airspace (SUA) which includes restricted areas as defined in FAA Order 7400.2. As a cooperating agency, the FAA coordinated the NEPA environmental impact analysis reviews closely with the Navy, and actively participated in the preparation of the Navy's FEA. In accordance with FAA Order 1050.1F, *Environmental Impacts: Policies and Procedures,* and the Council on Environmental Quality's (CEQ) NEPA implementing regulations at 40 CFR part 1500, the FAA conducted an independent evaluation and analysis of the NAWCWD FEA and adopted it in support of FAA's decision to establish R–2511 for implementation of the Navy's proposed action. Based on the environmental impact analyses in the Navy's FEA, the FAA has determined that it's Proposed Action of establishing R–2511, and the Navy's use of R–2511, would not result in any significant environmental impacts.

List of Subjects in 14 CFR Part 73

Airspace, Prohibited areas, Restricted areas.

The Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 73 as follows:

PART 73—SPECIAL USE AIRSPACE

■ 1. The authority citation for part 73 continues to read as follows:

Authority: 49 U.S.C. 106(f), 106(g); 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 73.25 California [Amended]

■ 2. Section 73.25 is amended as follows:

* * * * *

R–2511 Trona, CA [New]

Boundaries. Beginning at lat. 35°37′30″ N; long. 117°35′33″ W; to lat. 35°40′30″ N; long. 117°25′03″ W; to lat. 35°36′00″ N; long. 117°16′55″ W; to lat. 35°36′00″ N; long. 117°26′03″ W; to lat. 35°27′40″ N; long. 117°26′03″ W; to the point of beginning.

Designated Altitudes. 6,000 feet MSL, to but not including, FL 200.

Time of Designation. Intermittent, 0700–1700 local time, Monday–Friday; as activated by NOTAM at least 7-days in advance. Activation limited to no more than 36 days per year, and a maximum of two 2-hour blocks each day.

Controlling Agency. FAA, Joshua Control Facility, Edwards Air Force Base, CA.

Using Agency. Naval Air Warfare Center Weapons Division, China Lake, CA.

* * * * *

Issued in Washington, DC, on October 17, 2022.

Scott M. Rosenbloom,

Manager, Airspace Rules and Regulations.

[FR Doc. 2022–22783 Filed 10–19–22; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1

[Docket No. FDA–2022–D–1126]

Laboratory Accreditation for Analyses of Foods; Small Entity Compliance Guide; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the availability of a guidance for industry entitled ''Laboratory Accreditation for Analyses of Foods: What You Need to Know About the FDA Regulation: Guidance for Industry—Small Entity Compliance Guide.'' The small entity compliance guide (SECG) is intended to help small entities comply with the final rule entitled ''Laboratory Accreditation for Analyses of Foods.''

DATES: The announcement of the guidance is published in the **Federal Register** on October 20, 2022.

ADDRESSES: You may submit either electronic or written comments on Agency guidances 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–2022–D–1126 for ''Laboratory Accreditation for Analyses of Foods: What You Need to Know About the FDA Regulation: Guidance for Industry—Small Entity Compliance Guide.'' 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, 240–402–7500.

- 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.govinfo.gov/content/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, 240–402–7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the SECG to the Office of Strategic Planning and Operational Policy, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Element Building, Rockville, MD 20857. Send two self-addressed adhesive labels to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the SECG.

FOR FURTHER INFORMATION CONTACT: Stacie Hammack, Food and Feed Laboratory Operations, Office of Regulatory Affairs, Food and Drug Administration, 60 8th Street NE, Atlanta, GA 30309, 301–796–5817.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of December 3, 2021 (86 FR 68728), we issued a final rule establishing a program for the testing of food in certain circumstances by accredited laboratories, as required under the Federal Food, Drug, and Cosmetic Act (the final rule). Establishing this program will help FDA improve the safety of the U.S. food supply and protect U.S. consumers by helping to ensure that certain food testing of importance to public health is conducted subject to appropriate oversight and in accordance with appropriate model standards to produce reliable and valid test results. The final rule, which is codified at 21 CFR part 1, subpart R (21 CFR 1.1101 through 1.1201), became effective February 1, 2022.

We examined the economic implications of the final rule as required by the Regulatory Flexibility Act (5 U.S.C. 601–612) and determined that the final rule will have a significant economic impact on a substantial number of small entities. In compliance with section 212 of the Small Business Regulatory Enforcement Fairness Act (Pub. L. 104–121, as amended by Pub. L. 110–28), we are making available the SECG to explain the actions that a small entity must take to comply with the rule.

We are issuing the SECG consistent with our good guidance practices regulation (21 CFR 10.115(c)(2)). The SECG represents the current thinking of FDA on this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

II. Paperwork Reduction Act of 1995

While this guidance contains no collection of information, it does refer to previously approved FDA collections of information. Therefore, clearance by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521) is not required for this guidance. The previously approved collections of information are subject to review by OMB under the PRA. The collections of information in 21 CFR part 1, subpart R have been approved under OMB control number 0910–0898.

III. Electronic Access

Persons with access to the internet may obtain the SECG at *https://www.fda.gov/food/food-safety-modernization-act-fsma/fsma-rules-guidance-industry, https://www.fda.gov/regulatory-information/search-fda-guidance-documents,* or *https://www.regulations.gov.* Use the FDA website listed in the previous sentence to find the most current version of the guidance.

Dated: October 14, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–22706 Filed 10–19–22; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 165

[Docket Number USCG–2022–0867]

RIN 1625–AA00

Safety Zone; Corpus Christi Shipping Channel, Corpus Christi, TX

AGENCY: Coast Guard, Department of Homeland Security (DHS).

ACTION: Temporary final rule.

SUMMARY: The Coast Guard is establishing a temporary safety zone for all navigable waters of the Corpus Christi Shipping Channel in a zone defined by the following coordinates; 27°50′31.28″ N, 97°04′17.23″ W; 27°50′31.73″ N, 97°04′15.44″ W; 27°50′29.06″ N, 97°04′16.61″ W; 27°50′29.32″ N, 97°04′14.82″ W. The safety zone is needed to protect personnel, vessels, and the marine environment from potential hazards created by pipelines that will be removed from the floor of the Corpus Christi Shipping Channel. Entry of vessels or persons into this zone is prohibited unless specifically authorized by the Captain of the Port Sector Corpus Christi or a designated representative.

DATES: This rule is effective from 7 p.m. on October 19, 2022, through 5 a.m. on October 22, 2022. For the purposes of enforcement, actual notice will be used from 7 p.m. on October 19, 2022, through October 20, 2022.

FOR FURTHER INFORMATION CONTACT: If you have questions on this rule, call or email Lieutenant Commander Anthony Garofalo, Sector Corpus Christi Waterways Management Division, U.S. Coast Guard; telephone 361–939–5130, email *CCWaterways@uscg.mil.*

SUPPLEMENTARY INFORMATION:

I. Table of Abbreviations

CFR Code of Federal Regulations
DHS Department of Homeland Security
FR Federal Register
NPRM Notice of proposed rulemaking
§ Section
U.S.C. United States Code

II. Background Information and Regulatory History

The Coast Guard is issuing this temporary rule without prior notice and opportunity to comment pursuant to authority under section 4(a) of the Administrative Procedure Act (APA) (5 U.S.C. 553(b)). This provision authorizes an agency to issue a rule