final rule for any proposal to amend the standard of identity for canned Pacific salmon that may result from the National Food Processors Association's petition, submitted on behalf of Bumble Bee Seafoods, Inc., and other salmon packers holding temporary permits, or 30 days after termination of such proposal.

In the **Federal Register** of April 24, 2020 (85 FR 23047), we issued a notice announcing that we amended the temporary permit to allow for the canned skinless and boneless chunk salmon packed in water with or without sodium tripolyphosphate and to allow the test product to be manufactured only at one plant, Pataya Food Industries Ltd., located at 90/6 Moo 7, Settakit Road, Tambol Tarsai, Amphur Maung, Samutsakorn 74000, Thailand.

Under our regulations at 21 CFR 130.17(f), we are amending the temporary permit issued to Bumble Bee Seafoods, Inc., to allow the test product to be manufactured at an additional plant, RS Cannery Company Limited, located at 255/1 Industrial Soi 3, Bangpoo Industrial Estate, Samutprakarn 10280, Thailand. All other conditions and terms of this permit remain the same.

Dated: November 23, 2020.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2020–26533 Filed 12–1–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2013–N–1119; FDA–2010–N–0622; FDA–2011–N–0016; FDA–2009–N–0501; and FDA–2019–N–6098]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7726, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507). The OMB control number and expiration date of OMB approval for each information collection are shown in table 1. Copies of the supporting statements for the information collections are available on the internet at *https://www.reginfo.gov/public/do/PRAMain.* An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control No.	Date approval expires
Food Canning Establishment Registration, Process Filing and Recordkeeping for Acidified and Thermally Processed Low-Acid Foods	0910–0037	10/31/2023
Color Additive Certification Requests and Recordkeeping	0910–0216	10/31/2023
Recordkeeping and Records Access Requirements for Food Facilities	0910–0560	10/31/2023
Reporting and Recordkeeping Requirements for Reportable Food	0910–0643	10/31/2023
Focus Groups as Used by the Food and Drug Administration	0910–0497	11/30/2023

Dated: November 25, 2020.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2020–26571 Filed 12–1–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0190]

Agency Information Collection Activities; Proposed Collection; Comment Request; Infant Formula Requirements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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 collection of information, including 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 provisions of our infant formula regulations, including infant formula labeling, quality control procedures, notification requirements, and recordkeeping. The notice also invites comment on electronic Form FDA 3978 that allows manufacturers of infant formula to submit reports and notifications in a standardized format.

DATES: Submit either electronic or written comments on the collection of information by February 1, 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 February 1, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of February 1, 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.

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