Affected Public: Insured state nonmember banks and state savings associations.

Burden Estimate:

Type of burden	Number of respondents	Estimated time per response	Frequency of response	Total estimated annual burden (hours)
Reporting	1	16	On Occasion	16
Recordkeeping	1	166	On Occasion	166
Disclosure	1	1,332	On Occasion	1,332
Total Estimated Annual Burden				1,514

General Description of Collection: This information collection implements section 742(c)(2) of the Dodd-Frank Act (7 U.S.C. 2(c)(2)(E) and FDIC regulations governing retail foreign exchange transactions as set forth at 12 CFR part 349, subpart B. The regulation allows banking organizations under FDIC supervision to engage in off-exchange transactions in foreign currency with retail customers provided they comply with various reporting, recordkeeping and third-party disclosure requirements specified in the rule. If an institution elects to conduct such transactions, compliance with the information collection is mandatory.

Reporting Requirements—Part 349, subpart B requires that, prior to initiating a retail foreign exchange business; a banking institution must provide the FDIC with a notice certifying that the institution has written policies and procedures, and risk measurement and management systems and controls in place to ensure that retail foreign exchange transactions are conducted in a safe and sound manner. The institution must also provide information about it intends to manage customer due diligence, new product approvals and haircuts applied to noncash margin.

Recordkeeping Requirements—Part 349 subpart B requires that institutions engaging in retail foreign exchange transactions keep full, complete and systematic records of account, financial ledger, transaction, memorandum orders and post execution allocations of bunched orders. In addition, institutions are required to maintain records regarding their ratio of profitable accounts, possible violations of law, records of noncash margin and monthly statements and confirmations issued.

Disclosure Requirements—The regulation requires that, before opening an account that will engage in retail foreign exchange transactions, a banking institution must obtain from each retail foreign exchange customer an acknowledgement of receipt and understanding of a written disclosure specified in the rule and of disclosures about the banking institution's fees and other charges and of its profitable accounts ratio. The institution must also provide monthly statements to each retail foreign exchange customer and must send confirmation statements following every transaction. The customer dispute resolution provisions of the regulation require certain endorsements, acknowledgements and signature language as well as the timely provision of a list of persons qualified to handle a customer's request for arbitration.

There is no change in the method or substance of the collection. At present no FDIC-supervised institution is engaging in activities that would make them subject to the information collection requirements. FDIC originally estimated that 3 institutions would be impacted by the rule. The agency is reducing the estimated number of respondents to one (1) as a placeholder in case an institution elects to engage in covered activities in the future. There has been no change in the frequency of response or in the estimated number of hours required to respond. Because of the reduction in the estimated number of respondents from three (3) to one (1), the estimated annual burden has decreased.

Request for Comment

Comments are invited on: (a) Whether the collections of information are necessary for the proper performance of the FDIC's functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collections, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collections of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Dated at Washington, DC, this 12th day of May 2017.

Federal Deposit Insurance Corporation.

Ralph E. Frable,

Assistant Executive Secretary.

[FR Doc. 2017–09992 Filed 5–16–17; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL MARITIME COMMISSION

Notice of Agreements Filed

The Commission hereby gives notice of the filing of the following agreements under the Shipping Act of 1984. Interested parties may submit comments on the agreements to the Secretary, Federal Maritime Commission, Washington, DC 20573, within twelve days of the date this notice appears in the **Federal Register**. Copies of the agreements are available through the Commission's Web site (*www.fmc.gov*) or by contacting the Office of Agreements at (202)-523–5793 or *tradeanalysis@fmc.gov.*

Agreement No.: 012395–002.

Title: MSC/ACL Trans-Atlantic Space Charter Agreement.

Parties: Atlantic Container Line A.B. and MSC Mediterranean Shipping Company S.A.

Filing Party: Wayne R. Rohde, Esq.; Cozen O'Connor; 1200 Nineteenth St. NW.; Washington, DC 200036.

Synopsis: The amendment revises Article 5.1 to clarify that the space to be provided to ACL will be on MSC's SAWC–USA–NWC service. The amendment also reinserts language that was inadvertently deleted by Amendment No. 1 and deletes language that was inadvertently added by Amendment No. 1. It also restates the Agreement.

Agreement No.: 012483.

Title: HLAG/CMA CGM U.S.-Mediterranean Slot Charter Agreement.

Parties: Hapag-Lloyd AG and CMA CGM S.A.

Filing Party: Wayne R. Rohde, Esq.; Cozen O'Connor; 1200 Nineteenth St. NW.; Washington, DC 200036.

Synopsis: The Agreement authorizes HLAG to sell space to CMA CGM on its MGX service in the trade between ports on the U.S. Gulf Coast on the one hand, and ports on the Gulf Coast of Mexico and in Italy, Spain and Jamaica on the other hand.

Agreement No.: 012484.

Title: Port of New York & New Jersey/OCEMA Discussion Agreement.

Parties: Port Authority of New York and New Jersey and the Ocean Carrier Equipment Management Association, Inc., FMC Agreement No. 011284 (OCEMA).

Filing Party: Sam Ruda; Port Authority of NY & NJ; 4 World Trade Center; 150 Greenwich Street—17th Floor; New York, NY 10007.

Synopsis: The Agreement authorizes the Port of New York and New Jersey and OCEMA to collect and exchange information, discuss, and reach agreement upon matters relating to the Cargo Facility Charge levied by the Port Authority of New York and New Jersey.

By Order of the Federal Maritime Commission.

Dated: May 12, 2017.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2017–09987 Filed 5–16–17; 8:45 am]

BILLING CODE 6731–AA–P

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 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 Web site. The guidances identified in this notice were developed using the process described in that guidance.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment(s) on these draft guidances before it begins work on the final version of such guidances, submit either electronic or written comments on the draft guidance by July 17, 2017.

ADDRESSES: You may submit comments 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):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, 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 Division of Dockets Management 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 Division of Dockets Management. 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 Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Submit written requests for single copies of the draft guidances 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 documents.

FOR FURTHER INFORMATION CONTACT: Xiaoqiu Tang, Center for Drug