

106TH CONGRESS  
2D SESSION

# H. R. 3636

To amend the Internal Revenue Code of 1986 with respect to the purchase of prescription drugs by individuals who have attained retirement age, and to amend the Federal Food, Drug, and Cosmetic Act with respect to the importation of prescription drugs and the sale of such drugs through Internet sites.

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## IN THE HOUSE OF REPRESENTATIVES

FEBRUARY 10, 2000

Mr. PAUL introduced the following bill; which was referred to the Committee on Commerce, and in addition to the Committee on Ways and Means, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned

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## A BILL

To amend the Internal Revenue Code of 1986 with respect to the purchase of prescription drugs by individuals who have attained retirement age, and to amend the Federal Food, Drug, and Cosmetic Act with respect to the importation of prescription drugs and the sale of such drugs through Internet sites.

1 *Be it enacted by the Senate and House of Representa-*  
 2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Pharmaceutical Free-  
 5 dom Act of 2000”.

6 **TITLE I—AMENDMENTS TO IN-**  
 7 **TERNAL REVENUE CODE OF**  
 8 **1986**

9 **SEC. 101. INCOME TAX CREDIT FOR PRESCRIPTION DRUGS**  
 10 **PURCHASED BY INDIVIDUALS WHO HAVE AT-**  
 11 **TAINED RETIREMENT AGE.**

12 (a) IN GENERAL.—Subpart A of part IV of sub-  
 13 chapter A of chapter 1 of the Internal Revenue Code of  
 14 1986 (relating to nonrefundable personal credits) is  
 15 amended by inserting after section 25A the following new  
 16 section:

17 **“SEC. 25B. PRESCRIPTION DRUGS PURCHASED BY INDIVID-**  
 18 **UALS WHO HAVE ATTAINED SOCIAL SECU-**  
 19 **RITY RETIREMENT AGE.**

20 “(a) IN GENERAL.—In the case of an individual who  
 21 has attained social security retirement age, there shall be  
 22 allowed as a credit against the tax imposed by this chapter  
 23 for the taxable year an amount equal to 80 percent of the  
 24 amount paid by the taxpayer during the taxable year (and  
 25 not compensated for by insurance or otherwise) for any

1 prescribed drug (as defined in section 213(d)(3)) for use  
 2 by such individual.

3 “(b) SOCIAL SECURITY RETIREMENT AGE.—For  
 4 purposes of this section, the term ‘social security retire-  
 5 ment age’ means retirement age (as defined in section  
 6 216(l)(1) of the Social Security Act).

7 “(c) DENIAL OF DOUBLE BENEFIT.—

8 “(1) COORDINATION WITH MEDICAL EXPENSE  
 9 DEDUCTION.—The amount which would (but for this  
 10 subsection) be taken into account by the taxpayer  
 11 under section 213 for the taxable year shall be re-  
 12 duced by the credit (if any) allowed by this section  
 13 to the taxpayer for such year.

14 “(2) COORDINATION WITH MEDICAL SAVINGS  
 15 ACCOUNTS.—No credit shall be allowed under this  
 16 section for amounts paid from any medical savings  
 17 account (as defined in section 220(d)).

18 “(d) ELECTION NOT TO HAVE CREDIT APPLY.—  
 19 This section shall not apply to a taxpayer for a taxable  
 20 year if the taxpayer elects not to have this section apply  
 21 for such year.”

22 (b) CLERICAL AMENDMENT.—The table of sections  
 23 for subpart A of part IV of subchapter A of chapter 1  
 24 of such Code is amended by inserting after the item relat-  
 25 ing to section 25A the following new item:

“Sec. 25B. Prescription drugs purchased by individuals who have attained social security retirement age.”

1 (c) EFFECTIVE DATE.—The amendments made by  
2 this section shall apply to taxable years beginning more  
3 than 1 year after the date of the enactment of this Act.

4 **TITLE II—AMENDMENTS TO FED-**  
5 **ERAL FOOD, DRUG, AND COS-**  
6 **METIC ACT**

7 **SEC. 201. FACILITATION OF IMPORTATION OF DRUGS AP-**  
8 **PROVED BY FOOD AND DRUG ADMINISTRA-**  
9 **TION.**

10 (a) IN GENERAL.—Section 801(d) of the Federal  
11 Food, Drug, and Cosmetic Act (21 U.S.C. 381(d)) is  
12 amended—

13 (1) by striking paragraph (2); and

14 (2) by striking “(d)(1)” and all that follows  
15 through the end of paragraph (1) and inserting the  
16 following:

17 “(d)(1)(A) A person who meets applicable legal re-  
18 quirements to be an importer of drugs described in sub-  
19 paragraph (B) may import such a drug (without regard  
20 to whether the person is a manufacturer of the drug) if  
21 the person submits to the Secretary an application to im-  
22 port the drug and the Secretary approves the application.

23 “(B) For purposes of subparagraph (A), the drugs  
24 described in this subparagraph are drugs that are subject

1 to section 503(b)(1) or that are composed wholly or partly  
2 of insulin.

3 “(C) The Secretary shall approve an application  
4 under subparagraph (A) if the application demonstrates  
5 that the drug to be imported meets all requirements under  
6 this Act for the admission of the drug into the United  
7 States, including demonstrating that—

8 “(i) an application for the drug has been ap-  
9 proved under section 505, or as applicable, under  
10 section 351 of the Public Health Service Act; and

11 “(ii) the drug is not adulterated or misbranded.

12 “(D) Not later than 60 days after the date on which  
13 an application under subparagraph (A) is submitted to the  
14 Secretary, the Secretary shall—

15 “(i) approve the application; or

16 “(ii) refuse to approve the application and pro-  
17 vide to the person who submitted the application the  
18 reason for such refusal.

19 “(E) This paragraph may not be construed as affect-  
20 ing any right secured by patent.”.

21 (b) CONFORMING AMENDMENTS.—Section 801(d) of  
22 the Federal Food, Drug, and Cosmetic Act (21 U.S.C.  
23 381(d)) is amended—

24 (1) by redesignating paragraphs (3) and (4) as  
25 paragraphs (2) and (3), respectively; and

1           (2) in paragraph (3) (as so redesignated) by  
2           striking “paragraph (3)” each place such term ap-  
3           pears and inserting “paragraph (2)”.

4   **SEC. 202. INTERNET SALES OF PRESCRIPTION DRUGS.**

5           Section 503(b) of the Federal Food, Drug, and Cos-  
6   metic Act (21 U.S.C. 353(b)) is amended by adding at  
7   the end the following paragraph:

8           “(6)(A) With respect to the interstate sale of a pre-  
9   scription drug through an Internet site, the Secretary may  
10   not with respect to such sale take any action under this  
11   Act against any of the persons involved if—

12           “(i) the sale was made in compliance with this  
13   Act and with State laws that are applicable to the  
14   sale of the drug; and

15           “(ii) accurate information regarding compliance  
16   with this Act and such State laws is posted on the  
17   Internet site.

18           “(B) For purposes of subparagraph (A), the sale of  
19   a prescription drug by a person shall be considered to be  
20   an interstate sale of the drug through an Internet site if—

21           “(i) the purchaser of the drug submits the pur-  
22   chase order for the drug, or conducts any other part  
23   of the sales transaction for the drug, through an  
24   Internet site; and

1           “(ii) pursuant to such sale, the person intro-  
 2           duces the drug into interstate commerce or delivers  
 3           the drug for introduction into such commerce.

4           “(C) Subparagraph (A) may not be construed as au-  
 5           thorizing the Secretary to enforce any violation of State  
 6           law.

7           “(D) For purposes of this paragraph, the term ‘pre-  
 8           scription drug’ means a drug that is subject to paragraph  
 9           (1).”.

10   **SEC. 203. REGULATIONS OF SECRETARY OF HEALTH AND**  
 11           **HUMAN SERVICES; EFFECTIVE DATE.**

12           (a) REGULATIONS.—Before the expiration of the pe-  
 13           riod specified in subsection (b), the Secretary of Health  
 14           and Human Services shall promulgate regulations to carry  
 15           out the amendments to the Federal Food, Drug, and Cos-  
 16           metic Act that are made by sections 201 and 202.

17           (b) EFFECTIVE DATE.—The amendments to the Fed-  
 18           eral Food, Drug, and Cosmetic Act that are made by sec-  
 19           tions 201 and 202 take effect upon the expiration of the  
 20           one-year period beginning on the date of the enactment  
 21           of this Act, without regard to whether the regulations re-  
 22           quired in subsection (a) have been promulgated.

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