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
Adult Interview Guide	45	15	1	1.5	23
Adolescent Interview Guide	20	7	1	.875	6
Child Interview Guide	30	10	1	.50	5
Phone Screener for Prospective Families	120	40	1	.50	20

Estimated Total Annual Burden Hours: 54.

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, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@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.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–15830 Filed 7–24–18; 8:45 am]

BILLING CODE 4184–09–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living; Notice of Federal Review of the Puerto Rico State Council on Developmental Disabilities (SCDD) and the Protection and Advocacy System (P&A)

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: Representatives of the Administration on Intellectual and Developmental Disabilities (AIDD), Administration for Community Living (ACL), will be conducting a federal review of the Puerto Rico State Council on Developmental Disabilities (SCDD) and the Protection and Advocacy System (P&A) on September 17–21, 2018.

AIDD is soliciting comments from interested parties on your experiences with the work, program, and strategies employed by P&A and SDCC in meeting the needs of individuals with developmental disabilities and their families in Puerto Rico. You are encouraged to share your experiences by way of any of the following methods:

Email: Clare.huerta@acl.hhs.gov.

Telephone: 202–795–7301.

Mail Comments To: Clare Huerta, Program Specialist, Administration on Intellectual and Developmental Disabilities, Administration for Community Living, 330 C Street SW, 1st Floor, Washington, DC 20201.

Comments should be received by September 10, 2018 in order to be included in the final report.

FOR FURTHER INFORMATION CONTACT: Clare Barnett Huerta, Administration for Community Living, Administration on Intellectual and Developmental Disabilities, Office of Program Support, 330 C Street SW, 1st Floor, Washington, DC 20201, 202–795–7301.

Dated: July 12, 2018.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2018–15905 Filed 7–24–18; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; New Data Collection; National Center on Law and Elder Rights (NCLER)

AGENCY: Administration for Community Living (ACL), HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to ACL's National Center on Law and Elder Rights.

DATES: Submit written comments on the collection of information by August 24, 2018.

ADDRESSES: Submit written comments on the collection of information by:

(a) Email to: *OIRA_submission@omb.eop.gov,* Attn: OMB Desk Officer for ACL;

(b) fax to 202.395.5806, Attn: OMB Desk Officer for ACL; or

(c) by mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

FOR FURTHER INFORMATION CONTACT: Omar Valverde at *omar.valverde@acl.hhs.gov* or 202–795–7460.

SUPPLEMENTARY INFORMATION: 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 proposed collection of information represents new information requested from aging/disability networks to fulfill requirements regarding the provision of services and overall performance of ACL legal assistance programs.

ACL contracts with a national legal assistance resource center, the National Center on Law and Elder Rights, to provide the required services. Through the contract, ACL provides aging, disability, and related legal professionals with training, case consultations and technical assistance for demonstration projects regarding contractually identified priority legal topics.

The purpose of the information requested is for ACL to ensure that the resource center creates and prioritizes the training, case consultations and technical assistance resources it was contracted to provide and to ensure that the center targets the contractually designated aging network practitioners about priority subject matters. This approach enables ACL to make data-

informed decisions about the deployment of its resource center assets. These data are necessary for ACL to evaluate contractual compliance with established performance indicators. These metrics include quantifiable increases in uptake by stakeholders of training, case consultation and technical assistance, and measures of satisfaction with and perceived benefit from these services. For example, the metrics measure successful problem resolution as a result of the services provided and quantifiable data on fulfillment of requests for training, technical assistance, and consultation related to the contractually designated legal and systems development topic areas.

The information requested by ACL from legal and aging/disability professionals falls into the following areas: (1) Requests for training, case consultation, and technical assistance through an online, secure Uniform Resource Support Request Tool; (2) general requests for Legal Training (including the volume of Webinar registrations); (3) Case Consultation and Technical Assistance; and (4) information about satisfaction and use of the services and support received in order to enable ACL to measure performance outcomes.

Comments in Response to the 60-Day Federal Register Notice

As required by 5 CFR 1320.8(d), a 60-day notice was published in the **Federal Register** on December 5, 2017 (Volume 82, Number 232, pp. 57458–57460). One email was received expressing support for the data collection as proposed. No modifications were made to the proposed data collection elements and associated data collection instruments.

Estimated Annualized Burden Hours

The total estimated burden is 460.78 hours per year for individuals requesting and/or receiving resource support through NCLER. This figure is based on ACL field testing of 8 providers working within aging/disability/legal networks who measured the time required to fully submit information by answering the required questions using standardized forms:

Respondent/data collection activity	Number of respondents	Minutes per response	Annual burden hours
Resource Support Requests	80	1 min 54 sec	2.53
Legal Training, Case Consultation, Technical Assistance Requests	14,000	1 min 42 sec	397
Outcome Measurement	3,500	1 min 3 sec	61.25
Total	17,580	4 min 39 sec	460.78

Dated: July 12, 2018.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2018–15906 Filed 7–24–18; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–2544]

Use of Liquids and/or Soft Foods as Vehicles for Drug Administration: General Considerations for Selection and In Vitro Methods for Product Quality Assessments; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry entitled ''Use of Liquids and/or Soft Foods as Vehicles for Drug Administration: General Considerations for Selection and In Vitro Methods for Product Quality Assessments.'' This draft guidance applies to orally administered drug products and provides recommendations to sponsors who will use or recommend use of liquids and/or soft foods as vehicles for drug administration in investigational new drug applications (INDs), new drug applications (NDAs), Biologics License Applications (BLAs), as applicable, and in supplements to these applications.

DATES: Submit either electronic or written comments on the draft guidance by September 24, 2018 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

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–2018–N–2544 for ''Use of Liquids and/or Soft Foods as Vehicles for Drug Administration: General Considerations for Selection and In Vitro Methods for Product Quality Assessments.'' Received comments 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.