Note: Cessna Caravan Service Bulletin No. CAB04–9, dated October 4, 2004, also addresses the installation of the pilot assist handle.

May I Request an Alternative Method of Compliance?

(f) The Manager, Wichita Aircraft Certification Office (ACO), FAA, has the authority to approve alternative methods of compliance for this AD, if requested using the procedures found in 14 CFR 39.19. For information on any already approved alternative methods of compliance, contact Paul Pellicano, Aerospace Engineer (Icing), FAA, Small Airplane Directorate, c/o Atlanta ACO, One Crown Center, 1985 Phoenix Boulevard, Suite 450, Atlanta, GA 30349; telephone: (770) 703–6064; facsimile: (770) 703–6097; or Robert P. Busto, Aerospace Engineer, Wichita ACO, FAA, 1801 Airport Road, Wichita, Kansas 67209; telephone: (316) 946–4157; facsimile: (316) 946–4107.

Does This AD Incorporate Any Material by Reference?

(g) You must do the actions required by this AD following the instructions in Cessna Caravan Service Kit No. SK208–146, dated October 4, 2004, and Cessna Caravan Accessory Kit No. AK208–6C, Revision C, dated August 27, 1993.

(1) On February 22, 2006 (71 FR 1941, January 12, 2006), and in accordance with 5 U.S.C. 552(a) and 1 CFR part 51, the Director of the Federal Register previously approved the incorporation by reference of Cessna Caravan Service Kit No. SK208–146, dated October 4, 2004, and Cessna Caravan Accessory Kit No. AK208–6C, Revision C, dated August 27, 1993.

(2) To get a copy of this service information, contact The Cessna Aircraft Company, Product Support, P.O. Box 7706, Wichita, Kansas 67277–7706; telephone: (316) 517–5800; facsimile: (316) 942–9006. To review copies of this service information, go to the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, go to: *http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html* or call (202) 741–6030. To view the AD docket, go to the Docket Management Facility; U.S. Department of Transportation, 400 Seventh Street, SW., Nassif Building, Room PL–401, Washington, DC 20590–001 or on the Internet at *http://dms.dot.gov*. The docket number is FAA–2005–21275; Directorate Identifier 2005–CE–28–AD.

Appendix 1 to AD 2006–01–11 R1 Changes to the Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual

Affected Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual (AFM)

Insert the following text after the ''OTHER LIMITATIONS'' in the LIMITATIONS section of the Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual (AFM): COLD WEATHER OPERATIONS.

The airplane must be equipped with the following equipment when operating at an airport in the ground icing conditions defined under ''Visual/Tactile Check'' in the LIMITATIONS section:

1. Pilot assist handle, Cessna P/N SK208–146–2 (or FAA-approved equivalent part number).

Appendix 2 to AD 2006–01–11 R1 Changes to the Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual

Affected Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual (AFM)

Add the following to the equipment listed under ''FLIGHT INTO KNOWN ICING'' in the ''KINDS OF OPERATION LIMITS'' in the LIMITATIONS section of the FAA-Approved Airplane Flight Manual:

Lower main landing gear leading edge deice boots

Cargo pod nosecap deice boot

Appendix 3 to AD 2006–01–11 R1 Changes to the Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual Supplement S1

Affected Cessna Models 208 or 208B Pilot's Operating Handbook (POH) and FAA-Approved Airplane Flight Manual (AFM) Supplement S1

Remove the paragraph under ''REQUIRED EQUIPMENT'' in the Limitations section of the FAA-Approved Flight Manual Supplement S1 ''Known Icing Equipment'', that currently reads as follows:

''The following additional equipment is not required for flight into icing conditions as defined by FAR 25, but may be installed on early serial airplanes by using optional accessory Kit AK208–6. On later serial airplanes, this equipment may be included with the flight into known icing package. If installed, this equipment must be fully operational:''

Issued in Kansas City, Missouri, on March 10, 2006.

Kim Smith,

Manager, Small Airplane Directorate, Aircraft Certification Service.

[FR Doc. 06–2546 Filed 3–15–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510, 520, 522, 524, and 529

New Animal Drugs; Change of Sponsor's Drug Labeler Code

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of drug labeler code for Med-Pharmex, Inc.

