THE PROPOSAL

The FAA is considering an amendment to Subpart G of Part 71 of the Federal Aviation Regulations (14 CFR Part 71) to alter the Riverside, California 700 foot transition area. This action will provide additional controlled airspace for radar vector service to be provided by Ontario Approach Control.

Accordingly, the Federal Aviation Administration proposes to amend § 71.181 of Part 71 of the Federal Aviation Regulations (14 CFR Part 71) as follows:

71.181 (AMENDED) RIVERSIDE, CALIF.

Delete all between latitude 33°38'00" N., longitude 117°09'00" W., and to latitude 33°46'00" N., longitude 117°45'00" W., and substitute therein:

to latitude 33°43'00" N., longitude 117°15'00" W.,
to latitude 33°43'00" N., longitude 117°25'00" W.,
to latitude 33°39'00" N., longitude 117°25'00" W.,
to latitude 33°39'00" N., longitude 117°30'00" W.

(Sec. 307 (a), Federal Aviation Act of 1958, as amended (49 U.S.C. 1348 (a)); sec. 6 (c) of the Department of Transportation Act (49 U.S.C. 1655 (c)); 14 CFR 11.65)

NOTE.—The Federal Aviation Administration has determined that this document does not contain a major proposal requiring preparation of an Economic Impact Statement under Executive Order 11821, as amended by Executive Order 11949, and OMB Circular A-107.

Issued in Los Angeles, Calif., on June 12, 1978.

LEON C. DAUGHERTY,
Deputy Director,
Western Region.

[FR Doc. 78-17526 Filed 6-23-78; 8:45 am]

[6320-01]

CIVIL AERONAUTICS BOARD

[14 CFR Part 371]

[Docket 32242]

ADVANCE BOOKING CHARTERS

Proposed Rulemaking To Liberalize Charter Rules

JUNE 20, 1978.

AGENCY: Civil Aeronautics Board.

ACTION: Notice of oral argument.

SUMMARY: On March 17, 1978, the Board issued a notice (EDR-348, SPDR-64, 43 FR 11215) proposing to replace most of the existing charter forms with a simplified form known as a "public charter." Comments were requested by April 26, 1978, with replying comments due May 16, 1978. On March 24 and June 1, 1978, the Board issued notices (43 FR 24542, 43 FR 25141) that oral argument on this proceeding would be held before the Board on June 30, 1978.

DATES: Oral argument is scheduled for June 30, 1978, in Room 1027, Universal Building, 1825 Connecticut Avenue NW., Washington, D.C. Because of the number of persons wishing to participate, the time the argument will begin has been changed to 9:30 a.m. (local time).

FOR FURTHER INFORMATION CONTACT:

Richard B. Dyson, Civil Aeronautics Board, Office of the General Counsel, 1825 Connecticut Avenue NW., Washington, D.C. 20428, 202-673-5444.

SUPPLEMENTARY INFORMATION: The argument will be structured to facilitate give and take among the participants, and between participants and Board members. Participants will be organized into four panels, so that after presentation of brief opening statements they will be conveniently situated to ask and answer questions. The four panels are:

PANEL 1

Eastern Airlines, Inc., Northwest Airlines, Inc., Trans World Airlines, Inc., Western Airlines, Inc.[1]
Department of Justice
Air Charter Tour Operators of America, Spantax, S.A., Davis Agency, Inc., Arthurs Travel Center, Inc.[1]
Pan American World Airways, Inc.
Trans International Airlines

PANEL 2

American Airlines
Suntour, Ltd., and Hamilton, Miller, Hudson & Fayne Travel Corp.[1]
American Society of Travel Agents
Capitol International Airways

PANEL 3

National Airlines
Department of Transportation
Evergreen International Airlines
Las Vegas Parties

PANEL 4

British Airways
World Airways
Council of International Educational Exchange
International Weekends
Bureau of Competition of the Federal Trade Commission

Panels 1 and 4, each of which has five participants, will be allotted a total of 95 minutes; panels 2 and 3, each of which has four participants, will be allotted a total of 75 minutes. Each panel participant will have 8 minutes for an opening statement; the rest of the time will be devoted to questions by the Board, and to answers to those questions and to each other's statements and comments.

