PART 311—[REMOVED]

In accordance with the above and under the authority of 49 U.S.C. 40113, DOT removes 14 CFR Part 311.

Issued in Washington, DC, on this 1st day of April, 1996.

Federico Peña,

Secretary.

[FR Doc. 96–9703 Filed 4–19–96; 8:45 am]

BILLING CODE 4910–62–P

14 CFR Part 399

[Docket No. OST–96–1260; Notice 96–10]

RIN 2105–AC42

Release of Internal Staff Memoranda Relating to Public Meetings of the Civil Aeronautics Board

AGENCY: Office of the Secretary, Department of Transportation (DOT).

ACTION: Final rule.

SUMMARY: DOT is removing provisions concerning release of internal staff memorandum after public meetings of the Civil Aeronautics Board (CAB). The CAB was sunset in 1985 and the provisions no longer have any relevancy. This action is taken on the Department's initiative in response to the President's Regulatory Reinvention Initiative.

EFFECTIVE DATE: Effective May 22, 1996.

FOR FURTHER INFORMATION CONTACT: Robert I. Ross, Office of the General Counsel, C–10, Department of Transportation, Washington, DC 20590, telephone (202) 366–9156, FAX (202) 366–9170.

SUPPLEMENTARY INFORMATION: In 1985, the Civil Aeronautics Board (CAB) ceased to exist and many of its functions and resources were transferred to DOT. Some of its regulations, although no longer relevant, have remained in the Code of Federal Regulations. Specifically, § 399.102 of title 14, Code of Federal Regulations, has no relevancy to DOT The section specifically sets forth procedures peculiar to an agency that, like the CAB, was subject to the Government in the Sunshine Act, which does not apply to DOT. We are, therefore, removing the section because this procedure is not used at DOT. Analogous information may continue to be sought under the Freedom of Information Act. As part of the President's Regulatory Reinvention Initiative, we will be removing other duplicative or obsolete parts in separate rulemakings. Because these changes are editorial in nature and do not change the substantive requirements, the Department finds that notice and comment are unnecessary and contrary to the public interest.

Analysis of Regulatory Impacts

This amendment is not a ''significant regulatory action'' within the meaning of Executive Order 12866. It is also not significant within the definition in DOT's Regulatory Policies and Procedures, 49 FR 11034 (1979), in part because it does not involve any change in important Departmental policies. There is no economic impact as a result of this change. Moreover, I certify that this rule will not have a significant economic impact on a substantial number of small entities.

This rule does not significantly affect the environment, and therefore an environmental impact statement is not required under the National Environmental Policy Act of 1969. It has also been reviewed under Executive Order 12612, Federalism, and it has been determined that it does not have sufficient implications for federalism to warrant preparation of a Federalism Assessment.

Finally, the rule does not contain any collection of information requirements, requiring review under the Paperwork Reduction Act of 1980.

List of Subjects in 14 CFR Part 399

Administrative practice and procedure, Air carriers, Air rates and fares, Air taxis, Consumer protection, Small businesses.

In accordance with the above, DOT amends 14 CFR Part 399 as follows:

PART 399—[AMENDED]

1. The authority citation to Part 399 continues to read as follows:

Authority: 49 U.S.C. 40101, 40102, 40105, 40109, 40113, 40114, 40115, 41101, 41102, 41104, 41105, 41106, 41107, 41108, 41109, 41110, 41301, 41302, 41303, 41304, 41305, 41306, 41307, 41309, 41310, 41501, 41503, 41504, 41506, 41507, 41508, 41509, 41510, 41511, 41701, 41702, 41705, 41706, 41707, 41708, 41709, 41711, 41712, 41713, 41901, 41902, 41903, 41904, 41905, 41906, 41907, 41908, 41909, 41910, 41911, 41912, 42111, 42112, 46101, 46102, 46301, 46501.

§ 399.102 [Removed]

2. Section 399.102 is removed.

Issued in Washington, DC, on this 1st day of April, 1996.

Federico Peña,

Secretary of Transportation.

[FR Doc. 96–9702 Filed 4–19–96; 8:45 am]

BILLING CODE 4910–62–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor Name and Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor name and address from Agribusiness Marketers, Inc., to Mallinckrodt Veterinary Operations, Inc.

EFFECTIVE DATE: April 22, 1996.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: Agribusiness Marketers, Inc., 2667 West Dual, Baton Rouge, LA 70815, has informed FDA of a change of sponsor name and address to Mallinckrodt Veterinary Operations, Inc., 421 East Hawley St., Mundelein, IL 60060.

Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change of sponsor name and address.

List of Subjects in 21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 510 is amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e).

