

section (a) of this section and the fact of publication of a notice under subsection (b) of this section shall be admissible into evidence in any judicial or administrative proceeding.

(2) No action by the Attorney General or the Commission taken pursuant to this section shall be admissible into evidence in any such proceeding for the purpose of supporting or answering any claim under the antitrust laws or under any State law similar to the antitrust laws.

(Pub. L. 98-462, §6, Oct. 11, 1984, 98 Stat. 1818; Pub. L. 103-42, §3(f), June 10, 1993, 107 Stat. 119; Pub. L. 108-237, title I, §107, June 22, 2004, 118 Stat. 664.)

AMENDMENTS

2004—Subsec. (a). Pub. L. 108-237, §107(1), designated existing provisions as par. (1), redesignated former pars. (1) to (3) as subpars. (A) to (C), respectively, of par. (1), and added par. (2).

Subsec. (b). Pub. L. 108-237, §107(2), inserted “, or a notice with respect to such standards development activity that identifies the standards development organization engaged in such activity and that describes such activity in general terms” before period at end of first sentence and “or available to such organization, as the case may be” before period at end of last sentence.

Subsec. (d)(2). Pub. L. 108-237, §107(3), inserted “, or the standards development activity,” after “venture”.

Subsec. (e). Pub. L. 108-237, §107(4), substituted “person or standards development organization that” for “person who” and inserted “or any standards development organization” after “on any person”.

Subsec. (g)(1). Pub. L. 108-237, §107(5), inserted “or standards development organization” after “person”.

1993—Pub. L. 103-42, §3(f)(1), substituted “joint venture” for “joint research and development venture” in section catchline.

Subsec. (a). Pub. L. 103-42, §3(f)(2), (3), substituted “joint venture” for “joint research and development venture” and “October 11, 1984” for “the date of the enactment of this Act” and added par. (3).

Subsecs. (d)(2), (e). Pub. L. 103-42, §3(f)(3), substituted “joint venture” for “joint research and development venture”.

REPORTS ON JOINT VENTURES AND UNITED STATES COMPETITIVENESS

Section 4 of Pub. L. 103-42 provided that:
 “(a) PURPOSE.—The purpose of the reports required by this section is to inform Congress and the American people of the effect of the National Cooperative Research and Production Act of 1993 [15 U.S.C. 4301 et seq.] on the competitiveness of the United States in key technological areas of research, development, and production.
 “(b) ANNUAL REPORT BY THE ATTORNEY GENERAL.—In the 30-day period beginning at each 1-year interval in the 6-year period beginning on the date of the enactment of this Act [June 10, 1993], the Attorney General shall submit to the Committee on the Judiciary of the House of Representatives and the Committee on the Judiciary of the Senate—
 “(1) a list of joint ventures for which notice was filed under section 6(a) of the National Cooperative Research and Production Act of 1993 [15 U.S.C. 4305(a)] during the 12-month period for which such report is made, including—
 “(A) the purpose of each joint venture;
 “(B) the identity of each party described in section 6(a)(1) of such Act; and
 “(C) the identity and nationality of each person described in section 6(a)(3) of such Act; and
 “(2) a list of cases and proceedings, if any, brought during such period under the antitrust laws by the Department of Justice, and by the Federal Trade

Commission, with respect to joint ventures for which notice was filed under such section at any time.

“(c) TRIENNIAL REPORT BY THE ATTORNEY GENERAL.—In the 30-day period beginning at each 3-year interval in the 6-year period beginning on the date of the enactment of this Act [June 10, 1993], the Attorney General, after consultation with such other agencies as the Attorney General considers to be appropriate, shall submit to the Committee on the Judiciary of the House of Representatives and the Committee on the Judiciary of the Senate a description of the technological areas most commonly pursued by joint ventures for production for which notice was filed under section 6(a) of the National Cooperative Research and Production Act of 1993 [15 U.S.C. 4305(a)] during the 3-year period for which such report is made, and an analysis of the trends in the competitiveness of United States industry in such areas.

“(d) REVIEW OF ANTITRUST TREATMENT UNDER FOREIGN LAWS.—In the three 30-day periods beginning 1 year, 3 years, and 6 years after the date of the enactment of this Act [June 10, 1993], the Attorney General, after consultation with such other agencies as the Attorney General considers to be appropriate, shall submit to the Committee on the Judiciary of the House of Representatives and the Committee on the Judiciary of the Senate a report on the antitrust treatment of United States businesses with respect to participation in joint ventures for production, under the law of each foreign nation any of whose domestic businesses disclosed its nationality under section 6(a)(3) of the National Cooperative Research and Production Act of 1993 [15 U.S.C. 4305(a)(3)] at any time.”

§ 4306. Application of section 4303 protections to production of products, processes, and services

Notwithstanding sections 4303 and 4305 of this title, the protections of section 4303 of this title shall not apply with respect to a joint venture’s production of a product, process, or service, as referred to in section 4301(a)(6)(D) of this title, unless—

- (1) the principal facilities for such production are located in the United States or its territories, and
- (2) each person who controls any party to such venture (including such party itself) is a United States person, or a foreign person from a country whose law accords antitrust treatment no less favorable to United States persons than to such country’s domestic persons with respect to participation in joint ventures for production.

