AIRAC date	State	City	Airport name	FDC No.	FDC date	Procedure name
25–Jan–24	GU	Guam	Guam Intl	3/1623	10/17/23	RNAV (RNP) Z RWY 6L, Orig-D.
25–Jan–24	TX	Weslaco	Mid Valley	3/1772	10/26/23	VOR–A, Orig-B.
25–Jan–24	VA	Roanoke	Roanoke/Blacksburg Rgnl (Woodrum Fld).	3/3252	11/20/23	LDA Z RWY 6, Orig.
25–Jan–24	VA	Roanoke	Roanoke/Blacksburg Rgnl (Woodrum Fld).	3/3255	11/20/23	RNAV (RNP) Z RWY 24, Orig.
25–Jan–24	VA	Roanoke	Roanoke/Blacksburg Rgnl (Woodrum Fld).	3/3639	11/20/23	RNAV (RNP) Z RWY 6, Orig.
25–Jan–24	SC	Summerville	Summerville	3/3863	10/12/23	RNAV (GPS) RWY 24, Orig-C.
25–Jan–24	CA	Mountain View	Moffett Federal Airfield ..	3/4042	11/9/23	ILS OR LOC RWY 32R, Amdt 2.
25–Jan–24	FL	Fort Lauderdale	Fort Lauderdale/Hollywood Intl.	3/4077	11/2/23	RNAV (GPS) Y RWY 28R, Amdt 5.
25–Jan–24	SD	Aberdeen	Aberdeen Rgnl	3/4485	11/21/23	RNAV (GPS) RWY 35, Amdt 1.
25–Jan–24	SD	Aberdeen	Aberdeen Rgnl	3/4487	11/21/23	RNAV (GPS) RWY 31, Orig-B.
25–Jan–24	KS	Norton	Norton Muni	3/5231	9/29/23	RNAV (GPS) RWY 16, Amdt 1B.
25–Jan–24	PA	Harrisburg	Harrisburg Intl	3/5232	10/23/23	RNAV (GPS) RWY 31, Amdt 1A.
25–Jan–24	MO	Higginsville	Higginsville Industrial Muni.	3/5990	10/13/23	RNAV (GPS) RWY 34, Amdt 1.
25–Jan–24	MO	Higginsville	Higginsville Industrial Muni.	3/5993	10/13/23	RNAV (GPS) RWY 16, Amdt 1.
25–Jan–24	FL	Panama City	Northwest Florida Beaches Intl.	3/6411	10/27/23	ILS OR LOC RWY 16, ILS RWY 16 (SA CAT I), ILS RWY 16 (SA CAT II), Amdt 3A.
25–Jan–24	MI	Lansing	Capital Region Intl	3/7469	10/18/23	ILS OR LOC RWY 28L, Amdt 28A.
25–Jan–24	FL	Orlando	Orlando Intl	3/8059	10/5/23	RNAV (GPS) RWY 17R, Orig-D.
25–Jan–24	FL	Orlando	Orlando Intl	3/8060	10/5/23	RNAV (GPS) RWY 35L, Amdt 1A.
25–Jan–24	AK	Cordova	Merle K (Mudhole) Smith.	3/8167	11/6/23	RNAV (GPS)–B, Amdt 2B.
25–Jan–24	FL	West Palm Beach	Palm Beach Intl	3/8298	10/20/23	RNAV (GPS) Y RWY 14, Amdt 4.
25–Jan–24	CA	Mountain View	Moffett Federal Airfield ..	3/8470	11/9/23	RNAV (GPS) RWY 32L, Amdt 1.
25–Jan–24	NY	Malone	Malone-Dufort	3/9901	10/5/23	RNAV (GPS) RWY 23, Orig-D.
25–Jan–24	KY	Henderson	Henderson City-County	3/9903	10/6/23	RNAV (GPS) RWY 9, Amdt 1A.

[FR Doc. 2023–27826 Filed 12–18–23; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

[Docket No. FAA–2022–1203]

14 CFR Chapter I

Policy Regarding Processing Land Use Changes on Federally Acquired or Federally Conveyed Airport Land; Correction

AGENCY: Federal Aviation Administration (FAA), Department of Transportation (DOT).

ACTION: Notification of final policy; correction.

SUMMARY: The Federal Aviation Administration published a document in the **Federal Register** of December 8, 2023, concerning its Policy Regarding Processing Land Use Changes on Federally Acquired or Federally Conveyed Airport Land. The document contained an incorrect FAA Docket Number.

DATES: This correction is effective January 8, 2024.

FOR FURTHER INFORMATION CONTACT: Michael Helvey, Airport Compliance and Management Analysis, ACO–1, Federal Aviation Administration, 800 Independence Avenue SW, Washington, DC 20591, telephone (202) 267–3085; facsimile: (202) 267–4629.

SUPPLEMENTARY INFORMATION:

Correction

In the **Federal Register** of December 8, 2023, in FR Doc. 2023–27017, on page 85474, in the second column, correct the ''Docket No.'' to read:

[Docket No. FAA–2022–1203]

Dated: December 12, 2023.

