online and all active EAS/Comprizon contracts were migrated into the new EASi application. A complete snapshot of the EAS/Comprizon data was taken and stored in the Business Intelligence (BI) database. The snapshot still exists in the BI database today and is used for querying and reporting purposes. None of the EAS/Comprizon records currently in the BI database contain PII. The replacement system for EAS, EASi, also does not contain PII. The vendor information is directly collected and stored in SAM.

SYSTEM NAME AND NUMBER:

GSA/PBS–6 Electronic Acquisition System (EAS).

HISTORY:

73 FR 22389.

Richard Speidel,

Chief Privacy Officer, Office of the Deputy Chief Information Officer, General Services Administration.

[FR Doc. 2020–24077 Filed 10–29–20; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–21–1277; Docket No. CDC–2020–0109]

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 the existing information collection project titled The Childcare Survey of Activity and Wellness (C–SAW) Pilot Study. The pilot study will determine the current practices and policies of early care and education (ECE) providers in four states around nutrition, physical activity, and wellness and will inform the development of a potential national surveillance system.

DATES: CDC must receive written comments on or before December 29, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0109 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

The Childcare Survey of Activity and Wellness (C–SAW) Pilot Study—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) works to promote optimal nutrition, physical activity, and wellness in early care and education (ECE) facilities for children 0–5 years of age. Consistent with this mission, and with clear evidence that ECE facilities can impact the habits and preferences of young children, CDC obtained OMB approval (OMB Control No. 0920–1277, Expiration Date 12/31/2020) to conduct a pilot survey to better understand ECE center practices related to nutrition, physical activity, and wellness. CDC was unable to collect the information as planned due to closures of ECE centers because of the COVID–19 pandemic. CDC is requesting an extension of the collection for two years due to the COVID–19 pandemic and its impact on information collection and to collect 10 additional COVID–19 related questions on the survey to understand the impact of COVID–19 on our topic areas of child care center status, nutrition, physical activity and wellness The additional questions are expected to minimally affect burden. These critical data are used to effectively inform state and national programs.

Data collected from this pilot survey will be used to understand the current practices of ECE centers in a representative sample in four states. This initial C–SAW will establish baseline measures of the prevalence of specific practices related to nutrition, physical activity, and wellness in a standard way across states. This baseline will also allow CDC and state partners to better understand ECE center needs and provide opportunities for collaboration and areas for improvement at the state and national levels. Second, the survey will be used to inform the development of a potential national

surveillance system enabling states and CDC to track changes over time and obtain data to guide the planning, implementation, and evaluation of national and state obesity prevention efforts.

A sample of approximately 1,200 ECE centers across four states will be recruited to participate in this one-time data collection effort. Each center will receive a recruitment letter introducing the survey, explaining its objectives and the importance of their participation, and instructions for completing the survey. It is anticipated that most responses will be submitted online via the internet. However, paper surveys will be available upon request. Approximately two weeks after the initial recruitment letter is mailed, all sampled centers will receive a reminder postcard. Approximately four weeks after the initial recruitment letter is mailed, non-respondents will be sent another letter along with a hardcopy of the questionnaire.

Participation in this study is completely voluntary and there are no costs to the respondent other than their time. The estimated annualized burden hours is 409. The approval request is for two years.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
ECE Director or Administrator	Recruitment Letter	1,140	1	5/60	95
ECE Director or Administrator	Web/Mail Survey	627	1	30/60	314
Total					409

Jeffrey M. Zirger,

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

[FR Doc. 2020–24230 Filed 10–28–20; 4:15 pm]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Decision To Evaluate a Petition To Designate a Class of Employees From the Pinellas Plant in Clearwater, Florida, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: NIOSH gives notice of a decision to evaluate a petition to designate a class of employees from the Pinellas Plant in Clearwater, Florida, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000.

FOR FURTHER INFORMATION CONTACT: Grady Calhoun, Director, Division of Compensation Analysis and Support, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, MS C–46, Cincinnati, OH 45226–1938, Telephone 513–533–6800. Information requests can also be submitted by email to *DCAS@CDC.GOV.*

SUPPLEMENTARY INFORMATION: Pursuant to 42 CFR 83.12, the initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: Pinellas Plant.

Location: Clearwater, Florida.

Job Titles and/or Job Duties: All employees who worked in any areas of Pinellas Plant.

Period of Employment: May 19, 1957 through December 31, 1997.

Authority: 42 CFR 83.9–83.12.

John J. Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. 2020–24123 Filed 10–29–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–381]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by December 29, 2020.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.