the 2011–12 crop year; (3) handlers are aware of this action which was unanimously recommended by the Committee at a public meeting and is similar to other assessment rate actions issued in past years; and (4) this interim rule provides a 60-day comment period, and all comments timely received will be considered prior to finalization of this rule.

List of Subjects in 7 CFR Part 993

Marketing agreements, Plums, Prunes, Reporting and recordkeeping requirements.

For the reasons set forth in the preamble, 7 CFR part 993 is amended as follows:

PART 993—DRIED PRUNES PRODUCED IN CALIFORNIA

■ 1. The authority citation for 7 CFR part 993 continues to read as follows:

Authority: 7 U.S.C. 601–674.

■ 2. Section 993.347 is revised to read as follows:

§ 993.347 Assessment rate.

On and after August 1, 2011, an assessment rate of $0.22 per ton is established for California dried prunes.

Dated: August 19, 2011.

David R. Shipman,

Acting Administrator, Agricultural Marketing Service.

[FR Doc. 2011–22119 Filed 8–29–11; 8:45 am]

BILLING CODE 3410–02–P

DEPARTMENT OF AGRICULTURE

Agricultural Marketing Service

7 CFR Part 1217

[Document Number AMS–FV–10–0015C; FR]

RIN 0581–AD03

Softwood Lumber Research, Promotion, Consumer Education and Industry Information Order; Correction

AGENCY: Agricultural Marketing Service.

ACTION: Corrections to final rule.

SUMMARY: This document contains corrections to the final rule published on August 2, 2011 (76 FR 46185), regarding softwood lumber. Corrections are made in the amendatory instruction section and in § 1217.88 of the final rule.

DATES: *Effective Date:* August 31, 2011.

FOR FURTHER INFORMATION CONTACT: Maureen T. Pello, Marketing Specialist, Research and Promotion Division, Fruit and Vegetable Programs, AMS, USDA, P.O. Box 831, Beavercreek, Oregon 97004; telephone: (503) 632–8848; facsimile (503) 632–8852; or electronic mail: *Maureen.Pello@ams.usda.gov.*

SUPPLEMENTARY INFORMATION:

Background

This rule establishes a Softwood Lumber Research, Promotion, Consumer Education and Industry Information Order (Order). The purpose of the Order is to strengthen the position of softwood lumber in the marketplace, maintain and expand markets for softwood lumber, and develop new uses for softwood lumber within the United States. The Order is issued pursuant to the Commodity Promotion, Research, and Information Act of 1996 (7 U.S.C. 7411–7425).

Corrections

In FR Doc. 2011–19491, published August 2, 2011 (76 FR 46185), make the following corrections.

1. On page 46193, in column 2, the words of issuance are corrected to read as follows:

For the reasons set forth in the preamble, Title 7, Chapter XI of the Code of Federal Regulations is amended by adding subpart A to part 1217 to read as follows:

2. On page 46194, column 1, the words ''Subpart B—[Reserved]'' are removed.

3. On page 46202 in column 1, § 1217.88 is revised to read as follows:

§ 1217.88 OMB Control numbers.

The control numbers assigned to the information collection requirements by the Office of Management and Budget pursuant to the Paperwork Reduction Act of 1995, 44 U.S.C. Chapter 35, are OMB control number 0505–0001 (Board nominee background statement) and OMB control number 0581–0264.

Dated: August 22, 2011.

David R. Shipman,

Acting Administrator.

[FR Doc. 2011–22150 Filed 8–29–11; 8:45 am]

BILLING CODE 3410–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 14

[Docket No. FDA–2011–N–0002]

Advisory Committee; Change of Name and Function; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the standing advisory committees' regulations to change the name and function of the Anesthetic and Life Support Drugs Advisory Committee. This action is being taken to reflect changes made to the charter for this advisory committee.

DATES: Effective September 6, 2011.

FOR FURTHER INFORMATION CONTACT: Philip Bautista, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001.

SUPPLEMENTARY INFORMATION: FDA is announcing that the name of the Anesthetic and Life Support Drugs Advisory Committee, which was established on May 1, 1978, has been changed. The Agency decided that the name ''Anesthetic and Analgesic Drug Products Advisory Committee'' would more accurately describe the subject areas for which the committee is responsible. The mandate of the committee is being expanded to include analgesics, *e.g.,* abuse-deterrent opioids, novel analgesics, and opioid abuse.

The Committee reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products including analgesics, *e.g.,* abuse-deterrent opioids, novel analgesics, and issues related to opioid abuse, and those for use in anesthesiology.

The Anesthetic and Life Support Drugs Advisory Committee name was changed and its functions expanded in the charter renewal dated June 9, 2011. FDA is hereby revising 21 CFR 14.100 (c)(1) to reflect these changes.

Publication of this final rule constitutes a final action on this change under the Administrative Procedure Act. Under 5 U.S.C. 553(b)(3)(B) and (d) and 21 CFR 10.40(d) and (e), the Agency finds good cause to dispense with notice and public procedure and to proceed to an immediately effective regulation. Such notice and procedures are unnecessary and are not in the public interest, because the final rule is merely codifying the new name and the expanded function of the advisory committee to reflect the current committee charter.

List of Subjects in 21 CFR Part 14

Administrative practice and procedure, Advisory committees, Color additives, Drugs, Radiation protection.

Therefore, under the Federal Food and Drug, and Cosmetic Act and under