Michael Helvey,

Director, Office of Airport Compliance and Management Analysis.

[FR Doc. 2023–27829 Filed 12–18–23; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Part 744

[Docket No. 231213–0301]

RIN 0694–AJ50

Removals From the Unverified List

AGENCY: Bureau of Industry and Security, Department of Commerce.

ACTION: Final rule.

SUMMARY: The Bureau of Industry and Security (BIS) is amending the Export Administration Regulations (EAR) by removing four persons, all under the destination of People's Republic of China (China), from the UVL because BIS was able to verify their bona fides.

DATES: *This rule is effective:* December 15, 2023.

FOR FURTHER INFORMATION CONTACT: Kevin J. Kurland, Deputy Assistant Secretary for Export Enforcement, Phone: (202) 482–4255 or by email at *UVLRequest@bis.doc.gov*.

SUPPLEMENTARY INFORMATION:

Background

The UVL, found in supplement no. 6 to part 744 of the EAR (15 CFR parts 730–774), contains the names and addresses of foreign persons who are or have been parties to a transaction, as described in § 748.5 of the EAR, involving the export, reexport, or transfer (in-country) of items subject to the EAR. These foreign persons are added to the UVL because BIS or federal officials acting on BIS's behalf were unable to verify their *bona fides* (*i.e.,* legitimacy and reliability relating to the end-use and end user of items subject to the EAR) through an end-use check. These checks, such as a pre-license check (PLC) or a post-shipment verification (PSV), cannot be completed satisfactorily for reasons outside the U.S. Government's control.

There are any number of reasons why these checks cannot be completed to the satisfaction of the U.S. Government. Section 744.15(c)(1) of the EAR provides illustrative examples of those circumstances, including reasons unrelated to the cooperation of the foreign party subject to the end-use check. Such examples include: (i) During the conduct of an end-use check, the subject of the check is unable to demonstrate the disposition of items subject to the EAR; (ii) The existence or authenticity of the subject of an end-use check cannot be verified (*e.g.,* the subject of the check cannot be located or contacted); (iii) Lack of cooperation by the host government authority prevents an end-use check from being conducted.

BIS's inability to confirm the *bona fides* of foreign persons subject to end-use checks raises concerns about the suitability of such persons as participants in future exports, reexports, or transfers (in-country) of items subject to the EAR; it also indicates a risk that such items may be diverted to prohibited end uses and/or end users. Under such circumstances, there may not be sufficient information to add the foreign person at issue to the Entity List under § 744.11 of the EAR. Therefore, BIS may add the foreign person to the UVL.

As provided in § 740.2(a)(17) of the EAR, the use of license exceptions for exports, reexports, and transfers (in-country) involving a party or parties to the transaction who are listed on the UVL is suspended. Additionally, under § 744.15(b) of the EAR, there is a requirement for exporters, reexporters, and transferors to obtain (and maintain a record of) a UVL statement from a party or parties to the transaction who are listed on the UVL before proceeding with exports, reexports, and transfers (in-country) to such persons, when the exports, reexports and transfers (in-country) are not subject to a license requirement. Finally, pursuant to § 758.1(b)(8), Electronic Export Information (EEI) must be filed in the Automated Export System (AES) for all exports of tangible items subject to the EAR where any party to the transaction, as described in § 748.5(d) through (f), is listed on the UVL.

Requests for the removal of a UVL entry must be made in accordance with § 744.15(d) of the EAR. Decisions regarding the removal or modification of a UVL entry will be made by the Deputy Assistant Secretary for Export Enforcement, based on a demonstration by the listed person of their *bona fides*.

Removals From the UVL

This final rule removes four persons from the UVL because BIS was able to verify their *bona fides*. This rule removes Chengde Oscillator Electronic Technology Co., China National Erzhong Group, Ningbo III Lasers Technology Co., Ltd., and Xinjiang East Hope New Energy Company Ltd., all under the destination of China. BIS is removing these four persons pursuant to § 744.15(c)(2) of the EAR.

Rulemaking Requirements

Executive Order Requirements

Executive Orders 13563 and 12866 direct agencies to assess all costs and benefits of available regulatory alternatives and, if regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety effects, distribute impacts, and equity). Executive Order 13563 emphasizes the importance of quantifying both costs and benefits, of reducing costs, of harmonizing rules, and of promoting flexibility. This final rule has not been designated a ''significant regulatory action'' under Executive Order 12866.

This rule does not contain policies with Federalism implications as that term is defined under Executive Order 13132.

Paperwork Reduction Act Requirements

Notwithstanding any other provision of law, no person is required to respond to, nor is subject to a penalty for failure to comply with, a collection of information, subject to the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*) (PRA), unless that collection of information displays a currently valid Office of Management and Budget (OMB) Control Number.

