

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than November 16, 2017.

*A. Federal Reserve Bank of Richmond* (Adam M. Drimer, Assistant Vice President) 701 East Byrd Street, Richmond, Virginia 23261-4528. Comments can also be sent electronically to [Comments.applications@rich.frb.org](mailto:Comments.applications@rich.frb.org):

1. *Hamilton Bancorp, Inc.*; to become a bank holding company by acquiring 100 percent of the voting shares of Hamilton Bank, both of Towson, Maryland.

*B. Federal Reserve Bank of Atlanta* (Kathryn Haney, Director of Applications) 1000 Peachtree Street, NE., Atlanta, Georgia 30309. Comments can also be sent electronically to [Applications.Comments@atl.frb.org](mailto:Applications.Comments@atl.frb.org):

1. *Lakeside Bancshares, Inc.*; to become a bank holding company by acquiring 100 percent of the outstanding shares of Lakeside Bank, both of Lake Charles, Louisiana.

*C. Federal Reserve Bank of Dallas* (Robert L. Triplett III, Senior Vice President) 2200 North Pearl Street, Dallas, Texas 75201-2272:

1. *Henderson Citizens Bancshares, Inc., Henderson, Texas*; to acquire voting shares of Union State Bancshares, Inc., Florence, Texas, and therefore indirectly acquire shares of Union State Bank, Florence, Texas.

Board of Governors of the Federal Reserve System, October 18, 2017.

**Ann Misback,**

*Secretary of the Board.*

[FR Doc. 2017-22963 Filed 10-23-17; 8:45 am]

**BILLING CODE P**

## GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090-0163; Docket 2017-0001; Sequence 5]

### Submission for OMB Review; General Services Acquisition Regulation; Information Specific to a Contract or Contracting Action (Not Required by Regulation)

**AGENCY:** Office of Acquisition Policy, General Services Administration (GSA).

**ACTION:** Notice of request for public comments regarding an extension to an existing OMB information collection.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act, 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 regarding Information Specific to a Contract or Contracting Action (not required by regulation). A notice was published in the **Federal Register** at 82 FR 33911 on July 21, 2017. No comments were received.

**DATES:** Submit comments on or before: November 24, 2017.

**ADDRESSES:** Submit comments identified by Information Collection 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), 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 3090-0163. Select the link "Comment Now" that corresponds with "Information Collection 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation)". Follow the instructions provided on the screen. Please include your name, company name (if any), and "Information Collection 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation)," on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Mr. Poe/IC 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation).

**Instructions:** Please submit comments only and cite Information Collection 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), in all

correspondence related to this collection. Comments received generally will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check [www.regulations.gov](http://www.regulations.gov), approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Ms. Jennifer Calik, Procurement Analyst, GSA Acquisition Policy Division, at telephone 312-353-6090 or email [jennifer.calik@gsa.gov](mailto:jennifer.calik@gsa.gov).

### SUPPLEMENTARY INFORMATION:

#### A. Purpose

GSA has various mission responsibilities related to the acquisition and provision of supplies, transportation, information technology, telecommunications, real property management, and disposal of real and personal property. These mission responsibilities generate requirements that are realized through the solicitation and award of public contracts. In Fiscal Year 2016, these contracts had values ranging from under \$100 to over \$777,000,000, including the base and all options.

Most GSA procurement-related information collections are required by the Federal Acquisition Regulation (FAR) or General Services Administration Acquisition Regulation (GSAR); each clause requiring such a collection must be individually approved by OMB. However, some solicitations require contractors to submit information specific to that contracting action, such as information needed to evaluate offers (e.g. specific instructions for technical and price proposals, references for past performance) or data used to administer resulting contracts (e.g. project management plans).

This information collection is currently associated with GSA's information collection requirements contained in solicitations issued in accordance with the Uniform Contract Format under FAR Part 14, Sealed Bidding (see GSAR 514.201-1); FAR Part 15, Contracting by Negotiation (see GSAR 515.204-1); and solicitations under FAR Part 12, Acquisition of Commercial Items, when issued in accordance with the policy and procedures of FAR Part 14 and FAR Part 15 (see GSAR 512.301). This includes information collection requirements found in GSA Federal Supply Schedule (FSS) solicitations.

**B. Annual Reporting Burden**

*Respondents:* 347,239.

*Responses per Respondent:* 1.

*Total Responses:* 347,239.

*Hours per Response:* 40.

*Total Burden Hours:* 138,896.

**C. Public Comments**

*Public comments are particularly invited on:* Whether this collection of information is necessary, whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

**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, Washington, DC 20405, telephone 202-501-4755.

Please cite OMB Control No. 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), in all correspondence.

**Jeffrey A. Koses,**

*Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.*

[FR Doc. 2017-23027 Filed 10-23-17; 8:45 am]

**BILLING CODE 6820-61-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Food and Drug Administration**

[Docket No. FDA-2017-N-5436]

**Electronic Study Data Submission; Data Standards; Support for Version Update of World Health Organization Drug Global**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) is announcing support for the most current B3-format annual version of the World Health Organization (WHO) Drug Global (WHODG) (formerly named WHO Drug Dictionary) (available at <https://www.who-umc.org>), end of support for earlier versions of WHODG, and an update to the FDA Data Standards

Catalog (Catalog) for study data provided in new drug applications (NDAs), abbreviated new drug applications (ANDAs), biologics license applications (BLAs), and certain investigational new drug applications (INDs) to the Center for Biologics Evaluation and Research (CBER) and the Center for Drug Evaluation and Research (CDER).

**ADDRESSES:** You may submit either electronic or written comments 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-2017-N-5436 for "Electronic Study Data Submission; Data Standards; Support for Version Update of World Health Organization Drug Global." 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.

- **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.gpo.gov/fdsys/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.

**FOR FURTHER INFORMATION CONTACT:** Ron Fitzmartin, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 1115, Silver Spring, MD 20993-0002, 301-796-5333, email: [cderdatastandards@fda.hhs.gov](mailto:cderdatastandards@fda.hhs.gov); or Stephen Ripley, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 7301, Silver Spring, MD 20993-0002, 240-402-7911.

**SUPPLEMENTARY INFORMATION:**