DATES: This rule is effective March 16, 2006.

FOR FURTHER INFORMATION CONTACT: Charles Eastin, Center for Veterinary Medicine (HFV–210), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9077, e-mail: *charles.eastin@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: FDA has found that the animal drug regulations do not reflect the correct drug labeler code for Med-Pharmex, Inc. Accordingly, the agency is amending the regulations in 21 CFR 510.600, 520.1044a, 520.1195, 520.1484, 520.1485, 520.2220a, 520.2345d, 522.900, 524.1044b, 524.1044f, 524.1044g, 524.1193, 524.1443, 524.1580b, 524.1580e, 524.1600a, 524.2481, and 529.1044b to correct this error. In addition, 21 CFR 524.1044b, 524.1044f, 524.1443, and 524.2481 are being revised to reflect a current format.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects

21 CFR Part 510

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

21 CFR Parts 520, 522, 524, and 529

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 parts 510, 520, 522, 524, and 529 are amended as follows:

PART 510—NEW ANIMAL DRUGS

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

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

■ 2. Amend § 510.600 in the table in paragraph (c)(1) in the entry for ''Med-Pharmex, Inc.'' by removing ''051259'' and by adding in its place ''054925''; and in the table in paragraph (c)(2) by removing the entry for ''051259'' and by

numerically adding a new entry for ''054925'' to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(2) * * *

Drug labeler code	Firm name and address
* *	* * * *
054925	Med-Pharmex, Inc., 2727 Thompson Creek Rd., Pomona, CA 91767–1861
* *	* * * *

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 3. The authority citation for 21 CFR part 520 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 520.1044a [Amended]

■ 4. In paragraph (b) of § 520.1044a, remove ''051259'' and add in its place ''054925''.

§ 520.1195 [Amended]

■ 5. In paragraph (b)(2) of § 520.1195, remove ''051259'' and add in its place ''054925''.

§ 520.1484 [Amended]

■ 6. In paragraph (b)(1) of § 520.1484, remove ''051259'' and add in its place ''054925''.

§ 520.1485 [Amended]

■ 7. In paragraph (b) of § 520.1485, remove ''051259'' and add in its place ''054925''.

§ 520.2220a [Amended]

■ 8. In paragraphs (a)(1) and (a)(2) of § 520.2220a, remove ''051259'' and add in its place ''054925''.

§ 520.2345d [Amended]

■ 9. In paragraphs (b)(5), (d)(1)(iii), and (d)(2)(iii) of § 520.2345d, remove ''051259'' and add in its place ''054925''.

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

■ 10. The authority citation for 21 CFR part 522 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 522.900 [Amended]

■ 11. In paragraph (b)(1) of § 522.900, remove ''051259'' and add in its place ''054925''.

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

■ 12. The authority citation for 21 CFR part 524 continues to read as follows:

Authority: 21 U.S.C. 360b.

■ 13. Revise § 524.1044b to read as follows:

§ 524.1044b Gentamicin sulfate, betamethasone valerate otic solution.

(a) *Specifications.* Each milliliter of solution contains gentamicin sulfate equivalent to 3 milligrams (mg) gentamicin base and betamethasone valerate equivalent to 1 mg betamethasone alcohol.

(b) *Sponsors.* See Nos. 000061 and 054925 in § 510.600(c) of this chapter.

(c) *Conditions of use*—(1) *Amounts and indications for use*—(i) For the treatment of acute and chronic otitis externa caused by bacteria sensitive to gentamicin in dogs, instill three to eight drops of solution into the ear canal twice daily for 7 to 14 days.

(ii) For the treatment of infected superficial lesions caused by bacteria sensitive to gentamicin in dogs and cats, apply a sufficient amount of the drug to cover the treatment area twice daily for 7 to 14 days.

(2) *Limitations.* Federal law restricts this drug to use by or on the order of a licensed veterinarian.

■ 14. Revise § 524.1044f to read as follows:

§ 524.1044f Gentamicin sulfate, betamethasone valerate topical spray.

(a) *Specifications.* Each milliliter of spray contains gentamicin sulfate equivalent to 0.57 milligram (mg) gentamicin base and betamethasone valerate equivalent to 0.284 mg betamethasone.

(b) *Sponsors.* See Nos. 000061 and 054925 in § 510.600(c) of this chapter.

(c) *Conditions of use in dogs*—(1) *Amount.* Hold bottle upright 3 to 6 inches from the lesion and depress the sprayer head twice. Administer two spray actuations two to four times daily for 7 days.

(2) *Indications for use.* For the treatment of infected superficial lesions caused by bacteria sensitive to gentamicin.

(3) *Limitations.* Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 524.1044g [Amended]

■ 15. In paragraph (b)(2) of § 524.1044g, remove ''051259'' and add in its place ''054925''.

§ 524.1193 [Amended]

■ 16. In paragraph (b)(2) of § 524.1193, remove ''051259, 051311'' and add in its place ''051311, 054925''.

■ 17. Revise § 524.1443 to read as follows:

§ 524.1443 Miconazole.

(a) *Specifications*—(1) Each gram of cream contains miconazole nitrate equivalent to 20 milligrams miconazole base.

(2) Each gram of lotion or spray contains miconazole nitrate equivalent to 1 percent miconazole base.

(b) *Sponsors.* See § 510.600(c) of this chapter for use as in paragraph (c) of this section:

(1) No. 000061 for use of cream, lotion, and spray;

(2) Nos. 054925 and 058829 for use of lotion and spray.

(c) *Conditions of use in dogs and cats*—(1) *Amount.* Apply once daily by rubbing into or spraying a light covering on the infected site and the immediate surrounding vicinity. Continue treatment for 2 to 4 weeks until infection is completely eradicated as determined by appropriate laboratory examination.

(2) *Indications for use.* For topical treatment of infections caused by *Microsporum canis*, *Microsporum gypseum*, and *Trichophyton mentagrophytes.*

(3) *Limitations.* Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 524.1580b [Amended]

■ 18. In paragraph (b)(1) of § 524.1580b, remove ''051259'' and add in its place ''054925''.

§ 524.1580e [Amended]

■ 19. In paragraph (b) of § 524.1580e, remove ''051259'' and add in its place ''054925''.

§ 524.1600a [Amended]

■ 20. In paragraph (b) of § 524.1600a, remove both occurrences of ''051259, and 053501'' and add in their places ''053501, and 054925''.

■ 21. Revise § 524.2481 to read as follows:

§ 524.2481 Triamcinolone cream.

(a) *Specifications.* The vanishing cream contains 0.1 percent triamcinolone acetonide.

(b) *Sponsor.* See Nos. 053501 and 054925 in § 510.600(c) of this chapter.

(c) *Conditions of use in dogs*—(1) *Amount.* Rub into affected areas two to four times daily for 4 to 10 days.

(2) *Indications for use.* As an anti-inflammatory, antipruritic, and antiallergic agent for topical treatment of allergic dermatitis and summer eczema.

(3) *Limitations.* Federal law restricts this drug to use only by or on the order of a licensed veterinarian.

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

■ 22. The authority citation for 21 CFR part 529 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 529.1044b [Amended]

■ 23. In paragraph (b) of § 529.1044b, remove ''*Sponsor.* See Nos. 000061 and 051259'' and add in its place ''*Sponsors.* See Nos. 000061 and 054925''.

Dated: March 7, 2006.

Steven D. Vaughn,

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

[FR Doc. 06–2554 Filed 3–15–06; 8:45 am]

BILLING CODE 4160–01–S

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[FRL–8022–1]

Approval and Promulgation of Air Quality Implementation Plans; Arkansas Update to Materials Incorporated by Reference

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule; Notice of administrative change.

SUMMARY: EPA is updating the materials submitted by the State of Arkansas that are incorporated by reference (IBR) into the State Implementation Plan (SIP). The regulations affected by this update have been previously submitted by Arkansas and approved by EPA. This update affects the SIP materials that are available for public inspection at the National Archives and Records Administration (NARA), the Air and Radiation Docket and Information Center located at EPA Headquarters in Washington, DC, and the EPA Regional Office.

DATES: This rule is effective on March 16, 2006.

ADDRESSES: SIP materials incorporated by reference into 40 CFR part 52 are available for inspection at the following locations: Environmental Protection Agency, Region 6, 1445 Ross Avenue, Suite 700, Dallas, Texas 75202–2733; the EPA, Air and Radiation Docket and Information Center, Air Docket (Mail Code 6102T), 1200 Pennsylvania Avenue, NW., Washington, DC 20460, and the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call 202–741–6030, or go to: *http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html.*

FOR FURTHER INFORMATION CONTACT: Carl Young, Air Planning Section (6PD–L), Environmental Protection Agency, Region 6, 1445 Ross Avenue, Suite 700, Dallas, Texas 75202–2733, telephone 214–665–6645; fax number 214–665–7263, e-mail address *young.carl@epa.gov.*

SUPPLEMENTARY INFORMATION: Each State has an extensive SIP containing the control measures and strategies—such as air pollution control regulations, emission inventories, monitoring networks, attainment demonstrations, and enforcement mechanisms—used to attain and maintain the national ambient air quality standards (NAAQS).

Each State must formally adopt the control measures and strategies after the public has had an opportunity to comment on them and then must submit them to EPA for approval. Once these control measures and strategies are approved by EPA, after notice and comment, they are incorporated into the federally approved SIP and are identified in part 52, ''Approval and Promulgation of Implementation Plans'', Title 40 of the Code of Federal Regulations (40 CFR part 52). The full texts of a State's control measures and strategies approved by EPA are not reproduced in their entirety in 40 CFR part 52, but are ''incorporated by reference.'' This means that EPA has approved the identified State control measures and strategies, each with a specific effective date. The public is referred to the locations of the full text versions should they want to know which measures are contained in a given SIP. The information provided in 40 CFR part 52 allows EPA and the public to monitor a State's progress in implementing its SIP (and thus in attaining and maintaining the NAAQS) and to take enforcement action if necessary.

The SIP is a living document that the State can revise as necessary to address the unique air pollution problems in the State. Therefore, to incorporate the State's revisions into the federally approved SIP, EPA from time to time must take action on SIP revisions containing new and/or revised measures. On May 22, 1997. (62 FR 27968), EPA revised the procedures for incorporating by reference federally approved SIPs into the Code of Federal Regulations. These procedural revisions changed the format for the identification of a SIP in 40 CFR part 52, revised the mechanisms for announcing EPA approval of revisions to an applicable SIP, and revised the mechanisms for EPA's updating of both the IBR document (or SIP compilation) and the CFR. The SIP compilations contain the full text of the federally approved materials (including regulations, source-specific permits, and nonregulatory provisions and quasi-regulatory measures) submitted by each State agency, whereas the ''Identification of plan'' sections in 40 CFR part 52 merely identify the submitted materials incorporated by reference into the applicable SIP. Under the revised IBR procedures, EPA periodically publishes an informational document in the rules section of the **Federal Register** when updates are made to a State's SIP compilation. EPA's 1997 revised IBR procedures were formally applied to Arkansas on October 23, 1998 (63 FR 56824).

This action notifies the public of an update to the Arkansas SIP compilation, available for public inspection at the locations listed in the **ADDRESSES** section of this **Federal Register** notice, and updates the Arkansas ''Identification of plan'' section, appearing in 40 CFR part 52 (subpart E). The Arkansas SIP compilation, which consists of submitted materials incorporated by reference into the Arkansas SIP, is being updated to include EPA-approved revisions to Arkansas' SIP that have occurred since EPA's revised IBR procedures were applied to Arkansas on October 23, 1998 (63 FR 56824); specifically, the SIP compilation update includes revisions to Arkansas Regulation 19, Regulations of the Arkansas Plan of Implementation for Air Pollution Control, which we approved on October 16, 2000 (65 FR 61103), and revisions to Arkansas Regulation 26, Regulations of the Arkansas Operating Permit Program, which we approved on October 9, 2001 (66 FR 51312). These revisions have previously undergone notice and comment rulemaking and are, therefore, already in effect as a matter of law; thus this SIP compilation update does not affect the substance of those rulemaking actions nor does it change the rights or obligations of any party.