DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention (CDC)
Requirements and Registration for Healthcare Associated Venous Thromboembolism Prevention Challenge; Amendment of Notice

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).
AWARD APPROVING OFFICIAL: Thomas R. Frieden, MD, MPH, Director, Centers for Disease Control and Prevention, and Administrator, Agency for Toxic Substances and Disease Registry.
ACTION: Notice.
SUMMARY: The Centers for Disease Control and Prevention (CDC) located within the Department of Health and Human Services (HHS) announces an amendment to its notice entitled, Announcement of Requirements and Registration for Healthcare Associated Venous Thromboembolism Prevention Challenge. This amendment is being made to reflect an increase in the number of Champions and change the maximum total prize disbursement. There are no other changes to the September 22, 2015 notice.
FOR FURTHER INFORMATION CONTACT: Michele Beckman, Division of Blood Disorders, National Center on Birth Defects and Developmental Disabilities, Centers for Disease Control and Prevention, 1600 Clifton Road NE., Mailstop E–64, Atlanta, GA 30329.
SUPPLEMENTARY INFORMATION: Subject of Challenge Competition: On September 22, 2015 CDC announced the Requirements and Registration for Healthcare Associated Venous Thromboembolism Prevention Challenge (80 FR 57187). This notice announces an increase in the number of Champions, from 7 to 8. The Champions were selected from the highest scoring U.S. hospitals, multi-hospital systems, hospital networks, and managed care organizations. Champions were recognized as HA–VTE Prevention Champions and will receive a cash award of $10,000. A maximum of $80,000 will now be awarded in this challenge, an increase of $10,000. Additional honorable mention awards were also made to deserving entries. Federal and international winners received non-monetary recognition but no prize.
Dated: June 7, 2016.
Sandra Cashman.
Executive Secretary, Centers for Disease Control and Prevention.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (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.

The estimated annual burden hours are 466. There are no costs to respondents other than their time.
## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services


### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid 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 the burden estimates or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by August 9, 2016.

**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.

   To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:


   2. **Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.**

   3. **Call the Reports Clearance Office at (410) 786–1326.**

**FOR FURTHER INFORMATION CONTACT:** Reports Clearance Office at (410) 786–1326.

### SUPPLEMENTARY INFORMATION:

**Contents**

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection’s supporting statement and associated materials (see ADDRESSES).


Under the 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. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

1. **Type of Information Collection Request:** Extension of a currently approved collection; Title of Information Collection: Examination and Treatment for Emergency Medical Conditions and Women in Labor; Use: In accordance with to regulation sections 488.18, 489.20 and 489.24, during Medicare surveys of hospitals and State agencies CMS will review hospital records for lists of on-call physicians, and will review and obtain the information which must be recorded on hospital medical records for individuals with emergency medical conditions and women in labor, and the emergency department reporting information Medicare participating hospitals and Medicare State survey agencies must pass on to CMS. Additionally, CMS will use the QIO Report assessing whether an individual had an emergency condition and whether the individual was stabilized to determine whether to impose a CMP or physician exclusion sanctions. Without such information, CMS will be unable to make the hospital emergency services compliance determinations that Congress expects CMS to make under sections 1154, 1866 and 1867 of the Act. Form Number: CMS–R–142 (OMB control number: 0938–0667); Frequency: Occasionally; Affected Public: Private Sector; Number of Respondents: 6,149; Total Annual Responses: 6,149; Total Annual Hours: 1. (For policy questions regarding this collection contact Renate Dombrowski at 410–786–4645.)

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
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<td>Screener Script Guide</td>
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<td>1</td>
<td>10/60</td>
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<tr>
<td>Child participants aged 6–12 years</td>
<td>Seat Belt Fit Measurements</td>
<td>142</td>
<td>1</td>
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**ESTIMATED ANNUALIZED BURDEN HOURS**