patients, and VTE prophylaxis monitoring and support will be used to assess the extent to which hospitals apply these components of the framework for HA–VTE prevention. The responses to specific VTE prevention practices can be used to assess VTE prevention practices by hospital characteristics (e.g., bed size, urban vs. rural location, teaching vs. non-teaching status) to better target efforts or interventions to improve HA–VTE prevention. Information collected on the barriers to establishing a hospital-wide VTE prevention policy will be helpful in addressing these challenges. Information will be collected on both adult general medical and surgical units since VTE prevention practices differ by specialty. Information on VTE risk assessment (e.g., who conducts the assessment, when is it performed, mandatory or optional, format, type of RAM) will improve understanding of real-world hospital VTE risk assessment practices. Information on the capacity of hospitals to collect data on VTE risk assessment will be helpful in determining the feasibility of VTE risk assessment as a VTE prevention performance measure. The data collected can also serve as a baseline for evaluation of future HA–VTE prevention initiatives.

### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>The Director of Patient Safety and Quality, the Chairperson of the Patient Safety Committee, other quality improvement professional</td>
<td>Recruitment material: Implementation evaluation and project information sheet.</td>
<td>384</td>
<td>1</td>
<td>15/60</td>
<td>96</td>
</tr>
<tr>
<td>The Director of Patient Safety and Quality, the Chairperson of the Patient Safety Committee, other quality improvement professional</td>
<td>Evaluation of Venous Thromboembolism Prevention Practices in U.S. Hospitals Questionnaire.</td>
<td>384</td>
<td>1</td>
<td>1</td>
<td>384</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>480</td>
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</table>

Jeffrey M. Zirger, Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention. [FR Doc. 2020–25574 Filed 11–18–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–21–0879]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled Information Collections to Advance State, Tribal, Local, and Territorial (STLT) Governmental Agency and System Performance, Capacity, and Program Delivery to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on 05/21/2020 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. 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. Direct written comments and/or suggestions regarding 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**

Information Collections to Advance State, Tribal, Local, and Territorial (STLT) Governmental Agency and System Performance, Capacity, and Program Delivery (OMB Control No. 0920–0879, Exp. 1/31/2021)—Extension—Center for State, Tribal, Local and Territorial Support (CSTLTS), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The mission of the Department of Health and Human Services is to enhance the health and well-being of all Americans. As part of HHS, CDC conducts critical science and provides health information to people and communities to save lives and protect people from health threats. To this end, CDC and HHS seek to accomplish their mission by collaborating with partners throughout the nation and the world to
monitor health, detect and investigate health problems, conduct research to enhance prevention, develop and advocate sound public health policies, implement prevention strategies, promote healthy behaviors, foster safe and healthful environments, and provide leadership and training.

CDC is requesting a three-year approval to extend a generic clearance to collect information related to domestic public health issues and services that affect and/or involve state, tribal, local and territorial (STLT) government entities.

The respondent universe is comprised of STLT governmental staff or delegates acting on behalf of a STLT agency involved in the provision of essential public health services in the United States. Delegate is defined as a governmental or non-governmental agent (agency, function, office or individual) acting for a principal or submitted by another to represent or act on their behalf. The STLT agency is represented by a STLT entity or delegate with a task to protect and/or improve the public’s health.

Information will be used to assess situational awareness of current public health emergencies; make decisions that affect planning, response and recovery activities of subsequent emergencies; fill CDC and HHS gaps in knowledge of programs and/or STLT governments that will strengthen surveillance, epidemiology, and laboratory science; and improve CDC’s support and technical assistance to jurisdictions. CDC and HHS will conduct brief data collections, across a range of public health topics related to essential public health services.

CDC estimates up to 30 data collections with State, territorial, or tribal governmental staff or delegates, and 10 data collections with local/county/city governmental staff or delegates will be conducted on an annual basis. Ninety-five percent of these data collections will be web-based and five percent telephone, in-person, and focus groups. The total annualized burden of 54,000 hours is based on the following estimates.

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per respondent (in hrs.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>State, Territorial, or Tribal government staff or delegate.</td>
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<td>30</td>
<td>1</td>
</tr>
<tr>
<td>Local/County/City government staff or delegate.</td>
<td>Web, telephone, in-person, focus group ..........</td>
<td>3,000</td>
<td>10</td>
<td>1</td>
</tr>
</tbody>
</table>

**SUPPLEMENTARY INFORMATION:** The Privacy Act of 1974, as amended (5 U.S.C. 552a) provides certain protections for individuals applying for and receiving federal benefits. The law governs the use of computer matching by federal agencies when records in a system of records (meaning, federal agency records about individuals retrieved by name or other personal identifier) are matched with records of other federal or non-federal agencies. The Privacy Act requires agencies involved in a matching program to:

1. Enter into a written agreement, which must be prepared in accordance with the Privacy Act, approved by the Data Integrity Board of each source and recipient federal agency, provided to Congress and the Office of Management and Budget (OMB), and made available to the public, as required by 5 U.S.C. 552a(o), (u)(3)(A), and (u)(4).

2. Notify the individuals whose information will be used in the matching program that the information they provide is subject to verification through matching, as required by 5 U.S.C. 552a(o)(1)(D).

3. Verify match findings before suspending, terminating, reducing, or making a final denial of an individual’s benefits or payments or taking other adverse action against the individual, as required by 5 U.S.C. 552a(p).

4. Report the matching program to Congress and the OMB, in advance and

Jeffrey M. Zirger,

[FR Doc. 2020–25573 Filed 11–18–20; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Medicare & Medicaid Services

Privacy Act of 1974; Matching Program

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

ACTION: Notice of new matching program.

SUMMARY: In accordance with subsection (e)(12) of the Privacy Act of 1974, as amended, the Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS) is providing notice of the re-establishment of a matching program between CMS and the Social Security Administration (SSA), “Determining Enrollment or Eligibility for Insurance Affordability Programs Under the Patient Protection and Affordable Care Act.”

DATES: The deadline for comments on this notice is December 21, 2020. The re-established matching program will commence not sooner than 30 days after publication of this notice, provided no comments are received that warrant a change to this notice. The matching program will be conducted for an initial term of 18 months (from approximately March 9, 2021 to September 8, 2022) and within three months of expiration may be renewed for one additional year if the parties make no change to the matching program and certify that the program has been conducted in compliance with the matching agreement.

ADDRESSES: Interested parties may submit comments on the new matching program to the CMS Privacy Officer by mail at: Division of Security, Privacy Policy & Governance, Information Security & Privacy Group, Office of Information Technology, Centers for Medicare & Medicaid Services, Location: N1–14–56, 7500 Security Blvd., Baltimore, MD 21244–1850, or walter.stone@cms.hhs.gov.

FOR FURTHER INFORMATION CONTACT: If you have questions about the matching program, you may contact Anne Pesto, Senior Advisor, Marketplace Eligibility and Enrollment Group, Center for Consumer Information and Insurance Oversight, Centers for Medicare & Medicaid Services, at 410–786–3492, by email at anne.pesto@cms.hhs.gov, or by mail at 7500 Security Blvd., Baltimore, MD 21244.