

ATTACHMENT 7

**FEDERAL REGISTER NOTICE
DECEMBER 3, 1985, VOLUME 50, P. 49617-49619**

Transaction	Waiting period terminated effective
86-0168—Dress Brothers, Inc.'s proposed acquisition of voting securities of Scott & Myers de Brasil Cigarettes Co. (Grand Metropolitan PLC UFG).	Co.
86-0178—Cott Industries, Inc.'s proposed acquisition of voting securities of Albar, Inc.	Co.
86-0181—Cott Industries, Inc.'s proposed acquisition of voting securities of Albar, Inc.	Co.
86-0188—Proposed consolidation of Inet Communication Services, Inc. and Lental Corporation.	Co.
86-0191—Morvin J. Goodman's proposed acquisition of voting securities of Inet Corp.	Co.
86-0193—Phyllis D. Goodman's proposed acquisition of voting securities of Inet Corp.	Co.
86-0195—Kansas City Southern Industries, Inc.'s proposed acquisition of voting securities of Alax Corp.	Co.
86-0198—Michael P. Richard's proposed acquisition of voting securities of Alax Corp.	Co.
86-0199—Bernard S. LeBow's proposed acquisition of voting securities of the WFA Foods, Inc.	Nov. 14, 1985.
86-0226—Eastern Gas and Fuel Association's proposed acquisition of voting securities of Nicor Mining, Inc. and Carmo Minerals, Inc. (Nico, Inc. UFG).	Co.
86-0227—Niac Inc.'s proposed acquisition of assets of Powderhorn Properties Co. and voting securities of Season Coal Corp.	Co.
86-0157—Hilberbrand Industries, Inc.'s proposed acquisition of voting securities and 25442 of Support Systems International.	Nov. 18, 1985.

For further information contact: Sandra M. Peay, Legal Technician, Merger Notification Office, Bureau of Competition, Room 301, Federal Trade Commission, Washington, DC 20580, (302) 523-3894.

By direction of the Commission.
Emily H. Rock,
Secretary.
[FR Doc. 85-28823 Filed 12-7-85; 8:45 am]
BILLING CODE 8750-01-2

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Notice Regarding Requirement for Submission of List of Ingredients Added to Tobacco in Cigarettes

AGENCY: Department of Health and Human Services (HHS).
ACTION: Notice.

SUMMARY: This notice implements the requirement of the Federal Cigarette Labeling and Advertising Act that each person who manufactures, packages, or imports cigarettes shall annually provide the Secretary of HHS with a list of ingredients added to tobacco in the manufacture of cigarettes.

DATES: The lists are required to be provided to HHS April 2, 1986, and

annually thereafter by December 31, beginning with December 31, 1986.
ADDRESS: The list shall be submitted to: Director, Office on Smoking and Health, Park Building, Room 1-10, 5800 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: Donald R. Shopland, Acting Director, Office on Smoking and Health, (301) 443-1575.

SUPPLEMENTARY INFORMATION: Section 5(a) of Pub. L. 98-474 added a new section 7 to the Federal Cigarette Labeling and Advertising Act. That Section requires manufacturers, importers, and packagers of cigarettes to provide the Secretary of HHS annually with a list of all ingredients added to tobacco in the manufacture of cigarettes. The list shall not identify the company which uses the ingredients or the brand of cigarettes which contain the ingredients.

The list shall be provided reporting each ingredient by chemical name and chemical abstract service (CAS) registry number. A person or group of persons required to provide a list may designate an individual or entity to provide the list on their behalf. In such case, the designated individual or entity shall identify the person or group of persons on whose behalf the list is submitted.

In accordance with section 7, procedures for assuring the confidentiality of the information are available. A copy of these procedures may be obtained by written request to the address stated above. The information submitted will be treated as trade secret or confidential information subject to 5 U.S.C. 552(b)(5) and 18 U.S.C. 1905. Access to the information will be limited to those authorized by the Secretary in carrying out their official duties and, upon their request, to duly authorize committees or subcommittees of the Congress.

