to Federal Maritime Commission, 800 North Capitol Street NW, Washington, DC 20573–0001, and replies shall be served on Petitioner's counsel, Richard D. Gluck, Foster Garvey PC, 1000 Potomac Street NW, Suite 200, Washington, DC 20007, *Richard.Gluck@foster.com.*

Non-confidential filings may be submitted in hard copy to the Secretary at the above address or by email as a PDF attachment to *Secretary@fmc.gov* and include in the subject line: P1–22 (Commenter/Company). Confidential filings should not be filed by email. A confidential filing must be filed with the Secretary in hard copy only and be accompanied by a transmittal letter that identifies the filing as ''Confidential-Restricted'' and describes the nature and extent of the confidential treatment requested. The Commission will provide confidential treatment to the extent allowed by law for confidential submissions, or parts of submissions, for which confidentiality has been requested. When a confidential filing is submitted, there must also be submitted a public version of the filing. Such public filing version shall exclude confidential materials and shall indicate on the cover page and on each affected page ''Confidential materials excluded.'' Public versions of confidential filings may be submitted by email. The Petition will be posted on the Commission's website at *http://www.fmc.gov/P1-22.* Replies filed in response to the Petition will also be posted on the Commission's website at this location.

William Cody,

Secretary.

[FR Doc. 2022–04437 Filed 3–2–22; 8:45 am]

BILLING CODE 6730–02–P

FEDERAL MARITIME COMMISSION

Notice of Agreements Filed

The Commission hereby gives notice of 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, 800 North Capitol Street, 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**, and the Commission requests that comments be submitted within 7 days on agreements that request expedited review. 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.: 201379.

Agreement Name: Crowley/King Ocean Dominican Republic/Haiti Space Charter Agreement.

Parties: Crowley Latin America Services, LLC and King Ocean Services Limited, Inc.

Filing Party: Wayne Rohde; Cozen O'Connor.

Synopsis: The Agreement authorizes the parties to charter space to/from one another on a space available/as used basis in the trade between the U.S. Atlantic Coast on the one hand and ports in the Dominican Republic and Haiti on the other hand.

Proposed Effective Date: 2/22/2022.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/59504.

Agreement No.: 201380.

Agreement Name: Crowley/Antillean Dominican Republic/Haiti Space Charter Agreement.

Parties: Antillean Marine Shipping Corporation and Crowley Latin America Services, LLC.

Filing Party: Wayne Rohde; Cozen O'Connor.

Synopsis: The Agreement authorizes the parties to charter space to/from one another on a space available/as used basis in the trade between the U.S. Atlantic Coast on the one hand and ports in the Dominican Republic and Haiti on the other hand.

Proposed Effective Date: 2/24/2022.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/59505.

Agreement No.: 201356–002.

Agreement Name: PFLG/NPDL Slot Charter Agreement.

Parties: Neptune Pacific Direct Line Pte. Ltd. and Pacific Forum Line (Group) Limited.

Filing Party: David Monroe; GKG Law.

Synopsis: The amendment updates the amount of space being chartered under the Agreement.

Proposed Effective Date: 2/17/2022.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/39510.

Agreement No.: 201378.

Agreement Name: NPDL/PFLG Slot Charter Agreement.

Parties: Neptune Pacific Direct Line Pte. Ltd. and Pacific Forum Line (Group) Limited.

Filing Party: David Monroe; GKG Law.

Synopsis: The purpose of this agreement is to allow NPDL to charter space to PFLG in the relevant trades.

Proposed Effective Date: 2/17/2022.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/59502.

Dated: February 18, 2022.

William Cody,

Secretary.

[FR Doc. 2022–04436 Filed 3–2–22; 8:45 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10398 #64]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. On October 23, 2011, OMB approved our initial request to use the generic clearance process under control number 0938–1148 (CMS–10398). It was last approved on April 26, 2021, via the standard PRA process which included the publication of 60- and 30-day **Federal Register** notices. The scope of the April 2021 umbrella accounts for Medicaid and CHIP State plan amendments, waivers, demonstrations, and reporting. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit comments regarding our burden estimates or any other aspect of this collection of information, including: The necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of

the estimated burden, ways to enhance the quality, utility and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by March 17, 2022.