The following is the approximate time schedule:

9:30-10:10 a.m. Opening statements by participants of Panel 1.
10:10-11:05 a.m. Question and answer period.
11:05-11:15 a.m. Break.
11:15-11:50 a.m. Opening statements by participants of Panel 2.
11:50-12:30 p.m. Question and answer period.
12:30-1:30 p.m. Lunch break.
1:30-2:05 p.m. Opening statements by participants of Panel 3.
2:05-2:45 p.m. Question and answer period.
2:45-2:55 p.m. Break.
2:55-3:35 p.m. Opening statements by participants of Panel 4.
3:35-4:30 p.m. Question and answer period.
4:30 Adjourn.

By the Civil Aeronautics Board.

PHYLLIS T. KAYLOR,
Secretary.

[FR Doc. 78-17676 Filed 6-23-78; 8:45 am]

[1] To be jointly represented by single counsel.

[4410-09]

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[21 CFR Part 1308]

SCHEDULES OF CONTROLLED SUBSTANCES

Proposed Placement Into Schedules IV and V of Preparations Containing Difenoxin in Combination with Atropine Sulfate

AGENCY: Drug Enforcement Administration.

ACTION: Notice of proposed rulemaking.

SUMMARY: This is a notice of proposed rulemaking issued by the Administrator of DEA to place two preparations, one combining 1.0 mg. difenoxin with 0.025 mg. atropine sulfate, and the other combining 0.5 mg. difenoxin with 0.025 mg. atropine sulfate into Schedules IV and V respectively, of the Controlled Substances Act.

The effect of the present proposal would be to require that the manufacture, distribution, dispensing, importation and exportation of the 1 mg. difenoxin-atropine sulfate preparation be subject to controls for Schedule IV narcotic substances and that the manufacture, distribution, dispensing, importation and exportation of the 0.5 mg. difenoxin-atropine sulfate preparation be subject to controls for Schedule V narcotic drugs. This action does not affect the Schedule I status of any other form or preparation of difenoxin.

DATES: Comments should be received on or before July 26, 1978.

ADDRESS: Send comments in quintuplicate to: Administrator, Drug Enforcement Administration, U.S. Department of Justice, 1405 I Street NW., Washington, D.C. 20537. Attention: DEA Federal Register Representative.

FOR FURTHER INFORMATION CONTACT:

Howard McClain, Jr., Chief, Regula-

tory Control Division, Drug Enforcement Administration, telephone 202-633-1366.

SUPPLEMENTARY INFORMATION: On January 23, 1976, the Assistant Secretary for Health, Department of Health, Education, and Welfare sent a letter to the then Acting Administrator of the Drug Enforcement Administration which recommended that several drugs be placed into, transferred between or removed from certain schedules of the Controlled Substances Act (21 U.S.C. 801-966). The following recommendations were among those contained in the letter:

"(8) *Difenoxine and Difenoxine-Atropine*—The FDA is currently reviewing a New Drug Application for a difenoxine-atropine drug product as an antidiarrheal agent.

Difenoxine is currently a Schedule I substance under the CSA.

Data on difenoxine and the combination product containing atropine were reviewed by the FDA Controlled Substances Advisory Committee and FDA staff.

Based on this review we recommend that, should a New Drug Application be approved, difenoxine be controlled in the same manner as is diphenoxylate.

That is, we recommend that difenoxine be controlled in Schedule II of the CSA, and that the marketed formulation containing atropine in combination with difenoxine be controlled in Schedule V of the CSA."

As to the single entity narcotic difenoxin, the Administrator defers initiating rulemaking proceedings on that substance until the required certifications are made by the Assistant Secretary that difenoxin has a currently accepted medical use. However, a New Drug Application is pending issuance to permit marketing of two formulations of difenoxin with atropine sulfate: one combines 0.5 mg. difenoxin with 0.025 mg. atropine sulfate; the second formulation combines 1 mg. difenoxin with 0.025 mg. atropine sulfate.

To assist the Administrator in initiating rulemaking proceedings, the Assistant Secretary enclosed with his letter his summarized considerations. These were fully considered by the Administrator.

In addition, and paramount to these considerations, the Administrator evaluated the current drug control obligations of the United States under the United Nations Single Convention on Narcotic Drugs, 1961, as amended, in an effort to carry out the Assistant Secretary's recommendations and remain consistent with the requirements of the Single Convention.

