

(i) GE GE90–100 Service Bulletin 72–0926, Revision 01, dated December 22, 2023.

(ii) [Reserved]

(3) For GE material identified in this AD, contact General Electric Company, 1 Neumann Way, Cincinnati, OH 45215; phone: (513) 552–3272; email: aviation.fleetsupport@ge.com; website: ge.com.

(4) You may view this material at the FAA, Airworthiness Products Section, Operational Safety Branch, 1200 District Avenue, Burlington, MA 01803. For information on the availability of this material at the FAA, call (817) 222–5110.

(5) You may view this material at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, visit www.archives.gov/federal-register/cfr/ibr-locations or email fr.inspection@nara.gov.

Issued on July 25, 2025.

Peter A. White,

Deputy Director, Integrated Certificate Management Division, Aircraft Certification Service.

[FR Doc. 2025–14287 Filed 7–28–25; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 172

[Docket No. FDA–2025–F–2137]

Spoonbill Foundation; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of petition.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that we have filed a food additive petition, submitted by the Spoonbill Foundation, proposing that we amend our food additive regulations to provide for the safe use of 4'-phosphopantetheine (4'-PPT) as a nutrient in medical food.

DATES: The food additive petition was filed on July 3, 2025.

ADDRESSES: For access to the docket to read background documents or 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.

FOR FURTHER INFORMATION CONTACT: Marissa Santos, Human Foods Program,

Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–8160.

SUPPLEMENTARY INFORMATION: Under section 409(b)(5) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 348(b)(5)), we are giving notice that we have filed a food additive petition (FAP 5A4842), submitted on behalf of Spoonbill Foundation by Immix Law Group, 500 NW Naito Pkwy, Unit G, Portland, OR 97209. The petition proposes that we amend our food additive regulations in part 172 (21 CFR 172), “Food Additives Permitted For Direct Addition to Food For Human Consumption,” to provide for the safe use of 4'-PPT as a nutrient in medical food.

The petitioner has claimed that this action is categorically excluded under 21 CFR 25.32(k) because granting of this petition would become effective “for substances added directly to food that are intended to remain in food through ingestion by consumers and are not intended to replace macronutrients in food.” In addition, the petitioner has stated that, to their knowledge, no extraordinary circumstances exist. If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is required. If FDA determines a categorical exclusion does not apply, we will request an environmental assessment and make it available for public inspection.

Dated: July 25, 2025.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2025–14339 Filed 7–28–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2025–F–2423]

APIX Biosciences US LLC; Filing of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of petition.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that we have filed a food additive petition, submitted by APIX Biosciences US LLC, proposing that we amend our food additive regulations to

provide for the safe use of cholesterol as a source of sterol in food for honeybees at a level between 0.009 and 0.5% by weight of the food.

DATES: The food additive petition was filed on July 11, 2025.

ADDRESSES: For access to the docket to read background documents or 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.

FOR FURTHER INFORMATION CONTACT:

Wasima Wahid, Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Place, Rockville, MD 20855, 240–402–5857, Wasima.Wahid@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: Under section 409(b)(5) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 348(b)(5)), we are giving notice that we have filed a food additive petition (FAP 2322), submitted by APIX Biosciences US LLC, 2580 NE Rivercrest Rd., Fayetteville, AR 72701. The petition proposes that we amend our food additive regulations in 21 CFR part 573—Food Additives Permitted in Feed and Drinking Water of Animals to provide for the safe use of cholesterol as a source of sterol in food for honeybees at a level between 0.009 and 0.5% by weight of the food.

The petitioner has claimed that this action is categorically excluded under 21 CFR 25.32(r) because it is of a type that does not individually or cumulatively have a significant effect on the human environment. In addition, the petitioner has stated that, to their knowledge, no extraordinary circumstances exist that may significantly affect the quality of the human environment. If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is required. If FDA determines a categorical exclusion does not apply, we will request an environmental assessment and make it available for public inspection.

Dated: July 25, 2025.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2025–14335 Filed 7–28–25; 8:45 am]

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