COGNU	WP	(Lat. 65°48′29″ N., long. 167°50′06″ W.)
PHO	NDB	(Lat. 68°20′41″ N., long. 166°47′51″ W.)
T–262 ODK to JOH [New]		
ODK	VORTAC	(Lat. 57°46′30″ N., long. 152°20′23″ W.)
WUXAN	WP	(Lat. 59°53′00″ N., long. 149°00′00″ W.)
JOH	VOR/DME	(Lat. 60°28′51″ N., long. 146°35′58″ W.)
T–264 ODK to MDO [New]		
ODK	VORTAC	(Lat. 57°46′30″ N., long. 152°20′23″ W.)
ZAXUM	WP	(Lat. 58°41′15″ N., long. 147°53′26″ W.)
MDO	VOR/DME	(Lat. 59°25′18″ N., long. 146°21′00″ W.)
T–266 CGL to FPN [New]		
CGL	NDB	(Lat. 58°21′33″ N., long. 134°41′58″ W.)
FPN	NDB	(Lat. 56°47′32″ N., long. 132°49′15″ W.)
T–268 FPN to ICK [New]		
FPN	NDB	(Lat. 56°47′32″ N., long. 132°49′15″ W.)
ICK	NDB	(Lat. 55°04′15″ N., long. 131°36′18″ W.)

* * * * *

Issued in Washington, DC, on June 16, 2005.

Edith V. Parish,

Acting Manager, Airspace and Rules.

[FR Doc. 05–12366 Filed 6–21–05; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Docket No. FAA–2004–19851; Airspace Docket No. 04–AAL–13]

RIN 2120–AA66

Modification and Revocation of Federal Airways; AK

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule; correction.

SUMMARY: This action corrects an error in the airspace description of a notice of a final rule that was published in the **Federal Register** on May 6, 2005 (70 FR 23934), Airspace Docket No. 04–AAL–13.

DATES: *Effective Date:* 0901 UTC, July 7, 2005.

FOR FURTHER INFORMATION CONTACT: Ken McElroy, Airspace and Rules, Office of System Operations and Safety, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591; telephone: (202) 267–8783.

SUPPLEMENTARY INFORMATION:

History

On May 6, 2005, Airspace Docket No. 04–AAL–13, was published in the **Federal Register** (70 FR 23934), revising Jet Route 133 (J–133), AK. In that rule, the airspace description was incomplete. This action corrects that error.

Correction to Final Rule

■ Accordingly, pursuant to the authority delegated to me, the legal description for J–133, as published in the **Federal Register** on May 6, 2005 (70 FR 23934), on page 23934 and incorporated by reference in 14 CFR 71.1, is corrected as follows:

PART 71—[AMENDED]

§ 71.1 [Amended]

Paragraph 2004—Jet Airways

* * * * *

J–133 [Corrected]

J–133: From Sitka, AK NDB; via INT Sitka, AK NDB 308° and Orca Bay, AK, NDB 114°; Orca Bay, AK; Johnstone Point, AK; Anchorage, AK; to Galena AK.

* * * * *

Issued in Washington, DC, on June 10, 2005.

Edith V. Parish,

Acting Manager, Airspace and Rules.

[FR Doc. 05–12126 Filed 6–21–05; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Parts 121 and 135

[Docket No. FAA–2004–18477; Amendment Nos. 121–312; 135–98]

Aircraft Assembly Placard Requirements

AGENCY: Federal Aviation Administration, DOT.

ACTION: Final rule; Notice of Office of Management and Budget approval for information collection and addition of amendment numbers.

SUMMARY: This notice announces the Office of Management and Budget's (OMB) approval of the information collection requirement in the final rule published on June 29, 2004 (FR 69 39292). This notice also provides the amendment numbers for the final rule that were absent when it was published.

DATES: Final rule; Aircraft Assembly Placard Requirement was published in the **Federal Register** on June 29, 2004. FAA received OMB approval for the information collection requirement on November 8, 2004. The final rule becomes effective June 22, 2005.

FOR FURTHER INFORMATION CONTACT: Gary Davis, Flight Standards Service, Air Transportation Division, AFS–201A, Federal Aviation Administration, 800 Independence Avenue SW., Washington, DC 20591; telephone (202) 267–8166; facsimile (202) 267–5229; e-mail: *gary.davis@faa.gov.*

SUPPLEMENTARY INFORMATION:

Background

On June 29, 2004, the FAA published the Final Rule, Aircraft Assembly Placard Requirements, as instructed by an act of Congress. The rule instructed affected air carriers to display a placard with information on where the aircraft was assembled. We instructed air carriers to provide that information in one sentence on the seat-pocket cards that inform passengers of emergency procedures.

As noted in the preamble, the final rule would not become effective until the FAA received approval from OMB for the information collection that was required in the rule. In the **DATES** section of the final rule, we said that when that approval was received we would publish a notice in the **Federal Register** announcing the effective date.

In accordance with the Paperwork Reduction Act, OMB approved the FAA's request for new information collection on November 8, 2004. Please note that 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. The OMB control number associated with this collection is 2120–0691. The request was approved by OMB without change and expires on November 30, 2007.

