

(2) Provide documentation of completion of such privacy training to the contracting officer if requested.

The contracting officer will use the information in contract administration and to establish that all applicable contractor and subcontractor employees comply with the privacy training requirements.

C. Annual Burden

Respondents/Recordkeepers: 4,282.

Total Annual Responses: 25,799.

Total Burden Hours: 12,873. (27 reporting hours + 12,846 recordkeeping hours)

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division by calling 202-501-4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 9000-0182, Privacy Training.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

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OFFICE OF MANAGEMENT AND BUDGET

Office of Federal Procurement Policy

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0001; Docket No. 2026-0166; Sequence No. 1]

Information Collection; Certain Federal Acquisition Regulation Part 28 Requirements

AGENCY: Office of Federal Procurement Policy (OFPP), Office of Management and Budget (OMB); Department of Defense (DOD); General Services Administration (GSA); and National Aeronautics and Space Administration (NASA).

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995 and OMB regulations, OFPP, DoD, GSA, and NASA invite the public to comment on an extension concerning certain Federal Acquisition Regulation (FAR) part 28 requirements. OFPP, DoD, GSA, and

NASA invite comments on: whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through January 31, 2027. OFPP, DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: OFPP, DoD, GSA, and NASA will consider all comments received by July 21, 2026.

ADDRESSES: OFPP, DoD, GSA, and NASA invite interested persons to submit comments on this collection through <https://www.regulations.gov> and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202-501-4755 or GSARegSec@gsa.gov.

Instructions: All items submitted must cite OMB Control No. 9000-0001, Certain Federal Acquisition Regulation Part 28 Requirements. Comments received generally will be posted without change to <https://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check www.regulations.gov, approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: FARPolicy@gsa.gov or call 202-969-4075.

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

OMB Control No. 9000-0001, Certain Federal Acquisition Regulation Part 28 Requirements, Standard Forms (SF) 24, 25, 25-A, 25-B, 28, 34, 35, 273, 274, 275, 1414, 1415, 1416, and 1418.

B. Need and Uses

This clearance covers the information that offerors or contractors must submit to comply with the following FAR requirements:

- FAR 52.228-1, Bid Guarantee. This provision (or clause) requires offerors or contractors to furnish a bid guarantee in the proper form and amount when a performance bond or a performance and payment bond is also required. (SF 24, Bid Bond; SF 34, Annual Bid Bond).

- FAR 52.228-2, Additional Bond Security. This clause requires contractors to furnish additional bond security under certain circumstances. This clause is used both for construction and other than construction contracts. (SF 1414 Consent of Surety and SF 1415, Consent of Surety and Increase of Penalty).

- FAR 52.228-13, Alternative Payment Protections. This clause requires contractors to submit one of the payment protections listed in the clause by the contracting officer, in construction contracts greater than \$35,000 but not exceeding \$150,000.

- FAR 52.228-14, Irrevocable Letter of Credit. This clause requires offerors or contractors to provide certain information when they intend to use an irrevocable letter of credit (ILC) in lieu of a required bid bond, or to secure other types of required bonds such as performance and payment bonds. This clause is required in solicitations and contracts when a bid guarantee, or performance bond, or performance and payment bonds are required.

- FAR 52.228-15, Performance and Payment Bonds-Construction. This clause requires contractors to provide performance and payment bonds in construction contracts exceeding \$150,000 (SF 25, Performance Bond; SF 25-A, Payment Bond; SF 25-B, Continuation Sheet (for SF's 24, 25, and 25-A); SF 273, Reinsurance Agreement for a Bonds Statute Performance Bond; SF 274, Reinsurance Agreement for a Bonds Statute Payment Bond).

- FAR 52.228-16, Performance and Payment Bonds-Other Than Construction. This clause requires contractors to furnish performance and payment bonds for other than construction contracts exceeding the simplified acquisition threshold only in certain circumstances. (SF 35, Annual Performance Bond; SF 275, Reinsurance Agreement in Favor of the United States; SF 1416, Payment Bond for Other Than Construction Contracts; SF 1418, Performance Bond for Other Than Construction Contracts).

- Standard Form (SF) 28, Affidavit of Individual Surety. This form is used by all executive agencies, including DoD, 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 or contract for supplies or services (including construction). It is an elective decision on the part of the offeror or contractor to use individual sureties instead of other available sources of surety or sureties for Government bonds.

The Government retains the bid guarantees, bonds, or alternative payment protections until the contractor's obligation is fulfilled. The contracting officer uses the information on the SF 28 to determine the acceptability of individuals proposed as sureties.

C. Annual Burden

Respondents: 22,916.

Total Annual Responses: 22,926.

Total Burden Hours: 22,912.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by calling 202-501-4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 9000-0001, Certain Federal Acquisition Regulation Part 28 Requirements.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2007-D-0369]

Product-Specific Guidances; Draft and Revised Draft Guidances for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of additional draft and revised draft product-specific guidances. The draft guidances provide product-specific recommendations on, among other things, the design of bioequivalence (BE) studies to support abbreviated new drug applications (ANDAs). In the **Federal Register** of June 11, 2010, FDA announced the availability of a guidance for industry entitled "Bioequivalence Recommendations for Specific Products" that explained the process that would be used to make product-

specific guidances available to the public on FDA's website. The draft guidances identified in this notice were developed using the process described in that guidance.

DATES: Submit either electronic or written comments on these draft guidances by July 21, 2026 to ensure that the Agency considers your comment on these draft guidances before it begins work on the final version of these guidances.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- **Mail/Hand Delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2007-D-0369 for "Product-Specific Guidances; Draft and Revised Draft Guidances for Industry." Received comments will be placed in the docket

and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **Confidential Submissions—**To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240-402-7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10001 New Hampshire Ave., Hillandale Building, 4th Floor, Silver Spring, MD 20993-0002. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.