

package (as defined in section 2 of the Poison Prevention Packaging Act of 1970 (15 U.S.C. 1471))—

- (i) from which nicotine in a solution or other form is accessible through normal and foreseeable use by a consumer; and
- (ii) that is used to hold soluble nicotine in any concentration.

(B) Exclusion

The term “liquid nicotine container” does not include a sealed, pre-filled, and disposable container of nicotine in a solution or other form in which such container is inserted directly into an electronic cigarette, electronic nicotine delivery system, or other similar product, if the nicotine in the container is inaccessible through customary or reasonably foreseeable handling or use, including reasonably foreseeable ingestion or other contact by children.

(3) Nicotine

The term “nicotine” means any form of the chemical nicotine, including any salt or complex, regardless of whether the chemical is naturally or synthetically derived.

(Pub. L. 114–116, §2, Jan. 28, 2016, 130 Stat. 3.)

REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in subsec. (b)(1), is act June 25, 1938, ch. 675, 52 Stat. 1040, which is classified generally to chapter 9 (§301 et seq.) of Title 21, Food and Drugs. For complete classification of this Act to the Code, see section 301 of Title 21 and Tables.

The Family Smoking Prevention and Tobacco Control Act, referred to in subsec. (b)(1), is div. A of Pub. L. 111–31, June 22, 2009, 123 Stat. 1776. For complete classification of this Act to the Code, see Short Title of 2009 Amendment note set out under section 301 of Title 21, Food and Drugs, and Tables.

The Poison Prevention Packaging Act of 1970, referred to in subsec. (b)(2), is Pub. L. 91–601, Dec. 30, 1970, 84 Stat. 1670, which is classified principally to this chapter. For complete classification of this Act to the Code, see Short Title note set out under section 1471 of this title and Tables.

CODIFICATION

Section was enacted as part of the Child Nicotine Poisoning Prevention Act of 2015, and not as part of the Poison Prevention Packaging Act of 1970 which comprises this chapter.

EFFECTIVE DATE

Pub. L. 114–116, §3, Jan. 28, 2016, 130 Stat. 5, provided that: “This Act [see Short Title of 2016 Amendment note set out under section 1471 of this title] shall take effect on the date that is 180 days after the date of the enactment of this Act [Jan. 28, 2016].”

§ 1473. Conventional packages, marketing

(a) Noncomplying packages for elderly or handicapped persons; labeling statements

For the purpose of making any household substance which is subject to a standard established under section 1472 of this title readily available to elderly or handicapped persons unable to use such substance when packaged in compliance with such standard, the manufacturer or packer, as the case may be, may package any household substance, subject to such a standard, in packaging of a single size which does not comply with such standard if—

(1) the manufacturer (or packer) also supplies such substance in packages which comply with such standard; and

(2) the packages of such substance which do not meet such standard bear conspicuous labeling stating: “This package for households without young children”; except that the Commission may by regulation prescribe a substitute statement to the same effect for packaging too small to accommodate such labeling.

(b) Noncomplying packages for substances dispensed pursuant to orders of medical practitioners

In the case of a household substance which is subject to such a standard and which is dispensed pursuant to an order of physician, dentist, or other licensed medical practitioner authorized to prescribe, such substance may be dispensed in noncomplying packages only when directed in such order or when requested by the purchaser.

(c) Exclusive use of special packaging; necessary circumstances

In the case of a household substance subject to such a standard which is packaged under subsection (a) in a noncomplying package, if the Commission determines that such substance is not also being supplied by a manufacturer (or packer) in popular size packages which comply with such standard, it may, after giving the manufacturer (or packer) an opportunity to comply with the purposes of this Act, by order require such substance to be packaged by such manufacturer (or packer) exclusively in special packaging complying with such standard if it finds, after opportunity for hearing, that such exclusive use of special packaging is necessary to accomplish the purposes of this Act.

(Pub. L. 91–601, §4, Dec. 30, 1970, 84 Stat. 1671; Pub. L. 92–573, §30(a), Oct. 27, 1972, 86 Stat. 1231.)

REFERENCES IN TEXT

For classification to the Code of “this Act”, referred to in subsec. (c), see References in Text note set out under section 1471 of this title.

TRANSFER OF FUNCTIONS

“Commission” substituted for “Secretary” in subsecs. (a) and (c) and “it” substituted for “he” in subsec. (c) pursuant to section 30(a) of Pub. L. 92–573, which is classified to section 2079(a) of this title and which transferred functions of Secretary of Health, Education, and Welfare under this chapter to Consumer Product Safety Commission.

§ 1474. Regulations for special packaging standards

(a) Rule making procedure; election and application of procedure under section 371 of title 21; publication of election and proposal

Proceedings to issue, amend, or repeal a regulation prescribing a standard under section 1472 of this title shall be conducted in accordance with the procedures prescribed by section 553 (other than paragraph (3)(B) of the last sentence of subsection (b) of such section) of title 5 unless the Commission elects the procedures prescribed by subsection (e) of section 371 of title 21, in which event such subsection and subsections (f)