Additionally, the Final Rule was published without amendment numbers. This notice adds those amendment numbers as shown in the heading.

49 U.S.C. 106(g), 40113, 40119, 41706, 44101, 44701–44702, 44705, 44709–44711, 44713, 44716–44717, 44722, 46105, grants authority to the Administrator to publish this notice. The final rule (FR 69 39292) is effective immediately.

Issued in Washington, DC, on June 15, 2005.

Anthony F. Fazio,

Director, Office of Rulemaking.

[FR Doc. 05–12239 Filed 6–17–05; 11:35 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 172

[Docket No. 2002F–0160]

Food Additives Permitted for Direct Addition to Food for Human Consumption; Vitamin D_3

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correcting amendments.

SUMMARY: The Food and Drug Administration (FDA) is responding to objections and is denying requests that it has received for a hearing on the final rule that amended the food additive regulations authorizing the use of vitamin D_3 as a nutrient supplement in calcium-fortified fruit juices and fruit drinks, excluding fruit juices and fruit juice drinks specially formulated or processed for infants, at levels not to exceed 100 International Units (IU) per serving. (In the final rule, FDA used the term ''fruit drink;'' however, the common or usual name of the product is ''fruit juice drink.'' Therefore, FDA is replacing the term ''fruit drink'' with ''fruit juice drink.'') In response to one of the objections, FDA is amending the vitamin D_3 regulation to replace the current 100 IU per serving limits on the vitamin D_3 fortification of fruit juices and fruit juice drinks with limits of 100 IU per 240 milliliters (mL). This document also corrects three errors that appeared in the codified portion of the vitamin D_3 final rule.

DATES: This rule is effective June 22, 2005. Submit written or electronic objections and requests for a hearing by July 22, 2005. See section IX of this document for information on the filing of objections.

ADDRESSES: You may submit written or electronic objections and requests for a hearing, identified by Docket No. 2002F–0160, by any of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov*. Follow the instructions for submitting comments.
- Agency Web site: *http://www.fda.gov/dockets/ecomments*. Follow the instructions for submitting comments on the agency Web site.
- E-mail: *fdadockets@oc.fda.gov*. Include Docket No. 2002F–0160 in the subject line of your e-mail message.
- FAX: 301–827–6870.
- Mail/Hand delivery/Courier [For paper, disk, or CD-ROM submissions]: Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

Instructions: All submissions received must include the agency name and docket number for this rulemaking. All objections received will be posted without change to *http://www.fda.gov/ohrms/dockets/default.htm*, including any personal information provided. For detailed instructions on submitting objections, see the ''Objections'' heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or objections received, go to *http://www.fda.gov/ohrms/dockets/default.htm* 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 Division of Dockets Management, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Judith L. Kidwell, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–1071.

SUPPLEMENTARY INFORMATION:

I. Introduction

In the **Federal Register** of April 25, 2002 (67 FR 20533), FDA published a notice announcing the filing of a food additive petition (FAP 2A4734) by the Minute Maid Co. (Minute Maid), to amend the food additive regulations in part 172 *Food Additives Permitted for Direct Addition to Food for Human Consumption* (21 CFR part 172) to provide for the safe use of vitamin D_3 as a nutrient supplement in calcium-fortified fruit juices and fruit juice drinks. In response to FAP 2A4734, in the **Federal Register** of February 27, 2003 (68 FR 9000), FDA issued a final rule permitting the safe use of vitamin D_3 as a nutrient supplement in calcium-fortified fruit juices and fruit juice drinks[1], excluding fruit juices and fruit juice drinks specially formulated or processed for infants, at levels not to exceed 100 IU per serving. This regulation was codified in § 172.380. FDA based its decision on data contained in the petition and in its files.

The preamble to the final rule advised that objections to the final rule and requests for a hearing were due within 30 days of the publication date, by March 31, 2003. FDA received several submissions within the 30-day objection period. Some of the submissions sought revocation of the final rule and requested a hearing. In response to one of the objections received during the 30-day objection period, FDA is amending the food additive regulation to replace those portions of the vitamin D_3 regulation that prescribe limits on vitamin D_3 fortification of fruit juices and fruit juice drinks of 100 IU per serving with limits of 100 IU per 240 mL. This document also corrects three errors that appeared in the codified portion of the vitamin D_3 final rule.

II. Objections and Requests for a Hearing

Section 409(f) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 348(f)), provides that, within 30 days after publication of an order relating to a food additive regulation, any person adversely affected by such order may file objections, specifying with particularity the provisions of the order ''* * * deemed objectionable, stating reasonable grounds therefore, and requesting a public hearing [based] upon such objections.'' FDA may deny a hearing request if the objections to the regulation do not raise genuine and substantial issues of fact that can be resolved at a hearing.

Under 21 CFR 171.110 of the food additive regulations, objections and requests for a hearing are governed by part 12 (21 CFR part 12) of FDA's regulations. Under § 12.22(a) each

[1] In the final rule (68 FR 9000), FDA used the term ''fruit drink.'' In 21 CFR 102.33, the common or usual name of the product is ''fruit juice drink.'' To be consistent with § 102.33, FDA is replacing the term ''fruit drink'' with ''fruit juice drink'' in § 172.380(d) and elsewhere in this document.