

dren's product subject to the consumer product safety standards described in subsection (a) or any rule promulgated under this section. The Commission shall grant such an exemption if the State or political subdivision standard or regulation—

(A) provides a significantly higher degree of protection from such risk of injury than the consumer product safety standard or rule under this section; and

(B) does not unduly burden interstate commerce.

In determining the burden, if any, of a State or political subdivision standard or regulation on interstate commerce, the Commission shall consider and make appropriate (as determined by the Commission in its discretion) findings on the technological and economic feasibility of complying with such standard or regulation, the cost of complying with such standard or regulation, the geographic distribution of the consumer product to which the standard or regulation would apply, the probability of other States or political subdivisions applying for an exemption under this subsection for a similar standard or regulation, and the need for a national, uniform standard under this Act for such consumer product.

(2) Effect of standards on established State laws

Nothing in this section or in section 2075 of this title shall prevent a State or political subdivision of a State from continuing in effect a safety requirement applicable to a toy or other children's product that is designed to deal with the same risk of injury as the consumer product safety standards established by this section and that is in effect on the day before August 14, 2008, if such State or political subdivision has filed such requirement with the Commission within 90 days after August 14, 2008, in such form and in such manner as the Commission may require.

(i) Judicial review

The issuance of any rule under this section is subject to judicial review as provided in section 2060(g) of this title, as added by section 236 of this Act.

(Pub. L. 110-314, title I, §106, Aug. 14, 2008, 122 Stat. 3033; Pub. L. 112-28, §4, Aug. 12, 2011, 125 Stat. 280.)

REFERENCES IN TEXT

This Act, referred to in subsec. (h)(1), is Pub. L. 110-314, Aug. 14, 2008, 122 Stat. 3016, known as the Consumer Product Safety Improvement Act of 2008. For complete classification of this Act to the Code, see Short Title of 2008 Amendment note set out under section 2051 of this title and Tables.

Section 2060(g) of this title, as added by section 236 of this Act, referred to in subsec. (i), is section 2060(g) of this title, as added by section 236 of Pub. L. 110-314.

CODIFICATION

Section was enacted as part of the Consumer Product Safety Improvement Act of 2008, and not as part of the Consumer Product Safety Act which comprises this chapter.

AMENDMENTS

2011—Subsec. (a). Pub. L. 112-28 inserted “or any provision that restates or incorporates a regulation pro-

mulgated by the Food and Drug Administration or any statute administered by the Food and Drug Administration” after “or by statute”.

DEFINITION

For definition of “Commission” used in this section, see section 2(a) of Pub. L. 110-314, set out as a note under section 2051 of this title.

§ 2057. Banned hazardous products

Whenever the Commission finds that—

(1) a consumer product is being, or will be, distributed in commerce and such consumer product presents an unreasonable risk of injury; and

(2) no feasible consumer product safety standard under this chapter would adequately protect the public from the unreasonable risk of injury associated with such product,

the Commission may, in accordance with section 2058 of this title, promulgate a rule declaring such product a banned hazardous product.

(Pub. L. 92-573, §8, Oct. 27, 1972, 86 Stat. 1215; Pub. L. 97-35, title XII, §1203(c), Aug. 13, 1981, 95 Stat. 713.)

AMENDMENTS

1981—Pub. L. 97-35 substituted “may, in accordance with” for “may propose and, in accordance with”.

EFFECTIVE DATE OF 1981 AMENDMENT

Amendment by Pub. L. 97-35 applicable with respect to regulations under this chapter and chapters 25 and 30 of this title for which notices of proposed rulemaking are issued after Aug. 14, 1981, see section 1215 of Pub. L. 97-35, set out as a note under section 2052 of this title.

§ 2057a. Banning of butyl nitrite

(a) In general

Except as provided in subsection (b) of this section, butyl nitrite shall be considered a banned hazardous product under section 2057 of this title.

(b) Lawful purposes

For the purposes of section 2057 of this title, it shall not be unlawful for any person to manufacture for sale, offer for sale, distribute in commerce, or import into the United States butyl nitrite for any commercial purpose or any other purpose approved under the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.].

(c) Definitions

For purposes of this section:

(1) The term “butyl nitrite” includes n-butyl nitrite, isobutyl nitrite, secondary butyl nitrite, tertiary butyl nitrite, and mixtures containing these chemicals.

