not limited to, the remedies provided under 19 U.S.C. 1641.

* * * * *

Susan S. Thomas,

Executive Assistant Commissioner, Office of Trade, U.S. Customs and Border Protection.

[FR Doc. 2026–22735 Filed 11–5–26; 8:45 am]

BILLING CODE 9111–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 571

[Docket No. FDA–2026–F–4956]

Food Additive Petitions; Change of Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is amending the regulations for food additive petitions (FAPs) to update addresses and simplify notification procedures. These technical amendments are being made to improve the accuracy of the Agency's regulations and are non-substantive.

DATES: This technical amendment is effective November 6, 2026.

FOR FURTHER INFORMATION CONTACT: Isabel Pocurull, Center for Veterinary Medicine, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–5877, *Isabel.Pocurull@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: FDA is amending the animal food and drug regulations for food additive petitions (FAPs) at 21 CFR 571.1 to update addresses and Agency notification procedure. This technical amendment is being made to improve the accuracy of the Agency's regulations and is non-substantive.

Publication of this document constitutes final action on the changes under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because this amendment to the regulations provides only technical or non-substantive, ministerial changes to update addresses and Agency notification procedure.

List of Subjects in 21 CFR Part 571

Administrative practice and procedure, Animal feeds, Animal food additives, Confidential business information, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 571 is amended as follows:

PART 571—FOOD ADDITIVE PETITIONS

■ 1. The authority citation for part 571 continues to read as follows:

Authority: 21 U.S.C. 321, 342, 348, 371; 42 U.S.C. 241.

■ 2. Amend § 571.1 by:
■ a. Revising paragraph (a);
■ b. In paragraph (c), revising the undesignated text above item A; and
■ c. Revising paragraph (d).

The revisions read as follows:

§ 571.1 Petitions.

(a) Petitions to be filed under the provisions of section 409(b) of the Federal Food, Drug, and Cosmetic Act (the Act) can be submitted using both CVM's eSubmitter program, *https://www.fda.gov/industry/fda-esubmitter/cvm-esubmitter-programs,* to generate the submission and FDA's electronic submission gateway to submit, *https://www.fda.gov/industry/electronic-submissions-gateway-next-generation-esg-nextgen,* or on compact disk with paper cover letter. If any part of the material submitted is in a foreign language, it shall be accompanied by an accurate and complete English translation. The petition shall state petitioner's mailing address and email address to which published notices or orders issued or objections filed pursuant to section 409 of the Act may be sent.

* * * * *

(c) * * *

(Date) _____

Name of petitioner _____

Post office address _____

Date _____

Name of food additive and proposed use _____

Food and Drug Administration, Center for Veterinary Medicine, *Director,* Division of Animal Food Ingredients, 5001 Campus Dr., College Park, MD 20740–3835

The undersigned, _____, submits this petition pursuant to section 409(b)(1) of the Federal Food, Drug, and Cosmetic Act with respect to _____ (Name of the food additive and proposed use)

Attached and constituting a part of this petition, are the following:

* * * * *

(d) The petitioner will be notified of the date on which the petition is filed or if an incomplete petition, it will usually be retained but not filed as a petition under section 409 of the Act. The petitioner will be notified in what respects the petition is incomplete.

* * * * *

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2026–22663 Filed 11–5–26; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 874

[Docket No. FDA–2026–N–12125]

Medical Devices; Ear, Nose, and Throat Devices; Classification of the External Mechanical Stimulator for the Relief of Congestion

AGENCY: Food and Drug Administration, HHS.

ACTION: Final amendment; final order.

SUMMARY: The Food and Drug Administration (FDA) is classifying the external mechanical stimulator for the relief of congestion into class II (special controls). The special controls that apply to the device type are identified in this order and will be part of the codified language for classification of the external mechanical stimulator for the relief of congestion. We are taking this action because we have determined that classifying the device into class II will provide a reasonable assurance of the safety and effectiveness of the device. We believe this action will also enhance patients' access to beneficial innovative devices, in part by reducing regulatory burdens.

DATES: This order is effective November 6, 2026. The classification was applicable on December 4, 2023.

FOR FURTHER INFORMATION CONTACT: Payton Lin, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1212, Silver Spring, MD 20993–0002, 240–402–6580, *Payton.Lin@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

Upon request, FDA (the Agency or we) has classified the external mechanical stimulator for the relief of congestion into class II (special controls), which we have determined will provide a reasonable assurance of the safety and effectiveness of the