ADDRESSES: When commenting, please reference the applicable form number (see below) and the OMB control number (0938–1148). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: CMS–10398 (#64)/OMB control number: 0938–1148, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may access CMS' website at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

Generic Information Collection

1. *Title of Information Collection:* Medicaid Section 1115 Substance Use Disorder (SUD) Demonstration: Federal Meta-Analysis Support; *Type of Information Collection Request:* Revision of a currently approved collection; *Use:* Starting in 2015, in response to the opioid epidemic, CMS offered states the flexibility to test Medicaid coverage of a full substance use disorder (SUD) treatment service array in the context of overall SUD service delivery transformation through the authority of section 1115 demonstrations. In 2017, CMS modified the requirements for SUD section 1115 demonstrations to improve access to clinically appropriate treatment for OUD and other SUDs, to better support the development and expansion of comprehensive treatment strategies, and to incorporate improved progress and outcome monitoring. In 2018, CMS awarded the Federal Meta-Analysis Support contract to RTI International to understand the overall effectiveness of the groups of demonstrations with similar features and how variations in state demonstration features and the context in which they are implemented contribute to differences in effectiveness. The meta-analysis includes multiple rounds of qualitative data collection. The first round of interviews (both, Characteristics Interviews and Implementation Interviews) have been completed. This March 2022 collection of information request seeks OMB's approval for a second round (State-level Stakeholder Virtual Interviews) of data collection activities. The purpose is to learn about the perspectives of other types of stakeholders important to implementing the demonstration. Respondents would include stakeholders with differing perspectives, including leadership of behavioral health service providers and leadership of MCOs or third-party administrators in states with fee-for-service SUD treatment services. *Form Number:* CMS–10398 (#64) (OMB control number: 0938–1148); *Frequency:* Once; *Affected Public:* State, Local, or Tribal Governments, and the Private sector; *Number of Respondents:* 90; *Total Annual Responses:* 90; *Total Annual Hours:* 83. (For policy questions regarding this collection contact: Danielle Daly at 410–786–0897.)

Dated: February 28, 2022.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2022–04445 Filed 3–2–22; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0964]

Eduardo Navarro: Final Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is issuing an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) permanently debarring Eduardo Navarro from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Mr. Navarro was convicted of a felony under Federal law for conduct relating to the development or approval, including the process for development or approval, of any drug product under the FD&C Act. Mr. Navarro was given notice of the proposed permanent debarment and was given an opportunity to request a hearing to show why he should not be debarred. As of December 24, 2021 (30 days after receipt of the notice), Mr. Navarro had not responded. Mr. Navarro's failure to respond and request a hearing within the prescribed timeframe constitutes a waiver of his right to a hearing concerning this action.

DATES: This order is applicable March 3, 2022.

ADDRESSES: Submit applications for termination of debarment to the Dockets Management Staff, Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500, or at *https://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Jaime Espinosa, Division of Enforcement (ELEM–4029), Office of Strategic Planning and Operational Policy, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20857, 240–402–8743, or at *debarments@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

Section 306(a)(2)(A) of the FD&C Act (21 U.S.C. 335a(a)(2)(A)) requires debarment of an individual from providing services in any capacity to a person that has an approved or pending drug product application if FDA finds that the individual has been convicted of a felony under Federal law for conduct relating to the development or approval, including the process of development or approval, of any drug product under the FD&C Act. On August 11, 2021, Mr. Navarro was convicted as defined in section 306(l)(1) of the FD&C Act in the U.S. District Court for the Southern District of Florida, Miami Division, when the court accepted his plea of guilty and entered judgment against him for one count of Conspiracy to Defraud the United States in violation of 18 U.S.C. 371.

The factual basis for this conviction is as follows: As contained in the Information, entered into the docket on March 16, 2021, and the Factual Proffer in Support of his guilty plea, entered