identify and keep out shellfish processed by uncertified processors in other States and foreign nations. Consequently, NSSP would not be able to control the distribution of uncertified and possibly unsafe shellfish in interstate and international commerce, and its effectiveness would be nullified.

In the **Federal Register** of March 9, 2018 (83 FR 10487), we published a notice seeking comment on a proposed determination that the European Union’s (EU’s) system of food safety control measures for raw bivalve molluscan shellfish intended for export into the United States, as adopted and implemented in Spain and the Netherlands, provides at least the same

level of sanitary protection as the United States equivalent. If finalized, such a determination would permit the importation of shellfish harvested from certain European production areas and processed by European establishments that have been listed by FDA on the Interstate Certified Shellfish Shippers List.

The March 9, 2018, notice also described the European Commission’s (EC’s) determination that the United States’ system is equivalent to its own, and as a result of that determination, its stated intent to accept shellfish from certain growing areas in the United States. On November 6, 2018, the EC published Commission Implementing Decision (EU) 2018/1668 which added the United States (MA and WA only) to the list of Third Countries from which molluscan shellfish imports are permitted. Shellfish harvested from growing areas with an Approved classification in those states are eligible for export to the EU.

As part of the equivalence determination, the EC identified the need for FDA to provide documentation collected from NSSP-participating shellfish control authorities seeking recognition under the EC’s equivalence determination. This documentation includes:

- A list of growing areas with an Approved classification,
- The most recent sanitary survey for each growing area with an Approved classification, and
- The most recent inspection report for each firm seeking to export shellfish to the EU.

For NSSP-Participants that do not produce live/raw shellfish required documentation is limited to the most recent Plant and Shipping Element Program Evaluation Report and the most recent inspection report for each shellfish processing firm to be listed for export to the EU.

In the **Federal Register** of June 8, 2018 (83 FR 26699), we published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

Activity	FDA form No.	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Submission of Interstate Shellfish Dealer’s Certificate.	3038	40	57	2,280	0.10 (6 minutes)	228