§ 500.49 [Removed]

4. Section 500.49 *Chlorofluorocarbon propellants* is removed.

PART 505—[REMOVED]

5. Part 505 is removed.

PART 507—[REMOVED]

6. Part 507 is removed.

PART 508—[REMOVED]

7. Part 508 is removed.

PART 510—NEW ANIMAL DRUGS

8. 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).

§ 510.120 [Removed]

9. Section 510.120 *Suspension of approval of new-drug applications for certain diethylstilbestrol and diethylstilbestrol-containing drugs* is removed.

§ 510.200 [Removed]

10. Subpart C, consisting of § 510.200, is removed and reserved.

§ 510.310 [Removed]

11. Section 510.310 *Records and reports for new animal drugs approved before June 20, 1963* is removed.

§ 510.413 [Removed]

12. Section 510.413 *Chloroform used as an ingredient (active or inactive) in animal drug products* is removed.

PART 570—FOOD ADDITIVES

13. The authority citation for 21 CFR part 570 continues to read as follows:

Authority: Secs. 201, 401, 402, 408, 409, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 341, 342, 346a, 348, 371).

§ 570.22 [Removed]

14. Section 570.22 *Safety factors to be considered* is removed.

Dated: July 3, 1996.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 96–18234 Filed 7–18–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Gonadorelin Diacetate Tetrahydrate Injection

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 an abbreviated new animal drug application (ANADA) filed by Intervet, Inc. The ANADA provides for intramuscular and intravenous use of a sterile injectable solution of gonadorelin diacetate tetrahydrate for treating ovarian cysts in female dairy cattle of breeding age.

EFFECTIVE DATE: July 19, 1996.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center for Veterinary Medicine (HFV–135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1643.

SUPPLEMENTARY INFORMATION: Intervet, Inc., 405 State St., P.O. Box 318, Millsboro, DE 19966–0318, filed ANADA 200–134, which provides for intramuscular and intravenous use of Fertagyl® (gonadorelin diacetate tetrahydrate injection) for treatment of ovarian cysts in female dairy cattle of breeding age.

Approval of ANADA 200–134 is as a generic copy of Rhone Merieux's NADA 98–379 for Cystorelin® (gonadorelin diacetate tetrahydrate injection). The ANADA is approved as of June 17, 1996, and the regulations are amended by revising 21 CFR 522.1078(b) to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of part 20 (21 CFR part 20) and § 514.11(e)(2)(ii) (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 Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.24(d)(1)(i) 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.

List of Subjects in 21 CFR Part 522

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 522 is amended as follows:

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

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

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

§ 522.1078 [Amended]

2. Section 522.1078 *Gonadorelin diacetate tetrahydrate injection* is amended in paragraph (b) by removing ''No. 050604'' and adding in its place ''Nos. 050604 and 057926''.

Dated: July 11, 1996.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 96–18350 Filed 7–18–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 801

[Docket No. 95N–310R]

RIN 0910–AA54

Revocation of Certain Device Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is issuing a final rule to remove certain device regulations that are obsolete or no longer necessary to achieve public health goals. These regulations have been identified for revocation as the result of a page-by-page review of the agency's regulations in response to the administration's ''Reinventing Government'' initiative, which seeks to streamline Government and ease the burden on regulated industry and consumers.

EFFECTIVE DATE: August 19, 1996.

FOR FURTHER INFORMATION CONTACT: Joseph M. Sheehan, Center for Devices and Radiological Health (HFZ–215), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–827–2974.

SUPPLEMENTARY INFORMATION:

I. Background

On March 4, 1995, President Clinton announced plans for the reform of the Federal regulatory system as part of the administration's ''Reinventing Government'' initiative. In his March 4, 1995, directive, entitled ''Regulatory Reinvention Initiative,''the President ordered all Federal agencies to conduct a page-by-page review of all of their regulations and to ''eliminate or revise