Dated November 4, 1985.
James O. Mason,
Acting Assistant Secretary for Health.

Dated November 25, 1985.
Margaret M. Heckler,
Secretary of Health and Human Services.

Guidelines To Control and Protect Documents That Contain Privileged Information Obtained in Accordance With Sec. 5(a) of Pub. L. 98-474

1. Purpose

This guide establishes minimum requirements to control and protect those documents that contain privileged information. Its objective is to establish individual responsibility for the accountability and protection of

privileged information provided to the Secretary, Department of Health and Human Services, specifically that information on the ingredients added to tobacco in the manufacture of cigarettes as called for under Pub. L. 98-474. This document is directed at setting forth specific conditions governing access to privileged information, including trade secret data.

2. Policy

The Department of Health and Human Services recognizes that trust placed in it under the requirements of the Federal Statutes with respect to safeguarding privileged information. Employees of the Department of Health and Human Services shall take such action as may be necessary to preclude a breach of this trust. Privileged information shall be released only to employees of the Department as described herein, unless otherwise authorized by law or by the source of the information. Any Freedom of Information Act request for information obtained by the Department under section 5(a) of Pub. L. 98-474 shall be referred to the Freedom of Information Officer of the Public Health Service. In accordance with the provisions of 5 U.S.C. 552(b)(3) and 552(b)(4), 18 U.S.C. 1905, section 7(b)(2)(A) of Pub. L. 98-474, and 42 CFR 5.71, the Freedom of Information Officer shall deny any such requests. Any request for such information that is not submitted under the Freedom of Information Act shall be referred to the Director of the Office on Smoking and Health. With the exception of duly authorized request by a committee or subcommittee of Congress made in accordance with section 7(b)(2)(B) of Pub. L. 98-474, any such request shall be denied.

3. Statutory Requirements

Statutory requirements for safeguarding privileged information entrusted to the Department of Health and Human Services are contained in the following:

- a. Section 7(b)(2)(A) of the Federal Cigarette Labeling and Advertising Act.
- b. Section 1905, Title 18 U.S.C. Crimes and Criminal Procedure (18 U.S.C. 1905).
- c. Section 552(b)(4), Title 5, U.S.C.

4. Definitions

a. **Document Control Officer.** That individual who has been designated in writing as having the responsibility for the organization's secret document control. The Document Control Officer shall be the Director, Office on Smoking and Health.

Signed: _____
 Date: _____
 Witnessed by: _____
 etc: _____

Attachment B—Receipt for Confidential Information on the Ingredients Added to Tobacco in the Manufacture of Cigarettes

To: Director, Office on Smoking and Health, Office of the Assistant Secretary for Health, Rockville, Maryland 20857

From: Receipt of the following privileged information is hereby acknowledged.

File # _____
 Description of Information _____
 Anticipated Date of Return _____
 Date: _____
 Signature: _____

Attachment C—Authority To Remove Confidential Information on the Ingredients Added to Tobacco in the Manufacture of Cigarettes

_____, (name) of _____ (government agency or office) is hereby granted the authority to have the following privileged information in his/her personal possession from _____ (hours), _____ (date) to _____ (hours), _____ (date).

Described Privileged Information: _____
 Document Number: _____
 Title: _____

This information will be used for:

Authorized by: _____
 Director, Office on Smoking and Health
 Date: _____

[FR Doc. 85-25715 Filed 12-2-85; 8:45 am]
 BILLING CODE 4160-17-M

Food and Drug Administration

Midicel® Tablets; Withdrawal of Approval of NADA

AGENCY: Food and Drug Administration.
 ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of a new animal drug application (NADA) sponsored by Parks-Davis covering use of Midicel® Tablets (sulfamethoxyppyridazine) in treating dogs and cats for sulfasusceptible, gram-positive and gram-negative, bacterial infections. The sponsor requested the withdrawal of approval.

EFFECTIVE DATE: December 13, 1985.

FOR FURTHER INFORMATION CONTACT: John K. Augsburg, Center for Veterinary Medicine (HFV-216), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4093.

SUPPLEMENTARY INFORMATION: Parks-Davis, Division of Warner-Lambert Co., 201 Tabor Rd., Morris Plains, NJ 07950, is sponsor of NADA 12-821 for use of Midicel® Tablets (sulfamethoxyppyridazine) in treating dogs and cats for sulfasusceptible, gram-positive and gram-negative, bacterial infections.

The application was originally approved February 8, 1963. In a letter dated August 7, 1985, the firm requested withdrawal of approval of the NADA because the drug is no longer being marketed.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(e), 82 Stat. 345-347 (21 U.S.C. 360b(e))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Center for Veterinary Medicine (21 CFR 5.84), and in accordance with § 514.115 *Withdrawal of approval of applications* (21 CFR 514.115), notice is given that approval of NADA 12-821 for Midicel® Tablets (sulfamethoxyppyridazine) is hereby withdrawn, effective December 13, 1985.

In a final rule published elsewhere in this issue of the Federal Register, the regulation reflecting this approval is removed.

Date: November 23, 1985.

Gerald B. Guert,
 Acting Director, Center for Veterinary Medicine.
 [FR Doc. 85-28294 Filed 12-2-85; 8:43 am]
 BILLING CODE 4160-01-M

Vortech Pharmaceuticals, Ltd.;
 Dichlorophene and Toluene Capsules;
 Withdrawal of Approval

AGENCY: Food and Drug Administration.
 ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of a new animal drug application (NADA) sponsored by Vortech Pharmaceuticals, Ltd. (formerly sponsored by North American Pharmascel), for dichlorophene and toluene capsules. The firm requested the withdrawal of approval.

EFFECTIVE DATE: December 13, 1985.

FOR FURTHER INFORMATION CONTACT: John Augsburg, Center for Veterinary Medicine (HFV-216), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4093.

SUPPLEMENTARY INFORMATION: Vortech Pharmaceuticals, Ltd., P.O. Box 189, Dearborn, MI 48121, informed FDA, by letter dated July 26, 1983, that it had purchased the assets of North American

Pharmaceutical, 8851 Chass Rd., Dearborn, MI 48128, in June 1982. At the time it was purchased, North American Pharmaceutical was the sponsor of NADA 110-736 for PETAVERM Capsules (dichlorophene and toluene) labeled as an anthelmintic for dogs and cats.

Vortech Pharmaceuticals, Ltd., stated, by letter dated July 8, 1985, that PETAVERM has never been brought to market and is not in use, and the company requested that the drug be withdrawn.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(e), 82 Stat. 345-347 (21 U.S.C. 360b(e))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Center for Veterinary Medicine (21 CFR 5.84) and in accordance with § 514.115 *Withdrawal of approval of applications* (21 CFR 514.115), notice is given that approval of NADA 110-736 and all supplements thereto is hereby withdrawn, effective December 13, 1985.

In a final rule published elsewhere in this issue of the Federal Register, FDA is removing that portion of the regulations that reflects this NADA approval.

Date: November 23, 1985.

Gerald B. Guert,
 Acting Director, Center for Veterinary Medicine.
 [FR Doc. 85-28232 Filed 12-2-85; 8:45 am]
 BILLING CODE 4160-01-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. 1-85-198]

Intended Environmental Impact Statement

The Department of Housing and Urban Development gives notice that an Environmental Impact Statement (EIS) is intended to be prepared by the cities of Auburn Hills and Rochester Hills, Oakland County, Michigan, for the Oakland Technology Park under the HUD programs as described in the appendix of the Notice. This notice is required by the Council on Environmental Quality under its rule (40 CFR Part 1500).

Interested individuals, governmental agencies, and private organizations are invited to submit information and comments concerning the particular project to the specific person or address indicated in the appropriate part of the appendix.

Particularly solicited is information on reports or other environmental studies planned or completed in the project