§ 510.600 [Amended]

2. Section 510.600 *Names, addresses, and drug labeler codes of sponsors of approved applications* is amended in the table in paragraph (c)(1) by removing the entry for ''Agribusiness Marketers, Inc.,'' and by alphabetically adding a new entry for ''Mallinckrodt Veterinary Operations, Inc., 421 East Hawley St., Mundelein, IL

60060.....015563'' and in the table in paragraph (c)(2) in the entry for ''015563'' by removing the sponsor name ''Agribusiness Marketers, Inc.,'' and adding in its place ''Mallinckrodt Veterinary Operations, Inc., 421 East Hawley St., Mundelein, IL 60060.''

Dated: April 4, 1996.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–9779 Filed 4–19–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 558

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect the change of sponsor name for a new animal drug application (NADA) from MAC–PAGE, Inc., to ADM Animal Health & Nutrition Div.

EFFECTIVE DATE: April 22, 1996.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 6, 1994 (59 FR 50828), FDA published a final rule amending the animal drug regulations to reflect the change of sponsors for all NADA's held by Central Soya, P. O. Box 1400, Fort Wayne, IN 46801–2508, including MAC–PAGE, Inc., 1600 South Wilson Ave., Dunn, NC 28334, a wholly-owned subsidiary of Central Soya, transferred to Premiere Agri Technologies, Inc. The subsidiaries retained their names and drug labeler codes. In the Federal Register of September 11, 1995 (60 FR 40752), FDA published a final rule amending the animal drug regulations to reflect the change of sponsor name from Premiere Agri Technologies, Inc., and the names of all wholly-owned subsidiaries, to ADM Animal Health & Nutrition Div., P.O. Box 2508, Fort Wayne, IN 46801–2508. Both final rules, which reflected these changes, inadvertently did not include NADA 131–957 (Tylosin). This document corrects that error. Accordingly, FDA is amending the regulations in 21 CFR 558.625 to reflect the change of sponsor.

List of Subject in 21 CFR Part 558

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.625 [Amended]

2. Section 558.625 *Tylosin* is amended in paragraph (b)(79) by removing ''047427'' and adding in its place ''012286''.

Dated: April 4, 1996.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–9784 Filed 4–19–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Parts 1313 and 1316

[DEA No. 112C]

Implementation of the Domestic Chemical Diversion Control Act of 1993 (Pub. L. 103–200); Correction

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Correction to final regulations.

SUMMARY: This document contains corrections to the final regulations which were published on Thursday, June 22, 1995 (60 FR 32447). The regulations related to the registration, recordkeeping and reporting requirements for manufacturers, distributors, importers and exporters of listed chemicals.

EFFECTIVE DATE: April 22, 1996.

FOR FURTHER INFORMATION CONTACT: G. Thomas Gitchel, Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, D.C. 20537, Telephone (202) 307–7297.

SUPPLEMENTARY INFORMATION: The final regulations that are the subject of these corrections implement the Domestic Chemical Diversion Control Act of 1993 (Pub. L. 103–200) (DCDCA). The regulations amend Title 21, Code of Federal Regulations, to add a new Part 1309 and revise certain sections in Parts 1310, 1313 and 1316. As published, the final regulations contain errors that could cause confusion in the regulated industry.

Accordingly, the publication June 22, 1995 of the final regulations to implement the DCDCA, which were the subject of Federal Register Document 95–14978, is corrected as follows:

PART 1313—[CORRECTED]

1. On page 32465, in the first column, the section heading which reads ''§ 1312.32 Requirement of authorization for international transactions.'' is corrected to read ''§ 1313.32 Requirement of authorization for international transactions.''

PART 1316—[CORRECTED]

2. On page 32465, in the third column, amendment Number 1 immediately following PART 1316—[AMENDED] is corrected to read as follows:

1. The authority citation for Part 1316, Subpart A is amended to read as follows:

Authority: 21 U.S.C. 822(f), 830(a), 871(b), 880, 958(f), 965.

Dated: April 16, 1996.

Stephen H. Greene,

Deputy Administrator, Drug Enforcement Administration.

[FR Doc. 96–9813 Filed 4–19–96; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF TRANSPORTATION

Federal Highway Administration

23 CFR Part 625

[FHWA Docket No. 95–12]

RIN 2125–AD38

Design Standards for Highways; Geometric Design of Highways and Streets

AGENCY: Federal Highway Administration (FHWA), DOT.

ACTION: Interim final rule; request for comments.

SUMMARY: The National Highway System (NHS) was established by the National Highway System Designation Act of 1995 (Pub. L. 104–59, 109 Stat. 568). To reflect the establishment of the NHS, the FHWA is revising several areas of the text in its regulation governing design standards for highways; updating the listing of standards; relocating the guides and references; and adopting as its interim