

requests for records under the Freedom of Information Act, and requests for correction or amendment under the Privacy Act. 5 U.S.C.(d)(3).

Because notice and opportunity for comment are not required pursuant to 5 U.S.C. 553 or any other law, the analytical requirements of the Regulatory Flexibility Act (5 U.S.C. 601 et seq.) are inapplicable. Therefore, a regulatory flexibility analysis is not required and has not been prepared.

**List of Subjects in 15 CFR Part 4**

Freedom of information, Privacy.

■ For the reasons above, amend 15 CFR Part 4 as follows:

**PART 4—DISCLOSURE OF GOVERNMENT INFORMATION**

■ 1. The authority citation for Part 4 continues to read as follows:

**Authority:** 5 U.S.C. 301; 5 U.S.C. 552; 5 U.S.C. 552a; 5 U.S.C. 553; 31 U.S.C. 3717; 44 U.S.C. 3101; Reorganization Plan No. 5 of 1950.

**Appendix B to Part 4—[Amended]**

■ 2. In Appendix B to part 4, under the heading **ECONOMICS AND STATISTICS ADMINISTRATION**, delete “Bureau of the Census: Manager, Freedom of Information Act” and replace with “Bureau of the Census: Freedom of Information Act Officer”.

Dated: June 28, 2007.

**Brenda Dolan,**

*Departmental Freedom of Information and Privacy Act Officer.*

[FR Doc. E7–13001 Filed 7–3–07; 8:45 am]

**BILLING CODE 3510–07–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Parts 510 and 524**

**New Animal Drugs; Change of Sponsor’s Name; Liquid Crystalline Trypsin, Peru Balsam, Castor Oil**

**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 sponsor’s name from Mylan Bertek Pharmaceuticals, Inc., to UDL Laboratories, Inc.

**DATES:** This rule is effective July 5, 2007.

**FOR FURTHER INFORMATION CONTACT:** David R. Newkirk, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6967, e-mail: *david.newkirk@fda.hhs.gov*.

**SUPPLEMENTARY INFORMATION:** Mylan Bertek Pharmaceuticals, Inc., 12720 Dairy Ashford, Sugar Land, TX 77478, has informed FDA that it has changed its name to UDL Laboratories, Inc., and is using a new drug labeler code. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c) to reflect these changes. A conforming change is being made in 21 CFR 524.2620 for this sponsor’s sole product.

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 Part 524*

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 and 524 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. In § 510.600, in the table in paragraph (c)(1), remove the entry for “Mylan Bertek Pharmaceuticals, Inc.” and alphabetically add a new entry for “UDL Laboratories, Inc.”; and in the table in paragraph (c)(2) remove the entry for “062749” and numerically add a new entry for “051079” to read as follows:

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

Firm name and address	Drug labeler code
* * * * *	* * * * *
(c) * * *	* * * * *
(1) * * *	* * * * *
* * * * *	* * * * *

Firm name and address	Drug labeler code
UDL Laboratories, Inc., 12720 Dairy Ashford, Sugar Land, TX 77478.	051079
* * * * *	* * * * *
(2) * * *	* * * * *
Drug labeler code	Firm name and address
* * * * *	* * * * *
051079	UDL Laboratories, Inc., 12720 Dairy Ashford, Sugar Land, TX 77478
* * * * *	* * * * *

**PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS**

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

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

**§ 524.2620 [Amended]**

■ 4. In paragraph (a)(2) of § 524.2620, remove “062794” and add in its place “051079”.

Dated: June 21, 2007.

**Bernadette Dunham,**

*Deputy Director, Center for Veterinary Medicine.*

[FR Doc. E7–13010 Filed 7–3–07; 8:45 am]

**BILLING CODE 4160–01–S**

**DEPARTMENT OF THE INTERIOR**

**Office of Surface Mining Reclamation and Enforcement**

**30 CFR Part 946**

**[VA–123–FOR]**

**Virginia Regulatory Program**

**AGENCY:** Office of Surface Mining Reclamation and Enforcement (OSM), Interior.

**ACTION:** Final rule; approval of amendment.

**SUMMARY:** We are approving an amendment to the Virginia regulatory program under the Surface Mining Control and Reclamation Act of 1977 (SMCRA or the Act). Virginia is revising its remaining regulations to make three of those provisions permanent by deleting a termination date of September 30, 2004, from the regulations. The amendment is intended to render the State regulations consistent with recent amendments to SMCRA.

**EFFECTIVE DATE:** July 5, 2007.