The UVL additions contain collections of information approved by OMB under the following control numbers:

- OMB Control Number 0694–0088—Simple Network Application Process and Multipurpose Application Form
- OMB Control Number 0694–0122—Miscellaneous Licensing Responsibilities and Enforcement
- OMB Control Number 0694–0134—Entity List and Unverified List Requests,
- OMB Control Number 0694–0137—License Exemptions and Exclusions.

BIS believes that the overall increases in burdens and costs will be minimal and will fall within the already approved amounts for these existing collections. Additional information regarding these collections of information—including all background materials—can be found at *https://www.reginfo.gov/public/do/PRAMain* by using the search function to enter either the title of the collection or the OMB Control Number.

Administrative Procedure Act and Regulatory Flexibility Act Requirements

Pursuant to Section 1762 of ECRA (50 U.S.C. 4821), this action is exempt from the Administrative Procedure Act (5 U.S.C. 553) requirements for notice of proposed rulemaking and opportunity for public participation.

Further, no other law requires notice of proposed rulemaking or opportunity for public comment for this final rule. Because a notice of proposed rulemaking and an opportunity for public comment are not required under the Administrative Procedure Act or by any other law, the analytical requirements of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*) are not applicable.

List of Subjects in 15 CFR Part 744

Exports, Reporting and recordkeeping requirements, Terrorism.

Accordingly, part 744 of the Export Administration Regulations (15 CFR parts 730 through 774) is amended as follows:

PART 744—END-USE AND END–USER CONTROLS

■ 1. The authority citation for 15 CFR part 744 continues to read as follows:

Authority: 50 U.S.C. 4801–4852; 50 U.S.C. 4601 *et seq.;* 50 U.S.C. 1701 *et seq.;* 22 U.S.C. 3201 *et seq.;* 42 U.S.C. 2139a; 22 U.S.C. 7201 *et seq.;* 22 U.S.C. 7210; E.O. 12058, 43 FR 20947, 3 CFR, 1978 Comp., p. 179; E.O. 12851, 58 FR 33181, 3 CFR, 1993 Comp., p. 608; E.O. 12938, 59 FR 59099, 3 CFR, 1994

Comp., p. 950; E.O. 13026, 61 FR 58767, 3 CFR, 1996 Comp., p. 228; E.O. 13099, 63 FR 45167, 3 CFR, 1998 Comp., p. 208; E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783; E.O. 13224, 66 FR 49079, 3 CFR, 2001 Comp., p. 786; Notice of November 8, 2022, 87 FR 68015, 3 CFR, 2022 Comp., p. 563; Notice of September 7, 2023, 88 FR 62439 (September 11, 2023).

■ 2. Supplement No. 6 to Part 744 is amended under CHINA, PEOPLE'S REPUBLIC OF, by removing the entries for ''Chengde Oscillator Electronic Technology Co.'', ''China National Erzhong Group'', ''Ningbo III Lasers Technology Co., Ltd.'', and ''Xinjiang East Hope New Energy Company Ltd''.

Thea D. Rozman Kendler,

Assistant Secretary for Export Administration.

[FR Doc. 2023–27932 Filed 12–15–23; 11:15 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2020–F–0151]

Food Additives Permitted in Feed and Drinking Water of Animals; Calcium Formate

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA, we, or the Agency) is amending the regulations for food additives permitted in feed and drinking water of animals to provide for the safe use of calcium formate as a feed acidifying agent, to lower the pH, in complete swine and poultry feeds at levels not to exceed 1.2 percent of the complete feed. This action is in response to a food additive petition filed by LANXESS Corp.

DATES: This rule is effective December 19, 2023. See section V for further information on the filing of objections. Either electronic or written objections and requests for a hearing on the final rule must be submitted by January 18, 2024.

ADDRESSES: You may submit objections and requests for a hearing as follows. Please note that late, untimely filed objections will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of January 18, 2024. Objections received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic objections in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting objections. Objections submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your objection will be made public, you are solely responsible for ensuring that your objection 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 objection, that information will be posted on *https://www.regulations.gov.*
- If you want to submit an objection with confidential information that you do not wish to be made available to the public, submit the objection 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 objections submitted to the Dockets Management Staff, FDA will post your objection, 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–2020–F–0151 for ''Food Additives Permitted in Feed and Drinking Water of Animals; Calcium Formate.'' Received objections, those filed in a timely manner (see **ADDRESSES**), 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 an objection with confidential information that you do not wish to be made publicly available, submit your objections only as a written/paper submission. You should submit two copies in 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 objections. 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 objections 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.

FOR FURTHER INFORMATION CONTACT: Wasima Wahid, Center for Veterinary Medicine (HFV–221), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–402–5857, *wasima.wahid@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

In a document published in the **Federal Register** of February 11, 2020 (85 FR 7682), FDA announced that we had filed a food additive petition (animal use) (FAP 2310) submitted by LANXESS Corp., 111 RIDC Park West Dr., Pittsburgh, PA 15275. The petition proposed that the regulations for food additives permitted in feed and drinking water of animals be amended to provide for the safe use of calcium formate as a feed acidifying agent, to lower the pH, in complete feeds for swine or poultry.