District	Customs stations	Port of entry having supervision
Great Falls, Mont....	Colorado Springs, Colo.	Denver.
	Wild Horse, Mont....	Great Falls.
	Willow Creek, Mont.	Do.

Michael H. Lane,
Acting Commissioner of Customs.

Approved:
Francis A. Keating, II,
Assistant Secretary of the Treasury.
February 25, 1987.
[FR Doc. 87-4890 Filed 3-6-87; 8:45 am]
BILLING CODE 4820-02-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 341

[Docket No. 76N-052B]

Bronchodilator Drug Products for Over-the-Counter Human Use; Final Monograph; OMB Approval of Requirements

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the Office of Management and Budget (OMB) has approved the collection of information requirement concerning its final rule on over-the-counter (OTC) bronchodilator drug products. The agency is amending that regulation to reflect OMB's approval.

EFFECTIVE DATE: October 2, 1987.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Center for Drugs and Biologics (HFN-210), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-295-8000.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 2, 1986 (51 FR 35326), FDA issued a final rule in the form of a final monograph effective October 2, 1987, establishing conditions under which OTC bronchodilator drug products (drug products used in the symptomatic treatment of wheezing and shortness of breath of asthma) are generally recognized as safe and effective and not misbranded. In that document (51 FR 35339), FDA announced that it had submitted the final rule to the Office of Management and Budget (OMB) for approval of the collection of information requirement contained in § 341.76(d)(2)(i)(b).

OMB has approved the collection of information requirement under OMB control number 0910-0237. This document announces OMB's approval and amends the regulation to reflect that approval.

Because this amendment is nonsubstantive, notice and public procedure are unnecessary (5 U.S.C. 553(b)(B) and (d)).

List of Subjects in 21 CFR Part 341

Labeling, Over-the-counter drugs.

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 as follows:

PART 341—COLD, COUGH, ALLERGY, BRONCHODILATOR, AND ANTIASTHMATIC DRUG PRODUCTS FOR OVER-THE-COUNTER HUMAN USE

1. The authority citation for 21 CFR Part 341 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 5.11.

2. In § 341.76 by adding a parenthetical statement at the end of the section, to read as follows:

§ 341.76 Labeling of bronchodilator drug products.

* * * * *

(Collection of information requirement approved by the Office of Management and Budget under number 0910-0237.)

Dated: March 2, 1987.

John M. Taylor,
Associate Commissioner for Regulatory Affairs.

[FR Doc. 87-4818 Filed 3-6-87; 8:45 am]
BILLING CODE 4160-01-M

DEPARTMENT OF TRANSPORTATION

Federal Highway Administration

23 CFR Part 655

[FHWA Docket Nos. 79-37, 83-26, 85-1, 85-3, 85-15, 85-27 and 87-8]

National Standards for Traffic Control Devices; Manual on Uniform Traffic Control Devices

AGENCY: Federal Highway Administration (FHWA), DOT.

ACTION: Final rule; amendments to material incorporated by reference and request for comments.

SUMMARY: This document contains notice of amendments to the Manual on Uniform Traffic Control Devices (MUTCD) which are being adopted by the Federal Highway Administration for inclusion therein. The MUTCD is incorporated by reference in 23 CFR Part 655, Subpart F and recognized as the national standard for traffic control devices on all public roads. The amendments affect various parts of the MUTCD and are intended to expedite traffic, improve safety and provide a more uniform application of highway signs, signals, and markings. Comments on certain editorial amendments to the MUTCD, specified in SUPPLEMENTARY INFORMATION under "Discussion of Editorial Changes", are requested. A conforming amendment to Part 655 is also being made to update the citation to the MUTCD.

DATES: Effective March 9, 1987. Comments on the editorial amendments to the MUTCD must be received on or before April 3, 1987.

ADDRESS: Submit written comments, preferably in triplicate, on editorial changes to Docket No. 87-8, to the Federal Highway Administration, Room 4205, 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. ET, Monday through Friday. Those desiring notification of receipt of comments must include a self-addressed, stamped postcard.

FOR FURTHER INFORMATION CONTACT: Mr. Philip O. Russell, Office of Traffic Operations, (202) 366-2184, or Mr. Michael J. Laska, Office of Chief Counsel, (202) 366-1383, 400 Seventh Street, SW., Washington, DC 20590. Office hours are from 7:45 a.m. to 4:15 p.m. ET, Monday through Friday, except legal holidays.

SUPPLEMENTARY INFORMATION: The MUTCD is available for inspection and copying as prescribed in 49 CFR Part 7, Appendix D. It may be purchased for $44.00 from the Superintendent of Documents, U.S. Government Printing Office, Washington, DC 20402, Stock No. 950-036-00000-1. The purchase of a MUTCD includes a subscription service for adopted revisions.

This document contains the dispositions of proposals for changes in the MUTCD which were received or originated by the FHWA. Previous Federal Register actions regarding these requests are listed in the following table: