documents regarding the agreements to the Secretary by email at Secretary@fmc.gov, or by mail, Federal Maritime Commission, Washington, DC 20573. Comments will be most helpful to the Commission if received within 12 days of the date this notice appears in the Federal Register. Copies of agreements are available through the Commission’s website (www.fmc.gov) or by contacting the Office of Agreements at (202)–523–5793 or tradeanalyzer@fmc.gov.

Agreement No.: 201357.

Agreement Name: Sallauln Lines/Liberty Global Logistics LLC Space Charter Agreement.

Parties: Sallauln Lines Switzerland SA and Liberty Global Logistics LLC.

Filing Party: Wayne Rohde; Cozen O’Connor.

Synopsis: The agreement authorizes the parties to charter space to/from one another on an “as needed/as available” basis in the trade between the U.S. Atlantic and Gulf Coasts on the one hand and ports in Ghana, Togo, Benin, Nigeria, Jordan, Saudi Arabia, the United Arab Emirates, Qatar, Kuwait, and Pakistan on the other hand.

Proposed Effective Date: 2/22/2021.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/39512.


Rachel E. Dickon, Secretary.

[FR Doc. 2021–04379 Filed 3–2–21; 8:45 am]

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**FEDERAL RESERVE SYSTEM**

**Agency Information Collection Activities: Announcement of Temporary Approval by the Board Under Delegated Authority and Submission to OMB**

**AGENCY:** Board of Governors of the Federal Reserve System.

**ACTION:** Temporary approval of information collection, request for comment.

**SUMMARY:** The Board of Governors of the Federal Reserve System (Board) has temporarily revised the Reporting and Disclosure Requirements Associated with Emergency Lending, pursuant to the authority delegated to the Board by the Office of Management and Budget (OMB).

**DATES:** Comments must be submitted on or before May 3, 2021.

**ADDRESSES:** You may submit comments, identified by FR A, by any of the following methods:

- **Email:** regs.comments@federalreserve.gov. Include the OMB number in the subject line of the message.
- **Fax:** (202) 452–3819 or (202) 452–3102.
- **Mail:** Ann E. Misback, Secretary, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551.

All public comments are available from the Board’s website at https://www.federalreserve.gov/apps/fioa/proposedregs.aspx as submitted, unless modified for technical reasons or to remove personally identifiable information at the commenter’s request. Accordingly, comments will not be edited to remove any identifying or contact information. Public comments may also be viewed electronically or in paper in Room 146, 1709 New York Avenue NW, Washington, DC 20006, between 9:00 a.m. and 5:00 p.m. on weekdays. For security reasons, the Board requires that visitors make an appointment to inspect comments. You may do so by calling (202) 452–3684. Upon arrival, visitors will be required to present valid government-issued photo identification and to submit to security screening in order to inspect and photocopy comments.

Additionally, commenters may send a copy of their comments to the Office of Management and Budget (OMB) Desk Officer—Shagufta Ahmed—Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Room 10235, 725 17th Street NW, Washington, DC 20503, or by fax to (202) 395–6974.

**FOR FURTHER INFORMATION CONTACT:** Federal Reserve Board Clearance Officer—Nuha Elmaghribi—Office of the Chief Data Officer, Board of Governors of the Federal Reserve System, Washington, DC 20551. (202) 452–3829.

**SUPPLEMENTARY INFORMATION:** On June 15, 1984, OMB delegated to the Board authority under the PRA to approve and assign OMB control numbers to collections of information conducted or sponsored by the Board. In exercising this delegated authority, the Board is directed to take every reasonable step to solicit comments. In determining whether to approve a collection of information, the Board will consider all comments received from the public and other agencies. Pursuant to its delegated authority, the Board may temporarily approve a revision to a collection of information, without providing opportunity for public comment, if the Board determines that a change in an existing collection must be instituted quickly and that public participation in the approval process would defeat the purpose of the collection or substantially interfere with the Board’s ability to perform its statutory obligation.

As discussed below, the Board has made certain temporary revisions to the FR A information collection. The Board’s delegated authority requires that the Board, after temporarily approving a collection, publish a notice soliciting public comment. Therefore, the Board is also inviting comment on a proposal to extend the FR A information collection for three years, with these revisions.

