supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above), between 9 a.m. and 4 p.m., Monday through Friday. This action was considered under FDA's final rule implementing the National Environmental Policy Act (21 CFR Part 25).

List of Subjects in 21 CFR Part 357

Cholecystokinetic drug products, Labeling, Over-the-counter drugs, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and the Administrative Procedure Act, Subchapter D of Chapter I of Title 21 of the Code of Federal Regulations is amended in Part 357 as follows:

PART 357—MISCELLANEOUS INTERNAL DRUG PRODUCTS FOR OVER-THE-COUNTER HUMAN USE

1. The authority citation for 21 CFR Part 357 continues to read as follows:

Authority: Secs. 201(p), 502, 505, 701, 52 Stat. 1041–1042 as amended, 1050–1053 as amended, 1055–1056 as amended by 70 Stat. 919 and 72 Stat. 948 (21 U.S.C. 321(p), 352, 355, 371); 5 U.S.C. 553, 21 CFR 5.10 and 5.11.

2. Section 357.210 is revised to read as follows:

§ 357.210 Cholecystokinetic active ingredients.

The active ingredient of the product consists of any of the following when used within the specified concentration and dosage form established for each ingredient:

(a) 50-percent aqueous emulsion of corn oil.

(b) Hydrogenated soybean oil in a suitable, water-dispersible powder. The hydrogenated soybean oil is food-grade, partially hydrogenated with a melting point of 41 to 43.5 °C, an iodine value of 65 to 69, and a fatty acid composition as follows:

Fatty acid	Percent composition
Myristic acid	0.1
Palmitic acid	10.0
Palmitoleic acid	0.1
Stearic acid	13.5
Oleic acid	72.0
Linoleic acid	3.8
Linolenic acid	0.1
Arachidic acid	0.5
Behenic acid	0.2

3. Section 357.250 is amended by revising paragraphs (d)(2) and (3) to read as follows:

§ 357.250 Labeling of cholecystokinetic drug products.

* * * * *

(d) * * *

(2) *For products containing 50-percent aqueous emulsion of corn oil.*

(i) "Shake well before using."

(ii) Oral dosage is 60 milliliters 20 minutes before diagnostic gallbladder x-ray or as directed by a doctor.

(3) *For products containing hydrogeneated soybean oil.* Oral dosage is 12.4 grams in a suitable, water-dispersible powder in 2 to 3 ounces of water. Stir briskly to prepare a suspension before using. Drink 20 minutes before diagnostic gallbladder x-ray or as directed by a doctor.

* * * * *

4. Section 357.280 is revised to read as follows:

§ 357.280 Professional labeling.

The labeling provided to health professionals (but not to the general public) may contain the following information for ingredients identified in § 357.210: *Indication.* "For visualization of biliary ducts during cholecystography."

Dated: December 23, 1989.

Frank E. Young,

Commissioner of Food and Drugs.

[FR Doc. 89-4613 Filed 2-27-89; 8:45 am]

BILLING CODE 4160-01-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of the Secretary

Office of the Assistant Secretary for Housing—Federal Housing Commissioner

Office of the Assistant Secretary for Community Planning and Development

Office of the Assistant Secretary for Public and Indian Housing

24 CFR Parts 15, 24, 203, 234, 510, 511, 570, 885, 904, 941, and 2002

[Docket No. N-89-1945]

Announcement of Effective Dates

AGENCY: Office of the Secretary; Office of the Assistant Secretary for Housing—Federal Housing Commissioner; Office of the Assistant Secretary for Community Planning and Development; Office of the Assistant Secretary for Public and Indian Housing; HUD.

ACTION: Notice of announcement of effective dates for certain recent final rules.

SUMMARY: Section 7(o)(3) of the Department of Housing and Urban Development Act, 42 U.S.C. 3535(o)(3), requires HUD to wait thirty calendar days of continuous session of Congress, after publication, before it makes a published rule effective. Thirty calendar days of continuous session of Congress have now expired in the present Congress since these rules were published. This notice announces the effective dates for certain recently published final rules. For an explanation of subject matter on the rules, see **"SUPPLEMENTARY INFORMATION".**

DATES: For effective dates, see **"SUPPLEMENTARY INFORMATION."**

FOR FURTHER INFORMATION CONTACT: Grady J. Norris, Assistant General Counsel for Regulations, Department of Housing and Urban Development, Room 10276, 451 Seventh Street, SW., Washington, DC 20410, telephone (202) 755-7055. (This is not a toll-free number).

SUPPLEMENTARY INFORMATION: The effective date provision of each of the published rules affected by this Notice stated that the rule would become effective upon expiration of the first period of 30 calendar days of continuous session of Congress after publication, and announced that future notice of the rule's effectiveness would be published in the **Federal Register.** Thirty calendar days of continuous session of Congress will have expired in the present Congress before March 3, 1989.

Accordingly, the purpose of this notice is to announce the effective dates for the rules listed below, as follows:

1. *24 CFR Parts 15 and 2002:* The Freedom of Information Reform Act of 1988; Fee Schedule and Fee Waiver Regulations, Final Rule published September 27, 1988 (53 FR 37546), Docket No. R-88-1348; FR-2362.

DATE: Effective Date: March 3, 1989.

2. *24 CFR Part 24:* Debarment Suspension and Limited Denial of Participation, Contractors and Participants, Final Rule published November 15, 1988 (53 FR 45903), Docket No. R-88-831; FR-1676.

DATE: Effective Date: March 3, 1989.

3. *24 CFR Parts 203 and 234:* Disclosure of Annual Rate Changes of Adjustable Rate Mortgages (ARMs) and Carryovers, Final Rule published January 4, 1989 (54 FR 110), Docket No. R-88-1427; FR-2542.

DATE: Effective Date: March 31, 1989.

4. *24 CFR Part 510:* Section 312 Rehabilitation Loan Program; Removal of Risk Premium and Application Fee Provisions, Final Rule published