Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee (CLIAC)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting for the Clinical Laboratory Improvement Advisory Committee (CLIAC). This meeting is open to the public, limited only by the space available. The meeting room accommodates approximately 100 people. The public is also welcome to view the meeting by webcast. Check the CLIAC website on the day of the meeting for the webcast link *www.cdc.gov/cliac.*

DATES: The meeting will be held on April 16, 2020, 8:30 a.m. to 5:00 p.m., EDT and April 17, 2020, 8:30 a.m. to 11:30 a.m., EDT.

ADDRESSES: Food and Drug Administration (FDA), White Oak Campus, 10903 New Hampshire Avenue, Building 31, Great Room, Silver Spring, Maryland 20993 and via webcast at *www.cdc.gov/cliac.*

FOR FURTHER INFORMATION CONTACT: Nancy Anderson, MMSc, MT(ASCP), Senior Advisor for Clinical Laboratories, Division of Laboratory Systems, Center for Surveillance, Epidemiology and Laboratory Services, Office of Public Health Scientific Services, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop V24–3, Atlanta, Georgia 30329–4018, telephone (404) 498–2741; *NAnderson@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services (HHS); the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention; the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare and Medicaid Services (CMS). The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory medicine practice and specific questions related to possible revision of the Clinical Laboratory Improvement Amendment (CLIA) standards. Examples include providing guidance on studies designed to improve safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods, the electronic transmission of laboratory information, and mechanisms to improve the integration of public health and clinical laboratory practices.

All people attending the CLIAC meeting in-person are required to register for the meeting online at least five business days in advance for U.S. citizens and at least 10 business days in advance for international registrants. Register at: *www.cdc.gov/cliac.* Register by scrolling down and clicking the ''Register for this Meeting'' button and completing all forms according to the instructions given. Please complete all the required fields before submitting your registration and submit no later than April 8, 2020, for U.S. registrants and April 1, 2020, for international registrants.

It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments on agenda items. Public comment periods for each agenda item are scheduled immediately prior to the Committee discussion period for that item. In general, each individual or group requesting to make oral comments will be limited to a total time of five minutes (unless otherwise indicated). To assure adequate time is scheduled for public comments, speakers should notify the contact person below at least 5 business days prior to the meeting date. For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, it is requested that comments be submitted at least 5 business days prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments should be provided to the contact person at the mailing or email address below, and will be included in the meeting's Summary Report.

The CLIAC meeting materials will be made available to the Committee and the public in electronic format (PDF) on the internet instead of by printed copy. Check the CLIAC website on the day of the meeting for materials: *www.cdc.gov/cliac.*

Matters to be Considered: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and discussions will focus on an update on CLIAC recommendations; an update on the Genetic Testing Reference Materials Coordination Program (GeT–RM); an update of the December 2019 CDC's Board of Scientific Counselors, Deputy Director for Infectious Diseases meeting; a report from the Office of the National Coordinator for Health Information Technology (ONC) Health Information Technology Advisory Committee; the laboratory response to the COVID–19 coronavirus disease outbreak; and technological advances in digital imaging. Agenda items are subject to change as priorities dictate.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–20–0493; Docket No. CDC–2020–0015]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the

general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled 2021 and 2023 National Youth Risk Behavior Surveys (YRBS). CDC is requesting a three-year approval to reinstate, with changes, the data collection for the national YRBS, a biennially school-based survey of high school students in the United States.

DATES: CDC must receive written comments on or before April 28, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0015 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments through the Federal eRulemaking portal (*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

2021 and 2023 National Youth Risk Behavior Surveys (OMB Contrtol No. 0920–0493)—Reinstatement with change—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 reinstate with change, the data collection for the National Youth Risk Behavior Survey (YRBS), a school-based survey that has been conducted biennially since 1991. OMB approval for the 2017 YRBS and 2019 YRBS expired September 30, 2019 (OMB Control No. 0920–0493). CDC seeks a three-year approval to conduct the YRBS in Spring 2021 and Spring 2023. Minor changes incorporated into this reinstatement request include: An updated title for the information collection to accurately reflect the years in which the survey will be conducted, minor changes to the data collection instrument, and the use of a tablet-based data collection methodology starting in 2023.

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.

In Spring 2021 and Spring 2023, the YRBS will be conducted among nationally representative samples of students attending public and private schools in grades 9–12. Information supporting the YRBS also will be collected from state-, district-, and school-level administrators and teachers. The table below reports the number of respondents annualized over the three-year project period. There are no costs to respondents except their time. The total estimated annualized burden hours are 6,259.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
State Administrators	State-level Recruitment Script for the Youth Risk Behavior Survey.	17	1	30/60	9
District Administrators	District-level Recruitment Script for the Youth Risk Behavior Survey.	80	1	30/60	40
School Administrators	District-level Recruitment Script for the Youth Risk Behavior Survey.	133	1	30/60	67

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Teachers	Data Collection Checklist for the Youth Risk Behavior Survey.	440	1	15/60	110
Students	Youth Risk Behavior Survey	8,045	1	45/60	6,034
Total					6,259

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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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Personal Responsibility Education Program (PREP) Performance Measures and Adulthood Preparation Subjects (PMAPS) Studies—Data Collection Related to the Performance Measures Study—Extension (OMB #0970–0497).

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

ACTION: Request for public comment.

SUMMARY: A goal of the Performance Measures and Adulthood Preparation Subjects (PMAPS) studies is to collect, analyze, and report on performance measure data for the Personal Responsibility Education Program (PREP) programs. The Office of Planning, Research, and Evaluation (OPRE) and the Family and Youth Services Bureau (FYSB) in the Administration for Children and Families (ACF) request a revision to a currently approved information collection (OMB No. 0970–0497; expiration date: 04/30/2020). The purpose of the request is to make adaptions to the participant entry and exit surveys, and continue the ongoing data collection of the performance measures from PREP grantees.

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: The PMAPS studies consist of two components: The Performance Measures Study and the Adulthood Preparation Subjects Study. The data collection for the Adulthood Preparation Subjects Study is complete. This notice is specific to a request for an extension of data collection activities for the Performance Measures Study only. The Performance Measures Study component includes collection and analysis of performance measure data from State PREP (SPREP), Tribal PREP (TPREP), Competitive PREP (CPREP), and Personal Responsibility Education Innovative Strategies (PREIS) grantees. Data will be used to determine if PREP and PREIS grantees are meeting performance benchmarks related to the program's mission and priorities.

This request includes the development of adapted participant entry and exit surveys for middle school students (6th, 7th, and 8th grade youth) that exclude the most sensitive questions pertaining to sexual behavior. This is because some of the PREP middle school curricula do not include topics on sexual behavior, *i.e.,* focus only on healthy relationship education. The adapted surveys will be used by all grantees that serve middle school youth. In addition, some minor edits have been made to the high school surveys.

Respondents: Performance measurement data collection instruments will be administered to individuals representing SPREP, TPREP, CPREP, and PREIS grantees, their sub-awardees, and program participants.

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
PREP Participant Entry Survey	319,673	106,558	1	0.15	15,984
PREP Participant Exit Survey	291,624	97,208	1	0.13333	12,961
Performance Reporting System Data Form—State grantees	153	51	2	18	1,836
Performance Reporting System Data Form—TPREP grantees	28	9	2	18	324
Performance Reporting System Data Form—CPREP grantees	75	25	2	14	700
Performance Reporting System Data Form—PREIS grantees	38	13	2	14	364
Performance Reporting System Data Form—State subawardees	987	329	2	14	9,212