(2) The term “commercial purpose” means any commercial purpose other than for the production of consumer products containing butyl nitrite that may be used for inhaling or otherwise introducing butyl nitrite into the human body for euphoric or physical effects.

(d) Effective date

This section shall take effect 90 days after November 18, 1988.

(Pub. L. 100-690, title II, §2404, Nov. 18, 1988, 102 Stat. 4231.)

REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in subsec. (b), is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, 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.

CODIFICATION

Section was enacted as part of the Anti-Drug Abuse Act of 1988 and also as part of the Comprehensive Alcohol Abuse, Drug Abuse, and Mental Health Amendments Act of 1988, and not as part of the Consumer Product Safety Act which comprises this chapter.

§ 2057b. Banning of isopropal nitrite and other nitrites

(a) In general

Except as provided in subsection (b) of this section, volatile alkyl nitrite shall be considered a banned hazardous product under section 2057 of this title.

(b) Lawful purposes

For the purposes of section 2057 of this title, it shall not be unlawful for any person to manufacture for sale, offer for sale, distribute in commerce, or import into the United States volatile alkyl nitrites for any commercial purpose or any other purpose approved under the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.].

(c) “Commercial purpose” defined

For purposes of this section, the term “commercial purpose” means any commercial purpose other than for the production of consumer products containing volatile alkyl nitrites that may be used for inhaling or otherwise introducing volatile alkyl nitrites into the human body for euphoric or physical effects.

(d) Effective date

This section shall take effect 90 days after November 29, 1990.

(Pub. L. 101-647, title XXXII, §3202, Nov. 29, 1990, 104 Stat. 4917.)

REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in subsec. (b), is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, 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.

CODIFICATION

Section was enacted as part of the Crime Control Act of 1990, and not as part of the Consumer Product Safety Act which comprises this chapter.

§ 2057c. Prohibition on sale of certain products containing specified phthalates

(a) Prohibition on the sale of certain products containing phthalates

Beginning on the date that is 180 days after August 14, 2008, it shall be unlawful for any person to manufacture for sale, offer for sale, distribute in commerce, or import into the United States any children’s toy or child care article that contains concentrations of more than 0.1 percent of di-(2-ethylhexyl) phthalate (DEHP),

dibutyl phthalate (DBP), or benzyl butyl phthalate (BBP).

(b) Prohibition on the sale of additional products containing certain phthalates

(1) Interim prohibition

Beginning on the date that is 180 days after August 14, 2008, and until a final rule is promulgated under paragraph (3), it shall be unlawful for any person to manufacture for sale, offer for sale, distribute in commerce, or import into the United States any children’s toy that can be placed in a child’s mouth or child care article that contains concentrations of more than 0.1 percent of diisononyl phthalate (DINP), diisodecyl phthalate (DIDP), or di-n-octyl phthalate (DnOP).

(2) Chronic Hazard Advisory Panel

(A) Appointment

Not earlier than 180 days after August 14, 2008, the Commission shall begin the process of appointing a Chronic Hazard Advisory Panel pursuant to the procedures of section 28 of the Consumer Product Safety Act (15 U.S.C. 2077) to study the effects on children’s health of all phthalates and phthalate alternatives as used in children’s toys and child care articles.

(B) Examination

The panel shall, within 18 months after its appointment under subparagraph (A), complete an examination of the full range of phthalates that are used in products for children and shall—

(i) examine all of the potential health effects (including endocrine disrupting effects) of the full range of phthalates;

(ii) consider the potential health effects of each of these phthalates both in isolation and in combination with other phthalates;

(iii) examine the likely levels of children’s, pregnant women’s, and others’ exposure to phthalates, based on a reasonable estimation of normal and foreseeable use and abuse of such products;

(iv) consider the cumulative effect of total exposure to phthalates, both from children’s products and from other sources, such as personal care products;

(v) review all relevant data, including the most recent, best-available, peer-reviewed, scientific studies of these phthalates and phthalate alternatives that employ objective data collection practices or employ other objective methods;

(vi) consider the health effects of phthalates not only from ingestion but also as a result of dermal, hand-to-mouth, or other exposure;

(vii) consider the level at which there is a reasonable certainty of no harm to children, pregnant women, or other susceptible individuals and their offspring, considering the best available science, and using sufficient safety factors to account for uncertainties regarding exposure and susceptibility of children, pregnant women, and other potentially susceptible individuals; and