received. As required by OMB regulations, 5 CFR part 1320, the FTC is providing this second opportunity for public comment.

**Estimated Annual Hours Burden:** 450 hours (300 testing-related hours; 150 disclosure-related hours).

**Likely Respondents and Estimated Burden:**

- **(a) Testing—High fidelity manufacturers—**300 new products/year × 1 hour each = 300 hours; and
- **(b) Disclosures—High fidelity manufacturers—**[(300 new products/year × 1 specification sheet) + (300 new products/year × 1 brochure)] × 15 minutes each = 150 hours.

**Frequency of Response:** Periodic.

**Estimated Annual Labor Cost:** $23,463 per year ($14,967 for testing + $8,496 for disclosures).

**Request for Comment**

You can file a comment online or on paper. For the FTC to consider your comment, we must receive it on or before January 29, 2018. Write “Amplifier Rule: FTC File No. P974222” on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission website, at http://www.ftc.gov/os/publiccomments.shtm.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online, or to send them to the Commission by courier or overnight service. To make sure that the Commission considers your online comment, you must file it at https://ftcpublic.commentworks.com/ftc/amplifierrule2p2 by following the instructions on the web-based form. When this Notice appears at http://www.regulations.gov, you also may file a comment through that website.

If you file your comment on paper, write “Amplifier Rule: FTC File No. P974222” on your comment and on the envelope, and mail it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610, Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service. Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW, Washington, DC 20503. Comments sent to OMB by U.S. postal mail are subject to delays due to heightened security precautions. Thus, comments can also be sent via email to wiberante@omb.eop.gov.

Because your comment will be placed on the publicly accessible FTC website at https://www.ftc.gov, you are solely responsible for making sure that your comment does not include any sensitive or confidential information. In particular, your comment should not include any sensitive personal information, such as your or anyone else’s Social Security number; date of birth; driver’s license number or other state identification number, or foreign country equivalent; passport number; financial account number; or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, your comment should not include any “trade secret or any commercial or financial information which, . . . is privileged or confidential”—as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled “Confidential,” and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c). Your comment will be kept confidential only if the General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted on the public FTC website—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from the FTC website. Submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before January 29, 2018. For information on the Commission’s privacy policy, including routine uses permitted by the Privacy Act, see https://www.ftc.gov/site-information/privacy-policy.

David C. Shonka,
Acting General Counsel.

[FR Doc. 2017–28064 Filed 12–27–17; 8:45 am]

**BILLING CODE 6750–01–P**

**DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION**

[OMB Control No. 9000–0001; Docket No. 2017–0053; Sequence 14]

**Submission for OMB Review; Standard Form 28, Affidavit of Individual Surety**

**AGENCY:** Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

**ACTION:** Notice.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act of 1995, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning the Standard Form (SF) 28, Affidavit of Individual Surety.

**DATES:** Submit comments on or before January 29, 2018.

**ADDRESSES:** Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: Office of Information and Regulatory Affairs of OMB, Attention: Desk Officer for GSA, Room 10236, NEOB, Washington, DC 20503. Additionally submit a copy to GSA by any of the following methods:

- Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by searching the OMB control number 9000–0001. Select the link that corresponds with “Information Collection 9000–0001, SF 28, Affidavit..."
of Individual Surety”. Follow the instructions provided on the screen. Please include your name, company name (if any), and “Information Collection 9000–0001, SF 28, Affidavit of Individual Surety”, on your attached document.


Instruction: Please submit comments only and cite Information Collection 9000–0001, SF 28, Affidavit of Individual Surety, in all correspondence related to this collection. All comments received will be posted without change to http://www.regulations.gov, including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Ms. Zenaida Delgado, Procurement Analyst, Federal Acquisition Policy Division, GSA, 202–960–7207 or zenaaida.deldago@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

The Standard Form (SF) 28, Affidavit of Individual Surety, is used by all executive agencies, including the Department of Defense, to obtain information from individuals wishing to serve as sureties to Government bonds. Offerors and contractors may use an individual surety as security for bonds required under a solicitation/contract for supplies or services (including construction). It is an elective decision on the part of the offeror/contractor to use individual sureties instead of other available sources of surety or sureties for Government bonds. The information on the SF 28 is used to assist the contracting officer in determining the acceptability of individuals proposed as sureties.

B. Public Comment

A 60 day notice was published in the Federal Register at 82 FR 48231, on October 17, 2017. No comments were received.

C. Annual Reporting Burden

The number of solicitations and contracts requiring the submission of bid guarantees, performance, or payment bonds, correlate roughly to the number of contract awards containing FAR clause 52.228–11, Pledge of Assets. Fiscal year 2016 data on the number of contracts containing FAR clause 52.228–11 was obtained from the Electronic Document Access system (DoD official contract file system) to estimate burdens for this information collection notice. The following is a summary of the FY 2016 data:

Respondents: 244.

Responses Per Respondent: 2.

Total Responses: 488.

Hours Per Response: 0.3.

Total Burden Hours: 146.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0001, SF 28, Affidavit of Individual Surety, in all correspondence.


Lorin S. Curit,
Director, Federal Acquisition Policy Division, Office of Government-wide Acquisition Policy, Office of Government-wide Policy.

[FPR Doc. 2017–28025 Filed 12–27–17; 8:45 am
BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control, (BSC, NCIPC)

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 Board of Scientific Counselors, National Center for Injury Prevention and Control, (BSC, NCIPC). This meeting is open to the public limited only by the 75 telephone ports available. There will be a public comment period at the end of the meeting day from 3:30 p.m.–3:45 p.m., February 26, 2018.

DATES: The meeting will be held on February 26, 2018, 01:00 p.m.–04:00 p.m., EST.


FOR FURTHER INFORMATION CONTACT: Gwendolyn H. Cattledge, Ph.D., M.S.E.H., Deputy Associate Director for Science, NCIPC, CDC, 4770 Buford Highway NE, Mailstop F–63, Atlanta, GA 30341, Telephone (770) 488–1430, Email address: GCattledge@cdc.gov.

SUPPLEMENTARY INFORMATION:

Purpose

The Board will: (1) Conduct, encourage, cooperate with, and assist other appropriate public health authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes, diagnosis, treatment, control, and prevention of physical and mental diseases, and other impairments; (2) assist States and their political subdivisions in preventing and suppressing communicable and non-communicable diseases and other preventable conditions and in promoting health and well-being; and (3) conduct and assist in research and control activities related to injury. The Board of Scientific Counselors makes recommendations regarding policies, strategies, objectives, and priorities; and reviews progress toward injury prevention goals and provides evidence in injury prevention-related research and programs. The Board also provides advice on the appropriate balance of intramural and extramural research, the structure, progress and performance of intramural programs. The Board is designed to provide guidance on extramural scientific program matters, including the: (1) Review of extramural research concepts for funding opportunity announcements; (2) conduct of Secondary Peer Review of extramural research grants, cooperative agreements, and contracts applications received in response to the funding opportunity announcements as it relates to the Center’s programmatic balance and mission; (3) submission of secondary review recommendations to the Center Director of applications to be considered for funding support; (4) review of research program strategies, objectives, and priorities; and (5) review of program proposals.

Matters to be Considered: The agenda will include updates from the Center and Policy Directors, as well as discussions on use of social media in suicide prevention. Agenda items are subject to change as priorities dictate.

The Director, Management Analysis and Services Office, 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.

Claudette Grant, Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–28070 Filed 12–27–17; 8:45 am]
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