(19) Notice of Rights Handout and Notice of Rights and Provision of Services: Care providers are required to provide to all UC under the Flores v. Reno Settlement Agreement.

(20) Legal Service Provider List for UC: List of organizations who offer free legal representation and help for UC with State and Federal courts, immigration hearings, and appeals. Required under the Flores Settlement Agreement.

(21) URM Application: Certain populations of children and youth in ORR custody may become eligible for the Unaccompanied Refugee Minors Program, which is a State administered foster care program. In such instances the care provider facility or other interested party may complete this application form on behalf of the child.

(22) Withdrawal of Application or Declination of Placement Form: If a youth who has submitted an application for the URM Program wishes to withdraw this application, or if he or she has been offered placement and wishes to decline this placement, the youth must complete this form.

(23) Standard Shelter Tour Request: Used by members of the public and the media to submit to care providers in order to tour a shelter facility.

Respondents

UC in ORR care and custody (they are generally referred to ORR from the DHS) and who are then referred to ORR's Network of Care Providers.

Staff in ORR's Care Provider Network, including those in shelter care, secure and staff secure care, foster care, and residential treatment centers.

Approved sponsors of UC released from ORR care.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
UC Portal Capacity Report	50	1	.16/hour	8
Further Assessment Swift Track (FAST) Placement Tool	2,320	1	.25/hour	580
Placement Authorization Form	58,000	1	.1/hour	5,800
Notice of Placement in Secure or Staff Secure Facility	2,320	1	.1/hour	232
Initial Intakes Form	58,000	1	.25/hour	14,500
UC Assessment	58,000	1	.50/hour	29,000
Individual Service Plan	58,000	1	.25	14,500
UC Case Review Form	58,000	1	.50/hour	29,000
New Sponsor Form	55,200	1	.25/hour	13,800
Transfer Request and Tracking Form	1,000	1	.25/hour	250
Long Term Foster Care Placement Memo	279	1	.1/hour	28
Travel Request Form for UC Long Term Foster Care	20	1	.25/hour	5
Notice of Transfer to ICE Chief Counsel and Change of Address.	2,320	1	.1/hour	232
Care Provider Release Checklist	55,200	1	.1	5,520
Release Request	55,200	3	.25 hour	41,400
Discharge Notification	716	1	.25/hour	179
Verification of Release	55,200	1	.1/hour	5,520
Child Advocate Referral and Appointment Form	250	1	.50	125
Notice of Rights Handout and Notice of Rights and Provision of Services.	58,000	1	.1/hour	5,800
Legal Service Provider List for UC	58,000	1	.1	5,800
URM Application	350	1	1	350
Withdrawal of Application or Declination of Placement Form	10	1	.1/hour	1
Standard Shelter Tour Request	60	1	.1/hour	6

Estimated Total Annual Burden Hours: 172,636.

In compliance with the requirements of Section 506(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. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–23978 Filed 9–21–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AGENCY: Administration for Community Living, HHS.

ACTION: 60-Day notice of submission of information collection approval from the Office of Management and Budget and request for comments.

SUMMARY: As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service delivery, the Administration for Community Living proposes to submitted a Generic Information Collection Request (Generic ICR): ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery'' to OMB for approval under the Paperwork Reduction Act (PRA) (44 U.S.C. 3501 et. seq.).

DATES: Submit written or electronic comments on the collection of information by November 23, 2015.

ADDRESSES: Submit electronic comments on the collection of information to: Susan Jenkins at *Susan.Jenkins@aoa.hhs.gov*.

Submit written comments on the collection of information to Administration for Community Living, Washington, DC 20201, Attn. Susan Jenkins.

FOR FURTHER INFORMATION CONTACT: Susan Jenkins at 202.357.3591.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), *Title:* Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

Abstract: The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: the target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

Dated: September 17, 2015.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2015–24066 Filed 9–21–15; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0114]

Agency Information Collection Activities; Proposed Collection; Submission for Office of Management and Budget Review; Request for Samples and Protocols

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 October 22, 2015.

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 0910–0206. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *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.

Request for Samples and Protocols—OMB Control Number 0910–0206—Extension

Under section 351 of the Public Health Service Act (42 U.S.C. 262), FDA has the responsibility to issue regulations that prescribe standards designed to ensure the safety, purity, and potency of biological products and to ensure that the biologics licenses for such products are only issued when a product meets the prescribed standards. Under 21 CFR 610.2, the Center for Biologics Evaluation and Research (CBER) or the Center for Drugs Evaluation and Research may at any time require manufacturers of licensed biological products to submit to FDA samples of any lot along with the protocols showing the results of applicable tests prior to distributing the lot of the product. In addition to § 610.2, there are other regulations that require the submission of samples and protocols for specific licensed biological products: 21 CFR 660.6 (Antibody to Hepatitis B Surface Antigen); 21 CFR 660.36 (Reagent Red Blood Cells); and 21 CFR 660.46 (Hepatitis B Surface Antigen).

Section 660.6(a) provides requirements for the frequency of submission of samples from each lot of Antibody to Hepatitis B Surface Antigen product, and § 660.6(b) provides the requirements for the submission of a protocol containing specific information along with each required sample. For § 660.6 products subject to official release by FDA, one sample from each filling of each lot is required to be submitted along with a protocol consisting of a summary of the history of manufacture of the product, including all results of each test for which test results are requested by CBER. After official release is no longer required, one sample along with a protocol is required to be submitted at 90-day intervals. In addition, samples, which must be accompanied by a protocol, may at any time be required to