A copy of the Paperwork Reduction Act (PRA) OMB submission, including the reporting form and instructions, supporting statement, and other documentation will be available at https://www.reginfo.gov/public/do/PRAMain. These documents will also be made available on the Board’s public website at https://www.federalreserve.gov/apps/reportforms/review.aspx or may be requested from the agency clearance officer, whose name appears above.

**Request for Comment on Information Collection Proposal**

The Board invites public comment on the following information collection, which is being reviewed under authority delegated by the OMB under the PRA. Comments are invited on the following:

a. Whether the proposed collection of information is necessary for the proper performance of the Board’s functions, including whether the information has practical utility.

b. The accuracy of the Board’s estimate of the burden of the proposed information collection, including the validity of the methodology and assumptions used.

c. Ways to enhance the quality, utility, and clarity of the information to be collected.

d. Ways to minimize the burden of information collection on respondents, including through the use of automated collection techniques or other forms of information technology; and

e. Estimates of capital or startup costs and costs of operation, maintenance, and purchase of services to provide information.

At the end of the comment period, the comments and recommendations received will be analyzed to determine the extent to which the Board should modify the proposal.
Legal authorization and confidentiality: The FR A is authorized pursuant to section 13(3) of the Federal Reserve Act, which sets out requirements for emergency lending. The obligation to respond is required to obtain a benefit. The information collected under the FR A may be kept confidential under exemption 4 of the Freedom of Information Act, which protects commercial or financial information obtained from a person that is privileged or confidential.

Current actions: The Board is revising the FR A information collection to address information collection requirements related to borrowers under the Main Street Lending Program, who participate in the PPP. Participating borrowers seeking to reduce the calculation of "existing outstanding and undrawn available debt" for purposes of determining the maximum allowable loan amount under the Main Street Lending Program must provide its eligible lender either with the Small Business Administration form it has already completed and submitted to its PPP lender (which may be the same lender), or must complete and submit a Board form to its Main Street lender during the Main Street loan underwriting process, as applicable. The FR A respondent counts for all parts of the information collection are being revised to reflect updated estimates of lender participation in the Main Street Lending Program.

Detailed Discussion of Public Comments: On March 2, 2020, the Board published a notice in the Federal Register (85 FR 12295) requesting public comment for 60 days on the extension, without revision, of the FR A. One comment was received; it did not address aspects of the information collection as described in 5 CFR 1320.8(d). On May 15, 2020, following the temporary approval of a first set of revisions to the FR A, the Board published a Federal Register notice (85 FR 29447) requesting public comment for 60 days on those temporary revisions. On June 4, 2020, following the temporary approval of a second set of revisions to the FR A, the Board published a Federal Register notice (85 FR 34448) requesting public comment for 60 days on those temporary revisions. On August 21, 2020, following the temporary approval of a third set of revisions to the FR A, the Board published a Federal Register notice (85 FR 51715) requesting public comment for 60 days on those temporary revisions. Comments in response to all of those requests for comment are expected to be considered, along with any comments received in response to this request for comment.


Michele Taylor Fennell, Deputy Associate Secretary of the Board.

[FR Doc. 2021–04362 Filed 3–2–21; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0362]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice for Manufacturing, Processing, Packing, and Holding of Finished Pharmaceuticals, Including Medical Gases, and Active Pharmaceutical Ingredients

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the Federal Register concerning each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection associated with current good manufacturing practice (CGMP) for drugs, finished pharmaceuticals, including medical gases, and active pharmaceutical ingredients (APIs).

DATES: Submit either electronic or written comments on the collection of information by May 3, 2021.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before May 3, 2021. The https://www.regulations.gov electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of May 3, 2021. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Legal authorization and confidentiality: The FR A is authorized pursuant to section 13(3) of the Federal Reserve Act, which sets out requirements for emergency lending. The obligation to respond is required to obtain a benefit. The information collected under the FR A may be kept confidential under exemption 4 of the Freedom of Information Act, which protects commercial or financial information obtained from a person that is privileged or confidential.

