noise sensitive areas; and increases in minimum altitudes and landing minima. As such, this action is not expected to result in any potentially significant environmental impacts. In accordance with FAA Order 1050.1F, paragraph 5–2 regarding Extraordinary Circumstances, the FAA has reviewed this action for factors and circumstances in which a normally categorically excluded action may have a significant environmental impact requiring further analysis. Accordingly, the FAA has determined that no extraordinary circumstances exist that warrant preparation of an environmental assessment or environmental impact study.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

The Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—DESIGNATION OF CLASS A, B, C, D, AND E AIRSPACE AREAS; AIR TRAFFIC SERVICE ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 14 CFR part 71 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.

§ 71.1 [Amended]

■ 2. The incorporation by reference in 14 CFR 71.1 of FAA Order JO 7400.11G, Airspace Designations and Reporting Points, dated August 19, 2022, and effective September 15, 2022, is amended as follows:

Paragraph 6011 United States Area Navigation Routes

* * * * *

T–378 BRION, AK to Fort Yukon, AK (FYU) [New]

BRION, AK	FIX	(Lat. 66°09′38.95″ N, long. 150°12′25.77″ W)
ZUSPA, AK	WP	(Lat. 66°18′20.43″ N, long. 147°51′04.14″ W)
DUTKE, AK	WP	(Lat. 66°25′02.96″ N, long. 146°57′36.10″ W)
Fort Yukon, AK (FYU)	VORTAC	(Lat. 66°34′27.31″ N, long. 145°16′35.97″ W)

* * * * *

Issued in Washington, DC, on October 26, 2022.

Scott M. Rosenbloom,

Manager, Airspace Rules and Regulations.

[FR Doc. 2022–23725 Filed 10–31–22; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Docket No. FAA–2021–1152; Airspace Docket No. 19–AAL–72]

RIN 2120–AA66

Amendment of United States Area Navigation (RNAV) Route T–269; Yakutat, AK

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule; correction.

SUMMARY: This action corrects a final rule published by the FAA in the **Federal Register** on October 24, 2022, that amends United States Area Navigation (RNAV) route T–269 in the vicinity of Yakutat, AK, in support of a large and comprehensive T-route modernization project for the state of Alaska. The final rule identified the KATAT, AK, route point as a waypoint (WP), in error. This action makes an editorial correction to the reference of the KATAT, AK, WP to change it to be reflected as a Fix and match the FAA's aeronautical database information.

DATES: Effective date 0901 UTC, December 29, 2022. The Director of the Federal Register approves this incorporation by reference action under 1 CFR part 51, subject to the annual revision of FAA Order 7400.11 and publication of conforming amendments.

ADDRESSES: FAA Order 7400.11G, Airspace Designations and Reporting Points, and subsequent amendments can be viewed online at *www.faa.gov/air_traffic/publications/*. For further information, you can contact the Rules and Regulations Group, Federal Aviation Administration, 800 Independence Avenue SW, Washington, DC 20591; telephone: (202) 267–8783.

FOR FURTHER INFORMATION CONTACT: Colby Abbott, Rules and Regulations Group, Office of Policy, Federal Aviation Administration, 800 Independence Avenue SW, Washington, DC 20591; telephone: (202) 267–8783.

SUPPLEMENTARY INFORMATION:

History

The FAA published a final rule in the **Federal Register** (87 FR 64159; October 24, 2022), amending T–269 in support of a large and comprehensive T-route modernization project for the state of Alaska. Subsequent to publication, the FAA determined that the KATAT, AK, route point was inadvertently identified as a WP, in error. This rule corrects that error by changing the reference of the KATAT, AK, WP to the KATAT, AK, Fix. This is an editorial change only to match the FAA's aeronautical database information and does not alter the alignment of the affected T–269 route.

United States Area Navigation Routes are published in paragraph 6011 of FAA Order JO 7400.11G, dated August 19, 2022, and effective September 15, 2022, which is incorporated by reference in 14 CFR 71.1. The RNAV T-route listed in this document will be published subsequently in FAA Order JO 7400.11.

Correction to Final Rule

Accordingly, pursuant to the authority delegated to me, reference to the KATAT, AK, WP that is reflected in Docket No. FAA–2021–1152, as published in the **Federal Register** of October 24, 2022 (87 FR 64159), FR Doc. 2022–22496, is corrected as follows:

■ 1. On page 64160, correct the table for T–269 Annette Island, AK (ANN) to MKLUK, AK [Amended] to read:

T–269 Annette Island, AK (ANN) to MKLUK, AK [Amended]

Annette Island, AK (ANN)	VOR/DME	(Lat. 55°03′37.47″ N, long. 131°34′42.24″ W)
Biorka Island, AK (BKA)	VORTAC	(Lat. 56°51′33.87″ N, long. 135°33′04.72″ W)
Yakutat, AK (YAK)	VOR/DME	(Lat. 59°30′38.99″ N, long. 139°38′53.26″ W)
MALAS, AK	WP	(Lat. 59°39′58.52″ N, long. 140°34′57.61″ W)
OXIDS, AK	WP	(Lat. 59°41′51.68″ N, long. 141°03′17.73″ W)
FOGNU, AK	WP	(Lat. 59°53′31.88″ N, long. 141°49′02.83″ W)
HORGI, AK	WP	(Lat. 60°00′04.68″ N, long. 142°35′23.34″ W)
ZIXIM, AK	WP	(Lat. 60°03′48.75″ N, long. 143°13′27.77″ W)

JOVOM, AK	WP	(Lat. 60°07′40.55″ N, long. 143°42′56.99″ W)
OXUGE, AK	WP	(Lat. 60°06′15.81″ N, long. 144°13′28.54″ W)
KATAT, AK	FIX	(Lat. 60°15′29.17″ N, long. 144°42′18.77″ W)
Johnstone Point, AK (JOH)	VOR/DME	(Lat. 60°28′51.43″ N, long. 146°35′57.61″ W)
Anchorage, AK (TED)	VOR/DME	(Lat. 61°10′04.32″ N, long. 149°57′36.51″ W)
MKLUK, AK	WP	(Lat. 60°26′40.04″ N, long. 165°55′17.28″ W)

* * * * *

Issued in Washington, DC, on October 25, 2022.

Scott M. Rosenbloom,

Manager, Airspace Rules and Regulations.

[FR Doc. 2022–23536 Filed 10–31–22; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1271

[Docket No. FDA–2022–D–0563]

Regulation of Human Cells, Tissues, and Cellular and Tissue-Based Products: Small Entity Compliance Guide; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing the availability of a final guidance entitled ''Regulation of Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps): Small Entity Compliance Guide.'' The small entity compliance guide (SECG) is intended to help small entity establishments that manufacture HCT/Ps better understand the comprehensive regulatory framework for HCT/Ps set forth in the regulations and comply with certain HCT/P-related final rules. The SECG announced in this notice supersedes the SECG of the same title dated August 2007.

DATES: The announcement of the guidance is published in the **Federal Register** on November 1, 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–0563 for ''Regulation of Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps): 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 Communication, Outreach and Development, Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 3128, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. The guidance may also be obtained by mail by calling CBER at 1–800–835–4709 or 240–402–8010. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the SECG.

FOR FURTHER INFORMATION CONTACT: Myrna Hanna, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 7301, Silver Spring, MD 20993–0002, 240–402–7911.

SUPPLEMENTARY INFORMATION: