Respondents: Potential sponsors of UC.

Annual Burden Estimates:

RESPONDENTS

Instrument title	Annual total number of respondents	Annual total number of responses per respondent	Average burden hours per response	Annual total burden hours
Authorization for Release of Information (Forms FRP–2 & FRP–2s)	81,532	1	0.50	40,766
Family Reunification Application (Forms FRP–3 & FRP–3s)	122,950	1	1.00	122,950
Fingerprinting Instructions (Forms FRP–7 & FRP–7s)	81,532	1	1.25	101,915
Letter of Designation for Care of Minor (Forms FRP–9 & FRP–9s)	41,181	1	0.75	30,886
Estimated Annual Burden Total				296,517

RECORD KEEPERS

Instrument title	Annual total number of record keepers	Annual total number of responses per record keeper	Average burden hours per response	Annual total burden hours
Authorization for Release of Information (Forms FRP–2 & FRP–2s)	235	347	0.25	20,386
Family Reunification Application (Forms FRP–3 & FRP–3s)	235	523	0.25	30,726
Fingerprinting Instructions (Forms FRP–7 & FRP–7s)	235	347	1.00	81,545
Letter of Designation for Care of Minor (Forms FRP–9 & FRP–9s)	235	175	0.25	10,281
Estimated Annual Burden Hours Total				142,938

Authority: 6 U.S.C. 279; 8 U.S.C. 1232; *Flores* v. *Reno* Settlement Agreement, No. CV85–4544–RJK (C.D. Cal. 1996).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–13631 Filed 6–24–22; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Recordkeeping for New Vaccine and Mask Requirements To Mitigate the Spread of COVID–19 in Head Start (OMB #0970–0583)

AGENCY: Office of Head Start, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Head Start (OHS), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS) requests public comment on an extension with no changes to recordkeeping requirements for ACF Head Start grantees. An Interim Final Rule with Comment Period (IFC) was published on November 30, 2021 that established the COVID–19 vaccination requirements whereby all Head Start staff, certain contractors, and volunteers must be vaccinated for COVID–19 by January 31, 2022. OHS requested and received emergency approval from the Office of Management and Budget (OMB) to implement the associated recordkeeping requirements for 6 months. This request will extend approval beyond the current expiration date (6/30/2022). ACF is currently in the final rulemaking process. If the requirements in the final rule differ from the IFC in a way that alters recordkeeping requirements, ACF will make those changes in coordination with OMB.

DATES: *Comments due within 30 days of publication.* 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.

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. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: This request is for recipients of Head Start funding to continue to (1) collect and maintain records on the vaccination status of staff (including certain contractors) and volunteers in Head Start and Early Head Start programs and (2) develop and maintain a written COVID–19 testing protocol for individuals granted vaccine exemptions that was established through the IFC (86 FR 68052). There is no standard instrument required to be used to meet these recordkeeping requirements. Burden estimates have been updated to reflect more recent data available.

Respondents: Recipients of Head Start funding.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Responses per respondent	Average annual burden hours	Annual burden hours
Staff, Contractor, and Volunteer Reporting of New Vaccination	75,000	1	0.6667	50,002.5
Staff, Contractor, and Volunteer Reporting of Existing Vaccination	320,000	1	0.0833	26,656
Staff, Contactor, and Volunteer Requesting and Processing Vaccination Exemption	5,000	1	0.5000	2,500
Grant Recipient Maintaining Vaccination Records	1,573	1	6.3573	10,000
Grant Recipient Establishing COVID–19 Testing Protocol	1,573	1	3.3333	5,243.3
Grant Recipient Maintaining COVID–19 Testing Protocol	1,573	1	1	1,573

Estimated Total Annual Burden Hours: We estimate the one-time and ongoing burden to maintain records on staff and volunteer vaccination rates and establish and maintain a written COVID–19 testing protocol will result in 95,974.8 total annual burden hours.

(Authority: IFC [86 FR 68052])

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–13615 Filed 6–24–22; 8:45 am]

BILLING CODE 4184–40–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0977]

Agency Information Collection Activities; Proposed Collection; Comment Request; Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco To Protect Children and Adolescents

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) 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 the collection of information entitled, ''Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents.''

DATES: Submit either electronic or written comments on the collection of information by August 26, 2022.

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 26, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of August 26, 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–2012–N–0977 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents.'' 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 available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed