

1 (A) by striking “credit” and inserting  
2 “credits”, and

3 (B) by striking “subsection (a)” and in-  
4 serting “subsection (a) or (b)”.

5 (c) AGGREGATION RULES.—Clause (ii) of section  
6 41(h)(5)(B) is amended by striking “the \$250,000  
7 amount” and inserting “each of the \$250,000 amounts”.

8 (d) EFFECTIVE DATE.—The amendments made by  
9 this section shall apply to taxable years beginning after  
10 December 31, 2021.

11 **SEC. 138520. IMPOSITION OF TAX ON NICOTINE.**

12 (a) IN GENERAL.—Section 5701 is amended by re-  
13 designating subsection (h) as subsection (i) and by insert-  
14 ing after subsection (g) the following new subsection:

15 “(h) NICOTINE.—On taxable nicotine, manufactured  
16 in or imported into the United States, there shall be im-  
17 posed a tax equal to the dollar amount specified in section  
18 5701(b)(1) (or, if greater, \$50.33) per 1,810 milligrams  
19 of nicotine (and a proportionate tax at the like rate on  
20 any fractional part thereof).”.

21 (b) TAXABLE NICOTINE.—Section 5702 is amended  
22 by adding at the end the following new subsection:

23 “(q) TAXABLE NICOTINE.—

24 “(1) IN GENERAL.—Except as otherwise pro-  
25 vided in this subsection, the term ‘taxable nicotine’

1 means any nicotine which has been extracted, con-  
2 centrated, or synthesized.

3 “(2) EXCEPTION FOR PRODUCTS APPROVED BY  
4 FOOD AND DRUG ADMINISTRATION.—Such term  
5 shall not include any nicotine if the manufacturer or  
6 importer thereof demonstrates to the satisfaction of  
7 the Secretary of Health and Human Services that  
8 such nicotine will be used in—

9 “(A) a drug—

10 “(i) that is approved under section  
11 505 of the Federal Food, Drug, and Cos-  
12 metic Act or licensed under section 351 of  
13 the Public Health Service Act; or

14 “(ii) for which an investigational use  
15 exemption has been authorized under sec-  
16 tion 505(i) of the Federal Food, Drug, and  
17 Cosmetic Act or under section 351(a) of  
18 the Public Health Service Act; or

19 “(B) a combination product (as described  
20 in section 503(g) of the Federal Food, Drug,  
21 and Cosmetic Act), the constituent parts of  
22 which were approved or cleared under section  
23 505, 510(k), or 515 of such Act.

24 “(3) COORDINATION WITH TAXATION OF OTHER  
25 TOBACCO PRODUCTS.—Tobacco products meeting

1 the definition of cigars, cigarettes, smokeless to-  
2 bacco, pipe tobacco, and roll-your-own tobacco in  
3 this section shall be classified and taxed as such de-  
4 spite any concentration of the nicotine inherent in  
5 those products or any addition of nicotine to those  
6 products during the manufacturing process.

7 “(4) REGULATIONS.—The Secretary shall pre-  
8 scribe such regulations or other guidance as is nec-  
9 essary or appropriate to carry out the purposes of  
10 this subsection, including regulations or other guid-  
11 ance for coordinating the taxation of tobacco prod-  
12 ucts and taxable nicotine to protect revenue and pre-  
13 vent double taxation.”.

14 (c) TAXABLE NICOTINE TREATED AS A TOBACCO  
15 PRODUCT.—Section 5702(c) is amended by striking “and  
16 roll-your-own tobacco” and inserting “roll-your-own to-  
17 bacco, and taxable nicotine”.

18 (d) MANUFACTURER OF TAXABLE NICOTINE.—Sec-  
19 tion 5702, as amended by subsection (b), is amended by  
20 adding at the end the following new subsection:

21 “(r) MANUFACTURER OF TAXABLE NICOTINE.—

22 “(1) IN GENERAL.—Any person who extracts,  
23 concentrates, or synthesizes nicotine shall be treated  
24 as a manufacturer of taxable nicotine (and as manu-  
25 facturing such taxable nicotine).

1           “(2) APPLICATION OF RULES RELATED TO  
2 MANUFACTURERS OF TOBACCO PRODUCTS.—Any  
3 reference to a manufacturer of tobacco products, or  
4 to manufacturing tobacco products, shall be treated  
5 as including a reference to a manufacturer of tax-  
6 able nicotine, or to manufacturing taxable nicotine,  
7 respectively.”.

8 (e) EFFECTIVE DATE.—

9           (1) IN GENERAL.—The amendments made by  
10 this section shall apply to articles removed in cal-  
11 endar quarters beginning after the date which is 180  
12 days after the date of the enactment of this Act.

13           (2) TRANSITION RULE FOR PERMIT AND BOND  
14 REQUIREMENTS.—A person which is lawfully en-  
15 gaged in business as a manufacturer or importer of  
16 taxable nicotine (within the meaning of subchapter  
17 A of chapter 52 of the Internal Revenue Code of  
18 1986, as amended by this section) on the date of the  
19 enactment of this Act, first becomes subject to the  
20 requirements of subchapter B of chapter 52 of such  
21 Code by reason of the amendments made by this  
22 section, and submits an application under such sub-  
23 chapter B to engage in such business not later than  
24 90 days after the date of the enactment of this Act,  
25 shall not be denied the right to carry on such busi-

1           ness by reason of such requirements before final ac-  
2           tion on such application.

3   **SEC. 138521. TERMINATION OF EMPLOYER CREDIT FOR**  
4                           **PAID FAMILY AND MEDICAL LEAVE.**

5           Section 45S(i) is amended by striking “December 31,  
6 2025” and inserting “December 31, 2023”.

7                           **Subtitle I—Drug Pricing**

8           **PART 1—LOWERING PRICES THROUGH DRUG**  
9                           **PRICE NEGOTIATION**

10   **SEC. 139001. PROVIDING FOR LOWER PRICES FOR CERTAIN**  
11                           **HIGH-PRICED SINGLE SOURCE DRUGS.**

12           (a) PROGRAM TO LOWER PRICES FOR CERTAIN  
13 HIGH-PRICED SINGLE SOURCE DRUGS.—Title XI of the  
14 Social Security Act is amended by adding after section  
15 1184 (42 U.S.C. 1320e–3) the following new part:

16   **“PART E—PRICE NEGOTIATION PROGRAM TO**  
17           **LOWER PRICES FOR CERTAIN HIGH-PRICED**  
18           **SINGLE SOURCE DRUGS**

19   **“SEC. 1191. ESTABLISHMENT OF PROGRAM.**

20           “(a) IN GENERAL.—The Secretary shall establish a  
21 Drug Price Negotiation Program (in this part referred to  
22 as the ‘program’). Under the program, with respect to  
23 each price applicability period, the Secretary shall—

24                           “(1) publish a list of negotiation-eligible drugs  
25                           and selected drugs in accordance with section 1192;