Current actions: The Board is revising the FR A information collection to address information collection requirements related to borrowers under the Main Street Lending Program, who participate in the PPP. Participating borrowers seeking to reduce the calculation of "existing outstanding and undrawn available debt" for purposes of determining the maximum allowable loan amount under the Main Street Lending Program must provide its eligible lender either with the Small Business Administration form it has already completed and submitted to its PPP lender (which may be the same lender), or must complete and submit a Board form to its Main Street lender during the Main Street loan underwriting process, as applicable. The FR A respondent counts for all parts of the information collection are being revised to reflect updated estimates of lender participation in the Main Street Lending Program.

Detailed Discussion of Public Comments: On March 2, 2020, the Board published a notice in the Federal Register (85 FR 12295) requesting public comment for 60 days on the extension, without revision, of the FR A. One comment was received; it did not address aspects of the information collection as described in 5 CFR 1320.8(d). On May 15, 2020, following the temporary approval of a first set of revisions to the FR A, the Board published a Federal Register notice (85 FR 29447) requesting public comment for 60 days on those temporary revisions. On June 4, 2020, following the temporary approval of a second set of revisions to the FR A, the Board published a Federal Register notice (85 FR 34448) requesting public comment for 60 days on those temporary revisions. On August 21, 2020, following the temporary approval of a third set of revisions to the FR A, the Board published a Federal Register notice (85 FR 51715) requesting public comment for 60 days on those temporary revisions. Comments in response to all of those requests for comment are expected to be considered, along with any comments received in response to this request for comment.


Michele Taylor Fennell, Deputy Associate Secretary of the Board.

[FR Doc. 2021–04362 Filed 3–2–21; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0362]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice for Manufacturing, Processing, Packing, and Holding of Finished Pharmaceuticals, Including Medical Gases, and Active Pharmaceutical Ingredients

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the Federal Register concerning each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection associated with current good manufacturing practice (CGMP) for drugs, finished pharmaceuticals, including medical gases, and active pharmaceutical ingredients (APIs).

DATES: Submit either electronic or written comments on the collection of information by May 3, 2021.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before May 3, 2021. The https://www.regulations.gov electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of May 3, 2021. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Legal authorization and confidentiality: The FR A is authorized pursuant to section 13(3) of the Federal Reserve Act, which sets out requirements for emergency lending. The obligation to respond is required to obtain a benefit. The information collected under the FR A may be kept confidential under exemption 4 of the Freedom of Information Act, which protects commercial or financial information obtained from a person that is privileged or confidential.

Current actions: The Board is revising the FR A information collection to address information collection requirements related to borrowers under the Main Street Lending Program, who participate in the PPP. Participating borrowers seeking to reduce the calculation of "existing outstanding and undrawn available debt" for purposes of determining the maximum allowable loan amount under the Main Street Lending Program must provide its eligible lender either with the Small Business Administration form it has already completed and submitted to its PPP lender (which may be the same lender), or must complete and submit a Board form to its Main Street lender during the Main Street loan underwriting process, as applicable. The FR A respondent counts for all parts of the information collection are being revised to reflect updated estimates of lender participation in the Main Street Lending Program.

Detailed Discussion of Public Comments: On March 2, 2020, the Board published a notice in the Federal Register (85 FR 12295) requesting public comment for 60 days on the extension, without revision, of the FR A. One comment was received; it did not address aspects of the information collection as described in 5 CFR 1320.8(d). On May 15, 2020, following the temporary approval of a first set of revisions to the FR A, the Board published a Federal Register notice (85 FR 29447) requesting public comment for 60 days on those temporary revisions. On June 4, 2020, following the temporary approval of a second set of revisions to the FR A, the Board published a Federal Register notice (85 FR 34448) requesting public comment for 60 days on those temporary revisions. On August 21, 2020, following the temporary approval of a third set of revisions to the FR A, the Board published a Federal Register notice (85 FR 51715) requesting public comment for 60 days on those temporary revisions. Comments in response to all of those requests for comment are expected to be considered, along with any comments received in response to this request for comment.


Michele Taylor Fennell, Deputy Associate Secretary of the Board.

[FR Doc. 2021–04362 Filed 3–2–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0362]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice for Manufacturing, Processing, Packing, and Holding of Finished Pharmaceuticals, Including Medical Gases, and Active Pharmaceutical Ingredients

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the Federal Register concerning each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection associated with current good manufacturing practice (CGMP) for drugs, finished pharmaceuticals, including medical gases, and active pharmaceutical ingredients (APIs).

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