Ala., to the route structure. This realinement will provide pilots with improved navigational guidance.

EFFECTIVE DATE: November 2, 1978.

FOR FURTHER INFORMATION CONTACT:

Mr. Lewis W. Still, Airspace Regulations Branch (AAT-230), Airspace and Air Traffic Rules Division, Air Traffic Service, Federal Aviation Administration, 800 Independence Avenue SW., Washington, D.C. 20591, telephone 202-426-8525.

SUPPLEMENTARY INFORMATION: The purpose of this amendment to subpart B of part 75 of the Federal Aviation Regulations (14 CFR Part 75) is to realign a portion of J-2 between New Orleans, La., and Montgomery, Ala., so that the intersection in the present description is replaced by the Semmes VORTAC which is at the same geographical location. Presently en route aircraft have a tendency to stray north of the course when proceeding inbound to Crestview, Fla., VORTAC. This creates an additional workload for controllers who must monitor each flight to insure against straying east of the course centerline. By designating Semmes VORTAC in the route alinement, in lieu of the intersection, pilots will be able to remain on the centerline of J-2. There is no change in the currently designated airspace. Section 75.100 was republished in the FEDERAL REGISTER on January 3, 1978 (43 FR 714).

Under the circumstances presented, the FAA concludes that there is an immediate need for the safety of flight and security benefits of this minor modification to the airspace designation in the affected area. In order to affect that action before the next aeronautical charting date on November 2, 1978, it is necessary to immediately adopt this regulation change; accordingly, I find good cause that notice and public procedure thereon is impracticable and unnecessary.

ADOPTION OF THE AMENDMENT

Accordingly, pursuant to the authority delegated to me by the Administrator, § 75.100 of the Federal Aviation Regulations (14 CFR Part 75) as republished (43 FR 714) is amended, effective 0901 G.m.t., November 2, 1978, as follows:

Under jet route No. 2, "New Orleans; INT of the New Orleans 066° and the Crestview, Fla., 266° radials; Crestview;" is deleted and "New Orleans; Semmes, Ala.; Crestview, Fla.;" is substituted therefor.

(Secs. 307(a), 313(a), Federal Aviation Act of 1958 (49 U.S.C. 1348(a), 1354(a)); sec. 6(c), Department of Transportation Act (49 U.S.C. 1655(c)); and 14 CFR 11.69.)

NOTE.—The FAA has determined that this document involves a proposed regulation which is not considered to be significant under the procedures and criteria prescribed by Executive Order 12044 and as implemented by interim Department of Transportation guidelines (43 FR 9582; Mar. 8, 1978).

Issued in Washington, D.C., on September 15, 1978.

WRAY R. MCCLUNG,
Acting Chief, Airspace and Air Traffic Rules Division.

[FR Doc. 78-26626 Filed 9-22-78; 8:45 am]

[4410-09]

Title 21—Food and Drugs

CHAPTER II—DRUG ENFORCEMENT ADMINISTRATION, DEPARTMENT OF JUSTICE

PART 1308—SCHEDULES OF CONTROLLED SUBSTANCES

Placement of N-ethyl-1-phenylcyclohexylamine and 1-(1-phenylcyclohexyl)pyrrolidine into Schedule I

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Final Rule.

SUMMARY: This rule requires that the manufacture, distribution, dispensing, importation and exportation of PCE, which is N-ethyl-1-phenyl-cyclohexylamine, and PHP, which is 1-(1-phenylcyclohexyl)pyrrolidine, be subject to the regulations applicable to substances in schedule I of the Controlled Substances Act. This rule results from a recommendation of the Assistant Secretary for Health, Department of Health, Education, and Welfare, on behalf of the Secretary that these substances be placed into schedule I, review thereof by the Drug Enforcement Administration (EDEA) and subsequent publication in the FEDERAL REGISTER (43 FR 35743, Aug. 11, 1978), of a notice of proposed rulemaking to place PCE, PHP, and salts, isomers and salts of isomers thereof into schedule I. No comments or objections were received in response to the notice.

EFFECTIVE DATE: The effective date of schedule I control is October 25, 1978, except as otherwise provided in the Supplementary Information section of this order.

FOR FURTHER INFORMATION CONTACT:

Howard McClain, Jr., Chief, Regulatory Control Division, Drug Enforcement Administration, telephone 202-633-1366.

SUPPLEMENTARY INFORMATION: A notice was published in the FEDERAL REGISTER on August 11, 1978 (43 FR 35734) proposing that any material, compound, mixture, or preparation which contains any quantity of N-ethyl-1-phenylcyclohexylamine or 1-(1-phenylcyclohexyl)pyrrolidine or which contains any of their salts, isomers, or salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation (the term "isomer" includes optical, position and geometric isomers) be placed into schedule I of the Comprehensive Drug Abuse Prevention and Control Act of 1970 (21 U.S.C. 801-966) and that title 21 of the Code of Federal Regulations, § 1308.11 (schedule D be amended accordingly. All interested persons were given until September 11, 1978 to submit their comments or objections in writing regarding this proposal.

No comments or objections were received, nor were there any requests for a hearing. In view thereof, and based upon the investigations and review conducted by the Drug Enforcement Administration and upon the recommendation of the Assistant Secretary for Health on behalf of the Secretary of Health, Education, and Welfare, the Administrator of the Drug Enforcement Administration finds, pursuant to the authority delegated to him by regulations of the Department of Justice, that:

1. N-ethyl-1-phenylcyclohexylamine and 1-(1-phenylcyclohexyl)pyrrolidine each has a high potential for abuse;

2. Neither substance has a currently accepted medical use in treatment in the United States;

3. N-ethyl-1-phenylcyclohexylamine and 1-(1-phenylcyclohexyl)pyrrolidine each lacks accepted safety for use under medical supervision.

Therefore, under the authority vested in him by the Act and by regulations of the Department of Justice, the Administrator of the Drug Enforcement Administration hereby orders that § 1308.11(d) of title 21 of the Code of Federal Regulations be amended to read as follows:

§ 1308.11 Schedule I.

* * * * *

(d) *Hallucinogenic substances.* * * *

(21) Ethylamine analog of phencyclidine	7455
Some trade or other names: N-ethyl-1-phenylcyclohexylamine, (1-phenylcyclohexyl)ethylamine, N-(1-phenylcyclohexyl)ethylamine, cyclohexamine, PCE.	
(22) Pyrrolidine analog of phencyclidine	7458
Some trade or other names: 1-(1-phenylcyclohexyl)-pyrrolidine, PCPy, PHP.	

(23) Thiophene analog of phencyclidine	7470
Some trade or other names: 1-[1-(2-thienyl)-cyclohexyl]-piperidine, 2-thienylanalog of phencyclidine, TPCP, TCP.	

* * * * *

EFFECTIVE DATES

As to N-ethyl-1-phenyl-cyclohexylamine and 1-(1-phenylcyclohexyl)-pyrrolidine:

1. *Registration.* Any person who manufactures, distributes, dispenses, imports or exports such substances or who proposes to engage in such activities, shall submit an application for registration to conduct such activities in accordance with parts 1301 and 1311 of title 21 of the Code of Federal Regulations on or before October 25, 1978;

2. *Security.* Such substances must be manufactured, distributed, and stored in accordance with §§ 1301.71, 1301.72 (a), (c), and (d), 1301.73, 1301.74(a)-(f), 1301.75(a), and 1301.76 of title 21 of the Code of Federal Regulations on or before November 24, 1978. From now until the effective date of this provision, it is expected that manufacturers and distributors of such substances will initiate whatever preparation as may be necessary in order to provide adequate security in accordance with DEA regulations so that substantial compliance with this provision can be met by November 24, 1978. In the event that this imposes special hardships, the Drug Enforcement Administration will entertain any justified requests for extensions of time;

3. *Labeling and packaging.* All labels on commercial containers of, and all labeling of such substances packaged after November 24, 1978, shall comply with the requirements of §§ 1302.03-1302.05, 1302.07, and 1302.08 of title 21 of the Code of Federal Regulations. In the event this effective date imposes special hardships on any manufacturer, as defined in section 102(14) of the Controlled Substances Act (21 U.S.C. 802(14)), the Drug Enforcement Administration will entertain any justified requests for an extension of time;

4. *Quotas.* All persons required to obtain quotas with respect to either of such substances shall submit applications pursuant to §§ 1303.12 and 1303.22 of title 21 of the Code of Federal Regulations on or before November 24, 1978.

