the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

The National YRBS Test-Retest Reliability Study—New—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of this request is to obtain OMB approval to conduct the National YRBS Test-Retest Reliability Study to establish the reliability of the National Youth Risk Behavior Survey (''YRBS'') questionnaire.

The YRBS assesses priority health risk behaviors related to the major preventable causes of mortality, morbidity, and social problems among both youth and young adults in the United States. Data on health risk behaviors of adolescents are the focus of approximately 65 national health objectives in Healthy People 2030, an initiative of the U.S. Department of Health and Human Services (HHS). The YRBS provides data to measure 13 of the proposed health objectives and one of the Leading Health Indicators currently under public comment to establish Healthy People 2030 objectives. In addition, the YRBS can identify racial and ethnic disparities in health risk behaviors. No other national source of data measures as many of the Healthy People 2030 objectives addressing adolescent health risk behaviors as the YRBS. The data also will have significant implications for policy and program development for school health programs nationwide. CDC seeks a one-year approval to conduct the National YRBS Test-Retest Reliability Study.

Between September and December of 2021, a sample of 2,000 students from 20 regular public secondary schools in the U.S. containing at least one of grades 9–12 will be selected in no more than 20 districts. This sample is expected to yield at least 1,000 participating students who completed both a Time 1 and Time 2 YRBS questionnaire. The table below reports the number of respondents annualized over the one-year project period. There are no costs to respondents except their time. The total estimated annualized burden hours are 1,540.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
District Administrators	District recruitment script	20	1	30/60
School Principals	School recruitment script	20	1	30/60
Classroom Teachers	Consent form checklist	80	1	15/60
Students	YRBS Questionnaire	1,000	2	45/60

Jeffrey M. Zirger,

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

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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Follow-up Study of Coaching Practices in Early Care and Education Settings (OMB #0970–0515)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: This is a primary data collection request for the Follow-up Study of Coaching Practices in Early Care and Education Settings (3), a follow-up to the previously approved Study of Coaching Practices in Early Care and Education Settings (SCOPE) survey (OMB #0970–0515). The study aims to examine, using surveys and qualitative interviews, the practice and processes of coaching and professional development in supporting early care and education (ECE) settings in their provision of care for preschool children and their families as COVID–19 has progressed. The study will focus on both centers and family child care (FCC) homes that serve low-income children, with a primary target of settings that serve children supported by Child Care and Development Fund subsidies or a Head Start grant.

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 *OPREinfocollection@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: Follow-up SCOPE will examine the practice of coaching and professional development more broadly provided in support of centers and FCC homes. The study will collect information on the following: How coaching and professional development are supporting centers and FCC homes; the perceived value and role of coaching, professional development, and quality improvement; the features of coaching and how they are delivered; and the role(s) of coaches and how they have been supported. The study will also examine the degree to which coaching has been sustained and/or changed compared to before COVID–19. In particular, there will be a focus on understanding the use of remote versus in-person strategies for coaching and professional development. This study aims to explore the implementation of coaching and professional development in ECE settings as COVID–19 has progressed. The study will not allow for statistical generalization to different sites or service populations.

Survey and interview questions will focus on the current status of these activities at the time of the data

collection, changes compared to before COVID–19 began, and what has been challenging or worked well. The study will use surveys and interviews with center directors, FCC providers, and coaches. The sample frame will be comprised of respondents to the 2019 survey.

Respondents: ECE center directors, coaches, and FCC providers who responded to 2019 SCOPE surveys.

Annual Burden Estimates

Data collection will be completed within a 1-year period.

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average Burden per response (in hours)	Total/Annual burden (in hours)
Coach Survey (Instrument 1)	100	1	.33	33
Center Director Survey (Instrument 2)	66	1	.33	22
FCC Provider Survey (Instrument 3)	38	1	.33	13
Coach Interview (Instrument 4)	12	1	.75	9
Center Director Interview (Instrument 5)	24	1	.75	18
FCC Provider Interview (Instrument 6): FCC providers	12	1	.75	9

Estimated Total Annual Burden Hours: 104.

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. 9858(a)(5), 42 U.S.C. 9835, and 42 U.S.C. 9844.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–28043 Filed 12–18–20; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0161]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Export of Food and Drug Administration-Regulated Products: Export Certificates

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) 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: Submit written comments (including recommendations) on the collection of information by January 20, 2021.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 0910–0498. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *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.

Export of Food and Drug Administration-Regulated Products: Export Certificates

OMB Control Number 0910–0498—Extension

Sections 801(e) and 802 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 381(e) and 382) pertain to the export of FDA-regulated products and are intended to ease restrictions on exportation. The provisions also require the Agency to issue written export certifications within 20 days of any request. In January 2011, section 801(e)(4)(A) was amended by the FDA Food Safety Modernization Act (Pub. L. 111–353) to provide authorization for export certification for food and animal feed, as well as certain unapproved products. To offset Agency resource expenditures for processing certifications requests, the statute provides that FDA may charge firms a fee not to exceed $175.

There are four FDA forms (Form FDA 3613, 3613a, 3613b, and 3613c) related to exporting FDA-regulated products. A description of each form is provided below. To obtain a fillable PDF file of each form, visit *https://www.fda.gov/vaccines-blood-biologics/exporting-cber-regulated-products/fda-forms-certificates-exporting.* To learn more about how to complete these forms, visit *https://www.fda.gov/vaccines-blood-biologics/exporting-cber-regulated-products/how-complete-fda-export-certificate-forms.*