Section 201(d) of the Act (21 U.S.C. 811(d)) authorizes the Administrator to consider the United States international drug control obligations, and to consequently determine the most appropriate schedule for the difenoxin-atropine sulfate preparations recommended for control by the Assistant Secretary. The Administrator finds that all forms of difenoxin are listed in Schedule I of the Single Convention except preparations of difenoxin which contain per dosage unit, not more than 0.5 milligrams of difenoxin and a quantity of atropine sulfate equivalent to at least 5 percent of the dose of difenoxin. Such preparations are in Schedule III of the Single Convention. The Administrator also finds that preparations other than those in Schedule III of the Single Convention are subject to the same measures of control as Single Convention Schedule I substances except that estimates (article 19) and statistics (article 20) distinct from those dealing with difenoxin are not required, and article 29, paragraph 2(c) (quotas on the manufacture of the preparations) and article 30, paragraph 1(b)(ii) (registration of distributors) need not apply.

In consideration of the Assistant Secretary's recommendation, and based upon the single convention, the Controlled Substances Act, and the Controlled Substances Import and Export Act (21 U.S.C. 951-966), the Administrator has determined that Schedule IV of the Controlled Substances Act is the most appropriate schedule for the 1 mg. difenoxin-atropine sulfate preparation and that Schedule V of that Act is the most appropriate schedule for the 0.5 mg. difenoxin-atropine sulfate preparation.

Therefore, pursuant to section 201(d) of the Controlled Substances Act (21 U.S.C. 811(d)) and the regulations of the Drug Enforcement Administration and of the Department of Justice, the Administrator of the Drug Enforcement Administration proposes that upon approval by the Food and Drug Administration of the New Drug Application for Motofen and Motofen Half-Strength, §§ 1308.14 and 1308.15 of Title 21, Code of Federal Regulations be amended to read as follows:

§1308.14 Schedule IV.

* * * * *

(b) *Narcotic drugs.* Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing limited quantities of any of the following narcotic drugs, or any salts thereof:

(1) Not more than 1 milligram of difenoxin and not less than 25 micrograms of atropine sulfate per dosage unit.

(c) *Depressants* * * *
(d) *Fenfluramine* * * *
(e) *Stimulants* * * *
(f) *Other Substances* * * *

§ 1308.15 Schedule V.

* * * * *

(b) *Narcotic drugs containing non-narcotic active medicinal ingredients.* Any compound, mixture, or preparation containing any of the following limited quantities of narcotic drugs or salts thereof, which shall include one or more non-narcotic active medicinal ingredients in sufficient proportion to confer upon the compound, mixture, or preparation valuable medicinal qualities other than those possessed by the narcotic alone:

* * * * *

(6) Not more than 0.5 milligram of difenoxin and not less than 25 micrograms of atropine sulfate per dosage unit.

* * * * *

All interested persons are invited to submit their comments in writing regarding this proposal. These comments should state with particularity the issues concerning which the person desires to be heard. Comments should be submitted in quintuplicate to the Administrator, Drug Enforcement Administration, United States Department of Justice, 1405 I Street, NW., Washington, D.C. 20537, Attention: DEA Federal Register Representative.

Dated: June 12, 1978.

PETER B. BENSINGER,
Administrator,
Drug Enforcement Administration.

[FR Doc. 78-17602 Filed 6-23-78; 8:45 am]

[4210-01]

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Federal Insurance Administration

[24 CFR Part 1917]

[Docket No. FI-3176]

NATIONAL FLOOD INSURANCE PROGRAM

Proposed Flood Elevation Determinations for the City of Milford, New Haven County, Connecticut; Correction

AGENCY: Federal Insurance Administration, HUD.

ACTION: Correction of proposed rule.

SUMMARY: This document corrects a proposed rule on base (100-year) flood elevations that appeared on page 38526 of the FEDERAL REGISTER of July 28, 1977.

EFFECTIVE DATE: July 28, 1977.

FOR FURTHER INFORMATION CONTACT:

Mr. Richard Krimm, Assistant Administrator, Office of Flood Insurance, Room 5270, 451 Seventh Street, SW., Washington, D.C. 20410, (202) 755-5581 or Toll Free Line (800) 424-8872.

The following additions are to be made: