

Public Law 95-633  
95th Congress

An Act

Nov. 10, 1978  
[S. 2399]

To amend the Comprehensive Drug Abuse Prevention and Control Act of 1970 and other laws to meet obligations under the Convention on Psychotropic Substances relating to regulatory controls on the manufacture, distribution, importation, and exportation of psychotropic substances, and for other purposes.

Psychotropic  
Substances Act  
of 1978.  
21 USC 801 note.

*Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,* That this Act may be cited as the "Psychotropic Substances Act of 1978".

TITLE I—ENABLING PROVISIONS FOR THE CONVENTION ON PSYCHOTROPIC SUBSTANCES

21 USC 801a.

SEC. 101. The Congress makes the following findings and declarations:

(1) The Congress has long recognized the danger involved in the manufacture, distribution, and use of certain psychotropic substances for nonscientific and nonmedical purposes, and has provided strong and effective legislation to control illicit trafficking and to regulate legitimate uses of psychotropic substances in this country. Abuse of psychotropic substances has become a phenomenon common to many countries, however, and is not confined to national borders. It is, therefore, essential that the United States cooperate with other nations in establishing effective controls over international traffic in such substances.

(2) The United States has joined with other countries in executing an international treaty, entitled the Convention on Psychotropic Substances and signed at Vienna, Austria, on February 21, 1971, which is designed to establish suitable controls over the manufacture, distribution, transfer, and use of certain psychotropic substances. The Convention is not self-executing, and the obligations of the United States thereunder may only be performed pursuant to appropriate legislation. It is the intent of the Congress that the amendments made by this Act, together with existing law, will enable the United States to meet all of its obligations under the Convention and that no further legislation will be necessary for that purpose.

(3) In implementing the Convention on Psychotropic Substances, the Congress intends that, consistent with the obligations of the United States under the Convention, control of psychotropic substances in the United States should be accomplished within the framework of the procedures and criteria for classification of substances provided in the Comprehensive Drug Abuse Prevention and Control Act of 1970. This will insure that (A) the availability of psychotropic substances to manufacturers, distributors, dispensers, and researchers for useful and legitimate medical and scientific purposes will not be unduly restricted; (B) nothing in the Convention will interfere with bona fide research activities; and (C) nothing in the Convention will interfere with ethical medical practice in this country as determined by the Sec-

21 USC 801 note.

retary of Health, Education, and Welfare on the basis of a consensus of the views of the American medical and scientific community.

Sec. 102. (a) Subsection (d) of section 201 of the Controlled Substances Act (21 U.S.C. 811(d)) is amended by inserting "(1)" after "(d)" and by adding the following new paragraphs at the end thereof:

"(2) (A) Whenever the Secretary of State receives notification from the Secretary-General of the United Nations that information has been transmitted by or to the World Health Organization, pursuant to article 2 of the Convention on Psychotropic Substances, which may justify adding a drug or other substance to one of the schedules of the Convention, transferring a drug or substance from one schedule to another, or deleting it from the schedules, the Secretary of State shall immediately transmit the notice to the Secretary of Health, Education, and Welfare who shall publish it in the Federal Register and provide opportunity to interested persons to submit to him comments respecting the scientific and medical evaluations which he is to prepare respecting such drug or substance. The Secretary of Health, Education, and Welfare shall prepare for transmission through the Secretary of State to the World Health Organization such medical and scientific evaluations as may be appropriate regarding the possible action that could be proposed by the World Health Organization respecting the drug or substance with respect to which a notice was transmitted under this subparagraph.

Publication in  
Federal  
Register.

"(B) Whenever the Secretary of State receives information that the Commission on Narcotic Drugs of the United Nations proposes to decide whether to add a drug or other substance to one of the schedules of the Convention, transfer a drug or substance from one schedule to another, or delete it from the schedules, the Secretary of State shall transmit timely notice to the Secretary of Health, Education, and Welfare of such information who shall publish a summary of such information in the Federal Register and provide opportunity to interested persons to submit to him comments respecting the recommendation which he is to furnish, pursuant to this subparagraph, respecting such proposal. The Secretary of Health, Education, and Welfare shall evaluate the proposal and furnish a recommendation to the Secretary of State which shall be binding on the representative of the United States in discussions and negotiations relating to the proposal.

Publication in  
Federal  
Register.

"(3) When the United States receives notification of a scheduling decision pursuant to article 2 of the Convention on Psychotropic Substances that a drug or other substance has been added or transferred to a schedule specified in the notification or receives notification (referred to in this subsection as a 'schedule notice') that existing legal controls applicable under this title to a drug or substance and the controls required by the Federal Food, Drug, and Cosmetic Act do not meet the requirements of the schedule of the Convention in which such drug or substance has been placed, the Secretary of Health, Education, and Welfare, after consultation with the Attorney General, shall first determine whether existing legal controls under this title applicable to the drug or substance and the controls required by the Federal Food, Drug, and Cosmetic Act, meet the requirements of the schedule specified in the notification or schedule notice and shall take the following action:

21 USC 301.

"(A) If such requirements are met by such existing controls but the Secretary of Health, Education, and Welfare nonetheless

believes that more stringent controls should be applied to the drug or substance, the Secretary shall recommend to the Attorney General that he initiate proceedings for scheduling the drug or substance, pursuant to subsections (a) and (b) of this section, to apply to such controls.

“(B) If such requirements are not met by such existing controls and the Secretary of Health, Education, and Welfare concurs in the scheduling decision or schedule notice transmitted by the notification, the Secretary shall recommend to the Attorney General that he initiate proceedings for scheduling the drug or substance under the appropriate schedule pursuant to subsections (a) and (b) of this section.

“(C) If such requirements are not met by such existing controls and the Secretary of Health, Education, and Welfare does not concur in the scheduling decision or schedule notice transmitted by the notification, the Secretary shall—

“(i) if he deems that additional controls are necessary to protect the public health and safety, recommend to the Attorney General that he initiate proceedings for scheduling the drug or substance pursuant to subsections (a) and (b) of this section, to apply such additional controls;

“(ii) request the Secretary of State to transmit a notice of qualified acceptance, within the period specified in the Convention, pursuant to paragraph 7 of article 2 of the Convention, to the Secretary-General of the United Nations;

“(iii) request the Secretary of State to transmit a notice of qualified acceptance as prescribed in clause (ii) and request the Secretary of State to ask for a review by the Economic and Social Council of the United Nations, in accordance with paragraph 8 of article 2 of the Convention, of the scheduling decision; or

“(iv) in the case of a schedule notice, request the Secretary of State to take appropriate action under the Convention to initiate proceedings to remove the drug or substance from the schedules under the Convention or to transfer the drug or substance to a schedule under the Convention different from the one specified in the schedule notice.

“(4) (A) If the Attorney General determines, after consultation with the Secretary of Health, Education, and Welfare, that proceedings initiated under recommendations made under paragraph (B) or (C) (i) of paragraph (3) will not be completed within the time period required by paragraph 7 of article 2 of the Convention, the Attorney General, after consultation with the Secretary and after providing interested persons opportunity to submit comments respecting the requirements of the temporary order to be issued under this sentence, shall issue a temporary order controlling the drug or substance under schedule IV or V, whichever is most appropriate to carry out the minimum United States obligations under paragraph 7 of article 2 of the Convention. As a part of such order, the Attorney General shall, after consultation with the Secretary, except such drug or substance from the application of any provision of part C of this title which he finds is not required to carry out the United States obligations under paragraph 7 of article 2 of the Convention. In the case of proceedings initiated under subparagraph (B) of paragraph (3), the Attorney General, concurrently with the issu-

Notice.

ance of such order, shall request the Secretary of State to transmit a notice of qualified acceptance to the Secretary-General of the United Nations pursuant to paragraph 7 of article 2 of the Convention. A temporary order issued under this subparagraph controlling a drug or other substance subject to proceedings initiated under subsections (a) and (b) of this section shall expire upon the effective date of the application to the drug or substance of the controls resulting from such proceedings.

“(B) After a notice of qualified acceptance of a scheduling decision with respect to a drug or other substance is transmitted to the Secretary-General of the United Nations in accordance with clause (ii) or (iii) of paragraph (3) (C) or after a request has been made under clause (iv) of such paragraph with respect to a drug or substance described in a schedule notice, the Attorney General, after consultation with the Secretary of Health, Education, and Welfare and after providing interested persons opportunity to submit comments respecting the requirements of the order to be issued under this sentence, shall issue an order controlling the drug or substance under schedule IV or V, whichever is most appropriate to carry out the minimum United States obligations under paragraph 7 of article 2 of the Convention in the case of a drug or substance for which a notice of qualified acceptance was transmitted or whichever the Attorney General determines is appropriate in the case of a drug or substance described in a schedule notice. As a part of such order, the Attorney General shall, after consultation with the Secretary, except such drug or substance from the application of any provision of part C of this title which he finds is not required to carry out the United States obligations under paragraph 7 of article 2 of the Convention. If, as a result of a review under paragraph 8 of article 2 of the Convention of the scheduling decision with respect to which a notice of qualified acceptance was transmitted in accordance with clause (ii) or (iii) of paragraph (3) (C)—

“(i) the decision is reversed, and

“(ii) the drug or substance subject to such decision is not required to be controlled under schedule IV or V to carry out the minimum United States obligations under paragraph 7 of article 2 of the Convention,

the order issued under this subparagraph with respect to such drug or substance shall expire upon receipt by the United States of the review decision. If, as a result of action taken pursuant to action initiated under a request transmitted under clause (iv) of paragraph (3) (C), the drug or substance with respect to which such action was taken is not required to be controlled under schedule IV or V, the order issued under this paragraph with respect to such drug or substance shall expire upon receipt by the United States of a notice of the action taken with respect to such drug or substance under the Convention.

“(C) An order issued under subparagraph (A) or (B) may be issued without regard to the findings required by subsection (a) of this section or by section 202(b) and without regard to the procedures prescribed by subsection (a) or (b) of this section.

“(5) Nothing in the amendments made by the Psychotropic Substances Act of 1978 or the regulations or orders promulgated thereunder shall be construed to preclude requests by the Secretary of Health, Education, and Welfare or the Attorney General through the

*Ante*, p. 3768.

Secretary of State, pursuant to article 2 or other applicable provisions of the Convention, for review of scheduling decisions under such Convention, based on new or additional information.”.

(b) Section 102 of the Controlled Substances Act (21 U.S.C. 802) is amended by adding at the end the following:

“Convention on Psychotropic Substances.”

“(29) The term ‘Convention on Psychotropic Substances’ means the Convention on Psychotropic Substances signed at Vienna, Austria, on February 21, 1971; and the term ‘Single Convention on Narcotic Drugs’ means the Single Convention on Narcotic Drugs signed at New York, New York, on March 30, 1961.”.

21 USC 812 note.

(c) For the purpose of carrying out the minimum United States obligations under paragraph 7 of article 2 of the Convention on Psychotropic Substances, signed at Vienna, Austria, on February 21, 1971, with respect to pipradrol and SPA (also known as (-)-1-dimethylamino-1,2-diphenylethane), the Attorney General shall by order, made without regard to sections 201 and 202 of the Controlled Substances Act, place such drugs in schedule IV of such Act.

21 USC 811, 812.

SEC. 103. Subsection (d) of section 202 of the Controlled Substances Act (21 U.S.C. 812(d)) is amended by striking out “and” before “(2)” and by adding the following before the period at the end thereof: “, and (3) such exception does not conflict with United States obligations under the Convention on Psychotropic Substances”.

SEC. 104. Section 307 of the Controlled Substances Act (21 U.S.C. 827) is amended by redesignating subsection (e) as subsection (f) and by inserting after subsection (d) the following new subsection:

21 USC 823.

“(e) In addition to the reporting and recordkeeping requirements under any other provision of this title, each manufacturer registered under section 303 shall, with respect to narcotic and nonnarcotic controlled substances manufactured by it, make such reports to the Attorney General, and maintain such records, as the Attorney General may require to enable the United States to meet its obligations under articles 19 and 20 of the Single Convention on Narcotic Drugs and article 16 of the Convention on Psychotropic Substances. The Attorney General shall administer the requirements of this subsection in such a manner as to avoid the unnecessary imposition of duplicative requirements under this title on manufacturers subject to the requirements of this subsection.”.

SEC. 105. Section 1002(b) of the Controlled Substances Import and Export Act (21 U.S.C. 952(b)) is amended by inserting immediately before the period at the end of paragraph (2) the following: “, except that if a nonnarcotic controlled substance in schedule III, IV, or V is also listed in schedule I or II of the Convention on Psychotropic Substances it shall be imported pursuant to such import permit requirements, prescribed by regulation of the Attorney General, as are required by the Convention”.

SEC. 106. Subsection (e) of section 1003 of the Controlled Substances Import and Export Act (21 U.S.C. 953(e)) is amended—

(1) by striking out “, and” at the end of paragraph (2) and inserting in lieu thereof a semicolon;

(2) by striking out the period at the end of paragraph (3) and inserting in lieu thereof “; and”; and

(3) by adding after paragraph (3) the following new paragraph:

“(4) in any case when a nonnarcotic controlled substance in schedule III, IV, or V is also listed in schedule I or II of the



Convention on Psychotropic Substances, it is exported pursuant to such export permit requirements, prescribed by regulation of the Attorney General, as are required by the Convention, instead of the invoice required by paragraphs (2) and (3) of this subsection."

SEC. 107 (a) Part D of the Controlled Substances Act (21 U.S.C. 841 et seq.) is amended by adding at the end thereof the following new section:

"APPLICATION OF TREATIES AND OTHER INTERNATIONAL AGREEMENTS

"SEC. 412. Nothing in the Single Convention on Narcotic Drugs, the Convention on Psychotropic Substances, or other treaties or international agreements shall be construed to limit the provision of treatment, education, or rehabilitation as alternatives to conviction or criminal penalty for offenses involving any drug or other substance subject to control under any such treaty or agreement." 21 USC 852.

(b) The table of contents of the Comprehensive Drug Abuse Prevention and Control Act of 1970 is amended by inserting—

21 USC 801 note.

"Sec. 412. Application of treaties and other international agreements." immediately after

"Sec. 411. Proceedings to establish previous convictions."

SEC. 108. (a) Section 502 of the Controlled Substances Act (21 U.S.C. 872) is amended by redesignating subsection (d) as subsection (e), and by adding after subsection (c) the following new subsection:

"(d) Nothing in the Single Convention on Narcotic Drugs, the Convention on Psychotropic Substances, or other treaties or international agreements shall be construed to limit, modify, or prevent the protection of the confidentiality of patient records or of the names and other identifying characteristics of research subjects as provided by any Federal, State, or local law or regulation."

(b) Section 303 of the Public Health Service Act (42 U.S.C. 242a) is amended by redesignating subsection (b) as subsection (c), and by adding after subsection (a) the following new subsection:

"(b) Nothing in the Single Convention on Narcotic Drugs, the Convention on Psychotropic Substances, or other treaties or international agreements shall be construed to limit, modify, or prevent the protection of the confidentiality of patient records or of the names and other identifying characteristics of research subjects as provided by any Federal, State, or local law or regulation."

SEC. 109. Subsection (f) of section 303 of the Controlled Substances Act (21 U.S.C. 823(f)) is amended by adding at the end the following sentence: "Article 7 of the Convention on Psychotropic Substances shall not be construed to prohibit, or impose additional restrictions upon, research involving drugs or other substances scheduled under the Convention which is conducted in conformity with this subsection and other applicable provisions of this title."

SEC. 110. Subsection (c) of section 307 of the Controlled Substances Act (21 U.S.C. 827(c)) is amended by adding the following after and below paragraph (3): "Nothing in the Convention on Psychotropic Substances shall be construed as superseding or otherwise affecting the provisions of paragraph (1)(B), (2), or (3) of this subsection."

SEC. 111. Subsection (n) of section 502 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 352(n)) is amended by adding the fol-

lowing new sentence at the end thereof: "Nothing in the Convention on Psychotropic Substances, signed at Vienna, Austria, on February 21, 1971, shall be construed to prevent drug price communications to consumers."

Effective date.  
21 USC 801a  
note.

SEC. 112. This title and the amendments made by this title shall take effect on the date the Convention on Psychotropic Substances, signed at Vienna, Austria on February 21, 1971, enters into force in respect to the United States.

## TITLE II—PCP CRIMINAL PENALTIES AND PIPERIDINE REPORTING

SEC. 201. Section 401 of the Controlled Substances Act (21 U.S.C. 841) is amended—

(1) by inserting ", except as provided in paragraphs (4) and (5) of this subsection," after "such person shall" in the first sentence of subsection (b) (1) (B);

(2) by adding after paragraph (4) of subsection (b) the following new paragraph:

"(5) Notwithstanding paragraph (1) (B) of this subsection, any person who violates subsection (a) of this section by manufacturing, distributing, dispensing, or possessing with intent to manufacture, distribute, or dispense, except as authorized by this title, phencyclidine (as defined in section 310(c)(2)) shall be sentenced to a term of imprisonment of not more than 10 years, a fine of not more than \$25,000, or both. If any person commits such a violation after one or more prior convictions of him for an offense punishable under paragraph (1) of this paragraph, or for a felony under any other provision of this title or title III or other law of the United States relating to narcotic drugs, marihuana, or depressant or stimulant substances, have become final, such person shall be sentenced to a term of imprisonment of not more than 20 years, a fine of not more than \$50,000, or both. Any sentence imposing a term of imprisonment under this paragraph shall, in the absence of such a prior conviction, impose a special parole term of at least 2 years in addition to such term of imprisonment and shall, if there was such a prior conviction, impose a special parole term of at least 4 years in addition to such term of imprisonment."; and

Post, p. 3775.

(3) by adding after subsection (c) the following new subsection:

"(d) Any person who knowingly or intentionally—

"(1) possesses any piperidine with intent to manufacture phencyclidine except as authorized by this title, or

"(2) possesses any piperidine knowing, or having reasonable cause to believe, that the piperidine will be used to manufacture phencyclidine except as authorized by this title,

shall be sentenced to a term of imprisonment of not more than 5 years, a fine of not more than \$15,000, or both."

SEC. 202. (a) Part C of the Controlled Substances Act is amended (1) by inserting "; PIPERIDINE REPORTING" at the end of its heading, and (2) by adding after section 309 (21 U.S.C. 829) the following new section:

## "PIPERIDINE REPORTING

"SEC. 310. (a) (1) Except as provided under paragraph (3), any person who distributes, sells, or imports any piperidine shall report to the Attorney General such information, in such form and manner, and within such time period or periods (of not less than seven days), concerning the distribution, sale, or importation as the Attorney General may require by regulation, and the person shall preserve a copy of each such report for 2 years. The Attorney General may include in the information required to be reported the following: 21 USC 830.

"(A) The quantity, form, and manner in which, and date on which, the piperidine was distributed, sold, or imported.

"(B) (i) In the case of the distribution or sale of piperidine to an individual, the name, address, and age of the individual and the type of identification presented to confirm the identity of the individual.

"(ii) In the case of the distribution or sale of piperidine to an entity other than an individual, the name and address of the entity and the name, address, and title of the individual ordering or receiving the piperidine and the type of identification presented to confirm the identity of the individual and of the entity.

"(2) Except as provided under paragraph (3), no person may distribute or sell piperidine unless the recipient or purchaser presents to the distributor or seller identification of such type, to confirm the identity of the recipient or purchaser (and any entity which the recipient or purchaser represents), as the Attorney General establishes by regulation.

"(3) Under such conditions and to such extent as the Attorney General establishes, paragraphs (1) and (2) shall not apply to—

"(A) the distribution of piperidine between agents or employees within a single facility (as defined by the Attorney General), if such agents or employees are acting in the lawful and usual course of their business or employment;

"(B) the delivery of piperidine to or by a common or contract carrier for carriage in the lawful and usual course of its business, or to or by a warehouseman for storage in the lawful and usual course of its business; but where such carriage or storage is in connection with the distribution, sale, or importation of the piperidine to a third person, this subparagraph shall not relieve the distributor, seller, or importer from compliance with paragraph (1) or (2); or

"(C) any distribution, sale, or importation of piperidine with respect to which the Attorney General determines that the report required by paragraph (1) or the presentation of identification required by paragraph (2) is not necessary for the enforcement of this title.

"(b) Any information which is reported to or otherwise obtained by the Department of Justice under this section and which is exempt from disclosure pursuant to subsection (a) of section 552 of title 5, United States Code, by reason of subsection (b) (4) thereof shall be considered confidential and shall not be disclosed, except that such information may be disclosed to officers or employees of the United States concerned with carrying out this title or title III or when relevant in any proceeding for the enforcement of this title or title III.



Definitions.  
21 USC 841,  
842.  
21 USC 951.

"(c) For purposes of this section, section 401(d), and section 402(a)(9):

"(1) The term 'import' has the meaning given such term in section 1001(a)(1).

"(2) The term 'phencyclidine' means 1-(1-phenylcyclohexyl) piperidine, its salts, or any immediate precursor, homolog, analog, or derivative (or salt thereof) of 1-(1-phenylcyclohexyl) piperidine that is included in schedule I or II of part B of this title.

"(3) The term 'piperidine' includes its salts and acyl derivatives."

(b)(1) Section 402(a) of such Act (21 U.S.C. 842(a)) is amended—

(A) by striking out "or" at the end of paragraph (7);

(B) by striking out the period at the end of paragraph (8) and inserting in lieu thereof "; or"; and

(C) by adding after paragraph (8) the following new paragraph:

"(9) to distribute or sell piperidine in violation of regulations established under section 310(a)(2), respecting presentation of identification."

*Ante*, p. 3775.

(2) Section 402(c)(2) of such Act (21 U.S.C. 842(c)(2)) is amended by adding after subparagraph (B) the following new subparagraph:

"(C) Subparagraphs (A) and (B) shall not apply to a violation of subsection (a)(5) with respect to a refusal or failure to make a report required under section 310(a) (relating to piperidine reporting)."

(3) Section 403(a)(4) of such Act (21 U.S.C. 843(a)(4)) is amended—

(A) by inserting "(A)" after "(4)", and

(B) by inserting before "; or" the following: ", or (B) to present false or fraudulent identification where the person is receiving or purchasing piperidine and the person is required to present identification under section 310(a)".

21 USC 801 note.

(c) The table of contents of the Comprehensive Drug Abuse Prevention and Control Act of 1970 is amended—

(1) by inserting "; PIPERIDINE REPORTING" at the end of the item relating to part C, and

21 USC 829.

(2) by adding immediately after the item relating to section 309 the following new item:

"Sec. 310. Piperidine reporting."

21 USC 830 note.

SEC. 203. (a)(1) Except as provided under paragraph (2), the amendments made by this title shall take effect on the date of the enactment of this Act.

(2) Any person required to submit a report under section 310(a)(1) of the Controlled Substances Act respecting a distribution, sale, or importation of piperidine during the 90 days after the date of the enactment of this Act may submit such report any time up to 97 days after such date of enactment.

(3) Until otherwise provided by the Attorney General by regulation, the information required to be reported by a person under section 310(a)(1) of the Controlled Substances Act (as added by section 202(a)(2) of this title) with respect to the person's distribution, sale, or importation of piperidine shall—

(A) be the information described in subparagraphs (A) and (B) of such section, and

(B) except as provided in paragraph (2) of this subsection, be reported not later than seven days after the date of such distribution, sale, or importation.

## (b) The Attorney General shall—

(1) first publish proposed interim regulations to carry out the requirements of section 310(a) of the Controlled Substances Act (as added by section 202(a)(2) of this title) not later than 30 days after the date of the enactment of this Act, and

(2) first promulgate final interim regulations to carry out such requirements not later than 75 days after the date of the enactment of this Act, such final interim regulations to be effective with respect to distributions, sales, and importations of piperidine on and after the ninety-first day after the date of the enactment of this Act.

(c) The Attorney General, after consultation with the Secretary of Health, Education, and Welfare, shall analyze and evaluate the impact and effectiveness of the amendments made by this title, including the impact on the illicit manufacture and use of phencyclidine and the impact of the requirements imposed by such amendments on legitimate distributions and uses of piperidine. Not later than March 1, 1980, the Attorney General shall report to the President and the Congress on such analysis and evaluation and shall include in such report such recommendations as the Attorney General deems appropriate.

Report to  
President and  
Congress.

(d) On January 1, 1981, section 310, subsection (d) of section 401, paragraph (9) of section 402(a), subparagraph (C) of section 402(c)(2), and clause (B) of section 403(a)(4) of the Controlled Substances Act (as added by this title) are repealed.

Repeals.

*Ante*, p. 3775,  
3776.

### TITLE III—FORFEITURE OF PROCEEDS OF ILLEGAL DRUG TRANSACTIONS

SEC. 301. (a) Section 511 of the Comprehensive Drug Abuse Prevention and Control Act (21 U.S.C. 881) is amended—

(1) by adding at the end of subsection (a) the following new paragraph:

“(6) All moneys, negotiable instruments, securities, or other things of value furnished or intended to be furnished by any person in exchange for a controlled substance in violation of this title, all proceeds traceable to such an exchange, and all moneys, negotiable instruments, and securities used or intended to be used to facilitate any violation of this title, except that no property shall be forfeited under this paragraph, to the extent of the interest of an owner, by reason of any act or omission established by that owner to have been committed or omitted without the knowledge or consent of that owner.”;

(2) in subsection (e)(2) by striking out “, but the proceeds” and all that follows through “court costs”; and

(3) by adding at the end of subsection (e) the following new sentence:

"The proceeds from any sale under paragraph (2) and any moneys forfeited under this title shall be used to pay all proper expenses of the proceedings for forfeiture and sale including expenses of seizure, maintenance of custody, advertising, and court costs. The Attorney General shall forward to the Treasurer of the United States for deposit in the general fund of the United States Treasury any amounts of such moneys and proceeds remaining after payment of such expenses."

(b) The second sentence of section 1015 of such Act (21 U.S.C. 965) is amended by inserting "or 511" after "510" each place it appears.

Approved November 10, 1978.

#### LEGISLATIVE HISTORY:

HOUSE REPORT No. 95-1193 accompanying H.R. 12008 (Comm. on Interstate and Foreign Commerce).

SENATE REPORT No. 95-959 (Comm. on the Judiciary).

CONGRESSIONAL RECORD, Vol. 124 (1978):

July 27, considered and passed Senate.

Sept. 18, considered and passed House, amended, in lieu of H.R. 12008.

Oct. 7, Senate concurred in House amendment with amendments.

Oct. 13, House concurred in Senate amendments.

WEEKLY COMPILATION OF PRESIDENTIAL DOCUMENTS, Vol. 14, No. 45:

Nov. 10, Presidential statement.