PART 135—OPERATING REQUIREMENTS: COMMUTER AND ON-DEMAND OPERATIONS AND RULES GOVERNING PERSONS ON BOARD SUCH AIRCRAFT

■ 3. The authority citation for part 135 continues to read as follows:

Authority: 49 U.S.C. 106(g), 44113, 44701–44702, 44705, 44709, 44711–44713, 44715–44717, 44722.

4. In § 135.128, amend paragraph (a)(2)(ii)(D) to read as follows:

§ 135.128 Use of safety belts and child restraint systems

(a) * * *

(2) * * *

(ii) * * *

(D) Except as provided in § 135.128(a)(2)(ii)(C)(*3*) and § 135.128(a)(2)(ii)(C)(*4*), booster-type child restraint systems (as defined in Federal Motor Vehicle Safety Standard No. 213 (49 CFR 571.213)), vest- and harness-type child restraint systems, and lap held child restraints are not approved for use in aircraft; and

* * * * *

Issued in Washington, DC on September 29, 2006.

Brenda D. Courtney,

Acting Director, Office of Rulemaking.

[FR Doc. E6–16622 Filed 10–6–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Omeprazole

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by Merial Ltd. The supplemental NADA provides for administration of omeprazole paste to horses for 8 or 28 days for the prevention of gastric ulcers.

DATES: This rule is effective October 10, 2006.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center for Veterinary Medicine (HFV–110), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7540, e-mail: *melanie.berson@.fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Merial Ltd., 3239 Satellite Blvd., Bldg. 500, Duluth, GA 30096–4640, filed a supplement to NADA 141–227 for ULCERGARD (omeprazole) Paste. The supplemental application provides for administration of omeprazole paste to horses for 8 or 28 days for the prevention of gastric ulcers. The supplemental NADA is approved as of September 15, 2006, and 21 CFR 520.1615 is amended to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

Under section 512(c)(2)(F)(iii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(iii)), this approval qualifies for 3 years of marketing exclusivity beginning September 15, 2006.

The agency has determined under 21 CFR 25.33(d)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

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 in 21 CFR Part 520

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 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

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

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

§ 520.1615 Omeprazole.

■ 2. In paragraph (d)(1)(ii) of § 520.1615, at the end of the first sentence remove ''for up to 28 days'' and add in its place ''for 8 or 28 days''.

Dated: September 27, 2006.

Steven D. Vaughn,

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

[FR Doc. E6–16604 Filed 10–6–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF DEFENSE

Office of the Secretary

[DOD–2006–OS–0134; RIN 0790–AG91]

32 CFR Part 284

Waiver Procedures for Debts Resulting from Erroneous Payments of Pay and Allowances

AGENCY: Department of Defense.

ACTION: Final rule.

SUMMARY: This rule implements policy and prescribes procedures for considering applications for the waiver of debts resulting from erroneous payments of pay and allowances (including travel and transportation allowances) to or on behalf of members of the Uniformed Services and civilian Department of Defense (DoD) employees. The Legislative Branch Appropriations Act of 1996 transferred to the Director of the Office of Management and Budget (OMB) the Comptroller General's authority to settle claims. The OMB Director subsequently delegated some of these authorities to the Department of Defense. Later, the General Accounting Office Act of 1996 codified many of these delegations to the Secretary of Defense and others and transferred to the OMB Director the authority of the Comptroller General to waive uniformed service member and employee debts arising out of the erroneous payment of pay or allowances exceeding $1,500. The OMB Director subsequently delegated the authority to waive such debts of uniformed service members and DoD employees to the Secretary of Defense. The Secretary of Defense further delegated his claims settlement and waiver authorities to the General Counsel. This rule implements the reassignment of the Comptroller General's former duties within the Department of Defense with little impact on the public.

DATES: *Effective Date:* October 10, 2006.

FOR FURTHER INFORMATION CONTACT: Michael Hipple, 703–696–8510.

SUPPLEMENTARY INFORMATION: On Thursday, November 14, 2002 (67 FR 68964), the Department of Defense published 32 CFR part 284 along with parts 281, 282, and 283 as proposed rules with request for public comments.