5. *Inventory.* Every registrant required to keep records who possesses any quantity of such substances shall take an inventory pursuant to §§ 1304.11-1304.19 of title 21 of the Code of Federal Regulations, of all stocks of such substances on hand on November 24, 1978;

6. *Records.* All registrants required to keep records pursuant to §§ 1304.21-1304.27 of title 21 of the Code of Federal Regulations shall do so regarding such substances commencing on the date on which the inventory of such substances is taken;

7. *Reports.* All registrants required to submit reports pursuant to §§ 1304.37-1304.41 of title 21 of the Code of Federal Regulations shall do so regarding such substances commencing on the date on which the inventory of such substances is taken;

8. *Order forms.* The order form requirements of §§ 1305.01-1305.16 of title 21 of the Code of Federal Regulations shall be in effect on the date which the initial inventory of these schedule I controlled substances is taken;

9. *Importation and exportation.* All importation and exportation of such substances shall, on or after November 24, 1978, be required to be in compliance with part 1312 of title 21 of the Code of Federal Regulations;

10. *Criminal liability.* The Administrator, Drug Enforcement Administration, hereby orders that any activity with respect to N-ethyl-1-phenylcyclohexylamine and 1-(1-phenylcyclohexyl)pyrrolidine, not authorized by or in violation of the Controlled Substances Act or the Controlled Substances Import and Export Act, conducted after October 25, 1978, shall be unlawful, except that any person who is entitled to registration under such acts may continue to conduct normal business, research or professional practice with such substances between the date on which this order is published and the date on which he obtains or is denied registration: *Provided,* That application for such registration is submitted on or before October 25, 1978;

11. *Other.* In all other respects, this order is effective October 25, 1978.

Dated: September 19, 1978.

PETER B. BENSINGER,
Administrator, Drug Enforcement Administration.

[FR Doc. 78-26835 Filed 9-22-78; 8:45 am]

[4410-01]

Title 28—Judicial Administration

CHAPTER I—DEPARTMENT OF JUSTICE

[Order No. 802-78]

PART 0—ORGANIZATION OF THE DEPARTMENT OF JUSTICE

Service in Customs Litigation

AGENCY: Department of Justice.

ACTION: Final rule.

SUMMARY: This order amends the Department of Justice organizational regulations to reflect the Civil Division reorganization by changing the official designated to receive service in customs litigation. The official is changed from the Chief of the Customs Service, to the Attorney-in-Charge of the Field Office for Customs Litigation.

EFFECTIVE DATE: September 5, 1978.

FOR FURTHER INFORMATION CONTACT:

Barbara Allen Babcock, Assistant Attorney General, Civil Division, U.S. Department of Justice, Washington, D.C. 20530, 202-739-3301.

By virtue of the authority vested in me by 28 U.S.C. 509, 510, and 516, and 5 U.S.C. 301, section 0.48 of subpart I of part 0 of chapter I of title 28, Code of Federal Regulations, is amended by inserting "The Attorney-in-Charge, Field Office for Customs Litigation" for "The Chief, Customs Section."

Dated: September 14, 1978.

GRIFFIN B. BELL,
Attorney General.

[FR Doc. 78-26858 Filed 9-22-78; 8:45 am]

[4410-01]

[Order No. 800-78]

PART 0—ORGANIZATION OF THE DEPARTMENT OF JUSTICE

PART 21—WITNESS FEES

Technical Amendments

AGENCY: Department of Justice.

ACTION: Final rule.

SUMMARY: The regulations governing the disposition of unclaimed property in the Department of Justice are presently published in § 0.76(o), and the appendix to subpart O, part 0 of title 28, Code of Federal Regulations. Those regulations are being replaced by revised regulations to be issued by the Assistant Attorney General for Administration, and will be codified in part 128-48 of title 41, Code of Federal Regulations. This order, therefore, revokes § 0.76(o), and the appendix to subpart O. The order also updates references to the Federal Travel Regulations in 28 CFR 0.76, 0.142 and 21.1.

EFFECTIVE DATE: August 30, 1978.

FOR FURTHER INFORMATION CONTACT:

Vincent A. Lobisco, Chief, Administrative Programs Group, Administrative Programs Management Staff, Office of Management and Finance, Department of Justice, Washington, D.C. 20530, 202-739-2971.

By virtue of the authority vested in me by 28 U.S.C. 509, 510, and 5 U.S.C. 301, chapter I of title 28, Code of Federal Regulations, is amended as follows: