Respondents: Business or other for-profit entities and State, Local or Tribal Government.

Frequency of Response: On occasion reporting requirement; Third party disclosure requirement.

Number of Respondents and Responses: 1 respondent; 1 response.

Estimated Time per Response: 12 hours.

Total Annual Burden: 12 hours.

Total Annual Cost: No cost.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in Sections 4(i) and 623 of the Communications Act of 1934, as amended.

Nature and Extent of Confidentiality: There is no need to confidentiality with this collection of information.

Privacy Impact Assessment(s): No impact(s).

Needs and Uses: The information collection requirements contained in 47 CFR 76.922(b)(5) provides that an eligible small system that elects to use the streamlined rate reduction process must implement the required rate reductions and provide written notice of such reductions to local subscribers, the local franchising authority (''LFA''), and the Commission.

OMB Control Number: 3060–0938.

Title: Application for a Low Power FM Broadcast Station License, FCC Form 319.

Form Number: FCC Form 319.

Type of Review: Extension of a currently approved collection.

Respondents: Not-for-profit institutions, State, local or Tribal Government.

Number of Respondents and Responses: 200 respondents and 200 responses.

Estimated Time per Response: 1 hour.

Frequency of Response: On occasion reporting requirement.

Total Annual Burden: 200 hours.

Total Annual Cost: $27,500.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection of information is contained in Sections 154(i), 303 and 308 of the Communications Act of 1934, as amended.

Privacy Impact Assessment(s): No impacts.

Needs and Uses: On January 20, 2000, the Commission adopted a Report and Order (R&O) in MM Docket No. 99–25, In the Matter of Creation of Low Power Radio Service. With the adoption of this R&O, the Commission authorized the licensing of two new classes of FM radio stations, generally referred to as low power FM stations (LPFM): A LP100 class for stations operating at 50–100 watts effective radiated power (ERP) at an antenna height above average terrain (HAAT) of 30 meters; and a LP10 class for stations operating at 1–10 watts ERP and an antenna height of 30 meters HAAT. These stations will be operated on a noncommercial educational basis by entities that do not hold attributable interests in any other broadcast station or other media subject to the Commission's ownership rules. The LPFM service authorized in this Report and Order provides significant opportunities for new radio services. The LPFM service creates a class of radio stations designed to serve very localized communities or underrepresented groups within communities.

In connection with this new service, the Commission developed a new FCC Form 319, Application for a Low Power FM Broadcast Station License. FCC Form 319 is required to apply for a license for a new or modified Low Power FM (LPFM) station.

Federal Communications Commission.

Cecilia Sigmund,

Federal Register Liaison Officer, Office of the Secretary.

[FR Doc. 2020–04970 Filed 3–10–20; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0997; FRS 16544]

Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995, the Federal Communications Commission (FCC or the Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees. The FCC may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid Office of Management and Budget (OMB) control number.

DATES: Written PRA comments should be submitted on or before May 11, 2020. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Nicole Ongele, FCC, via email *PRA@fcc.gov* and to *Nicole.Ongele@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Nicole Ongele at (202) 418–2991.

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060–0997.

Title: Section 52.15(k), Numbering Utilization and Compliance Audit.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Businesses or other for-profit.

Number of Respondents and Responses: 10 respondents; 10 responses.

Estimated Time Per Response: 33 hours.

Frequency of Response: Third party disclosure requirement.

Obligation to Respond: Mandatory. Statutory authority for this information collection is contained in 47 U.S.C. Section 251.

Total Annual Burden: 330 hours.

Total Annual Cost: No cost.

Privacy Act Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: Commission employees and the independent auditor are prohibited by 47 U.S.C. 220(f) from divulging any fact or information that may come to their knowledge in the course of performing the audit, except as directed by the Commission or a court.

Needs and Uses: The audit program, consisting of audit procedures and guidelines, is developed to conduct

random audits. The random audits are conducted on the carriers that use numbering resources in order to verify the accuracy of numbering data reported on FCC Form 502, and to monitor compliance with FCC rules, orders and applicable industry guidelines. Failure of the audited carriers to respond to the audits can result in penalties. Based on the final audit report, evidence of potential violations may result in enforcement action.

Federal Communications Commission.

Cecilia Sigmund,

Federal Register Liaison Officer, Office of the Secretary.

[FR Doc. 2020–04967 Filed 3–10–20; 8:45 am]

BILLING CODE 6712–01–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, relevant information, or 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 *tradeanalysis@fmc.gov.*

Agreement No.: 201334.

Agreement Name: COSCO/ONE/OOCL/YM EMED–USEC Vessel Sharing Agreement.

Parties: COSCO SHIPPING Lines Co., Ltd.; Ocean Network Express Pte. Ltd.; Orient Overseas Container Line Limited; and Yang Ming Marine Transport Corp., Yang Ming (UK) Ltd., and Yang Ming (Singapore) Pte. Ltd. (acting as a single party).

Filing Party: Joshua Stein; Cozen O'Connor.

Synopsis: The Agreement authorizes the parties to cooperate on the provision of a service operating between the U.S. East Coast and ports in the Mediterranean.

Proposed Effective Date: 3/4/2020.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/27479.

Agreement No.: 012056–002.

Agreement Name: WWOcean/EUKOR Joint Operating Agreement.

Parties: Wallenius Wilhelmsen Ocean AS and EUKOR Car Carriers, Inc.

Filing Party: Wayne Rohde; Cozen O'Connor.

Synopsis: This amendment revises Article 5.6(c) of the Agreement to exclude tug services from the services for which the parties are authorized to negotiate jointly.

Proposed Effective Date: 3/5/2020.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/2021.

Dated: March 6, 2020.

Rachel Dickon,

Secretary.

[FR Doc. 2020–05002 Filed 3–10–20; 8:45 am]

BILLING CODE 6731–AA–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–D–0420]

Providing Regulatory Submissions in Alternate Electronic Format; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is announcing the availability of a draft guidance for industry entitled ''Providing Regulatory Submissions in Alternate Electronic Format.'' Under the Federal Food, Drug, and Cosmetic Act (FD&C Act), Congress granted FDA the authority to implement the statutory electronic submission requirements in guidance. In response, FDA implemented binding guidance requiring that new drug applications (NDAs), abbreviated new drug applications (ANDAs), certain drug master files (DMFs), certain biologics license applications (BLAs), and certain investigational new drug applications (INDs) be submitted to the Agency in electronic common technical document (eCTD) format. Recognizing that some submissions are exempt from this requirement and that waivers of the requirement may be granted on a case-by-case basis, the Agency is issuing this draft guidance to describe the alternate electronic format sponsors or applicants should use for submissions covered under such exemptions and waivers.

DATES: Submit either electronic or written comments on the draft guidance by May 11, 2020 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

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–2020–D–0420 for ''Providing Regulatory Submissions in Alternate Electronic Format.'' 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