2. Kokoski, C. J., ''Regulatory Food Additive Toxicology,'' in ''Chemical Safety Regulation and Compliance,'' edited by F. Homburger and J. K. Marquis, S. Karger, New York, NY, pp. 24–33, 1985.

3. ''Bioassay of 1,4–Dioxane for Possible Carcinogenicity,'' National Cancer Institute, NCI–CG–TR–80, 1978.

4. Memorandum, Report of the Quantitative Risk Assessment Committee, June 30, 1994.

5. Dunkelberg, H., ''Carcinogenicity of Ethylene Oxide and 1,2–Propylene Oxide Upon Intragastric Administration to Rats,'' *British Journal of Cancer*, 46:924, 1982.

List of Subjects in 21 CFR Part 175

Adhesives, Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 175 is amended as follows:

PART 175—INDIRECT FOOD ADDITIVES: ADHESIVES AND COMPONENTS OF COATINGS

1. The authority citation for 21 CFR part 175 continues to read as follows:

Authority: Secs. 201, 402, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 348, 379e).

2. Section 175.105 is amended in the table in paragraph (c)(5) by alphabetically adding a new entry under the heading ''Substances'' to read as follows:

§ 175.105 Adhesives.

* * * * *

(c) * * *

(5) * * *

Substances	Limitations
* * * * *	
Ethoxylated primary linear alcohols of greater than 10 percent ethylene oxide by weight having molecular weights of 390 to 7,000 (CAS Reg. No. 97953–22–5).	
* * * * *	

Dated: September 1, 1995.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 95–22637 Filed 9–12–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 510

New Animal Drugs; Change of Sponsor Name

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 of approved applications from Animal Sciences Division of Monsanto Co. to Protiva, A Unit of Monsanto Co.

EFFECTIVE DATE: September 13, 1995.

FOR FURTHER INFORMATION CONTACT: Judith M. O'Haro, Center for Veterinary Medicine (HFV–238), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1737.

SUPPLEMENTARY INFORMATION: Animal Sciences Division of Monsanto Co., 800 North Lindbergh Blvd., St. Louis, MO 63167, has informed FDA of a change of sponsor name to Protiva, A Unit of Monsanto Co. Accordingly, FDA is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change of sponsor name.

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 ''Animal Sciences Division of Monsanto Co.'' and by alphabetically adding a new entry for ''Protiva, A Unit of Monsanto Company,'' and in the table in paragraph (c)(2) in the entry for ''059945'' by removing the sponsor name ''Animal Sciences Division of Monsanto Co.'' and adding in its place ''Protiva, A Unit of Monsanto Company.''

Dated: September 1, 1995.

Robert C. Livingston,

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

[FR Doc. 95–22638 Filed 9–12–95; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF TRANSPORTATION

Federal Highway Administration

23 CFR Part 640

[FHWA Docket No. 95–19]

RIN 2125–AD62

Certification Acceptance

AGENCY: Federal Highway Administration (FHWA), DOT.

ACTION: Interim final rule; request for comments.

SUMMARY: The FHWA is adopting an interim policy for certification acceptance (CA) which modifies the current FHWA policy. The interim policy streamlines and simplifies the existing procedures for CA applications to be consistent with the new program provisions in sections such as 1016(f) and 1105(e) of the Intermodal Surface Transportation Efficiency Act of 1991 (ISTEA), Pub. L. 102–240, 105 Stat. 1914. The modifications simplify the current regulations by eliminating unnecessary and prescriptive requirements. The new policy will allow State highway agencies (SHAs) to use the CA alternate procedures to supplement the administrative flexibility provided in the ISTEA for non-Interstate projects.

DATES: This regulation is effective September 13, 1995. Written comments must be received on or before December 12, 1995.

ADDRESSES: Submit written, signed comments to FHWA Docket No. 95–19, Federal Highway Administration, Room 4232, HCC–10, 400 Seventh Street SW., Washington, DC 20590. All comments received will be available for examination at the above address between 8:30 a.m. and 3:30 p.m., e.t., Monday through Friday. Those desiring notification of receipt of comments must include a self-addressed, stamped postcard.

FOR FURTHER INFORMATION CONTACT: Mr. Donald J. Marttila, Interstate and Program Support Branch, Federal-Aid and Design Division, Office of Engineering, (202) 366–4637, or Mr. Wilbert Baccus, Office of the Chief Counsel, (202) 366–0780, Federal Highway Administration, 400 Seventh Street SW., Washington, DC 20590.