(Pub. L. 98-462, §7, as added Pub. L. 103-42, §3(g), June 10, 1993, 107 Stat. 119.)

CHAPTER 70—COMPREHENSIVE SMOKELESS TOBACCO HEALTH EDUCATION

Sec.	
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§ 4401. Public education

(a) Development

(1) The Secretary of Health and Human Services shall establish and carry out a program to inform the public of any dangers to human

health resulting from the use of smokeless tobacco products. In carrying out such program the Secretary shall—

(A) develop educational programs and materials and public service announcements respecting the dangers to human health from the use of smokeless tobacco;

(B) make such programs, materials, and announcements available to States, local governments, school systems, the media, and such other entities as the Secretary determines appropriate to further the purposes of this chapter;

(C) conduct and support research on the effect of smokeless tobacco on human health; and

(D) collect, analyze, and disseminate information and studies on smokeless tobacco and health.

(2) In developing programs, materials, and announcements under paragraph (1) the Secretary shall consult with the Secretary of Education, medical and public health entities, consumer groups, representatives of manufacturers of smokeless tobacco products, and other appropriate entities.

(b) Assistance

The Secretary of Health and Human Services may provide technical assistance and may make grants to States—

(1) to assist in the development of educational programs and materials and public service announcements respecting the dangers to human health from the use of smokeless tobacco,

(2) to assist in the distribution of such programs, materials, and announcements throughout the States, and

(3) to establish 18 as the minimum age for the purchase of smokeless tobacco.

(Pub. L. 99-252, § 2, Feb. 27, 1986, 100 Stat. 30.)

EFFECTIVE DATE

Section 11 of Pub. L. 99-252 provided that:

“(a) IN GENERAL.—Except as provided in sections 3(f) and 5(b) [sections 4402(f) and 4404(b) of this title] and subsection (b), this Act [enacting this chapter and amending section 342 of Title 21, Food and Drugs] shall take effect one year after the date of enactment of this Act [Feb. 27, 1986].

“(b) EXCEPTION.—Sections 2, 3(b), 3(c), 3(d), 3(e), 4(b), 7, 8, 9 [sections 4401, 4402(b) to (e), 4403(b), and 4406 to 4408 of this title], and 10 [amending section 342 of Title 21] shall take effect on the date of the enactment of this Act [Feb. 27, 1986].”

SHORT TITLE

Section 1 of Pub. L. 99-252 provided that: “This Act [enacting this chapter and amending section 342 of Title 21, Food and Drugs] may be cited as the ‘Comprehensive Smokeless Tobacco Health Education Act of 1986’.”

§ 4402. Smokeless tobacco warning

(a) General rule

(1) It shall be unlawful for any person to manufacture, package, sell, offer to sell, distribute, or import for sale or distribution within the United States any smokeless tobacco product unless the product package bears, in accordance with the requirements of this chapter, one of the following labels:

WARNING: This product can cause mouth cancer.

WARNING: This product can cause gum disease and tooth loss.

WARNING: This product is not a safe alternative to cigarettes.

WARNING: Smokeless tobacco is addictive.

(2) Each label statement required by paragraph (1) shall be—

(A) located on the 2 principal display panels of the package, and each label statement shall comprise at least 30 percent of each such display panel; and

(B) in 17-point conspicuous and legible type and in black text on a white background, or white text on a black background, in a manner that contrasts by typography, layout, or color, with all other printed material on the package, in an alternating fashion under the plan submitted under subsection (b)(3), except that if the text of a label statement would occupy more than 70 percent of the area specified by subparagraph (A), such text may appear in a smaller type size, so long as at least 60 percent of such warning area is occupied by the label statement.

(3) The label statements required by paragraph (1) shall be introduced by each tobacco product manufacturer, packager, importer, distributor, or retailer of smokeless tobacco products concurrently into the distribution chain of such products.

(4) The provisions of this subsection do not apply to a tobacco product manufacturer or distributor of any smokeless tobacco product that does not manufacture, package, or import smokeless tobacco products for sale or distribution within the United States.

(5) A retailer of smokeless tobacco products shall not be in violation of this subsection for packaging that—

(A) contains a warning label;

(B) is supplied to the retailer by a license- or permit-holding tobacco product manufacturer, importer, or distributor; and

(C) is not altered by the retailer in a way that is material to the requirements of this subsection.

(b) Required labels

(1) It shall be unlawful for any tobacco product manufacturer, packager, importer, distributor, or retailer of smokeless tobacco products to advertise or cause to be advertised within the United States any smokeless tobacco product unless its advertising bears, in accordance with the requirements of this section, one of the labels specified in subsection (a).

(2)(A) Each label statement required by subsection (a) in smokeless tobacco advertising shall comply with the standards set forth in this paragraph.

(B) For press and poster advertisements, each such statement and (where applicable) any required statement relating to tar, nicotine, or other constituent yield shall comprise at least 20 percent of the area of the advertisement.

(C) The word “WARNING” shall appear in capital letters, and each label statement shall appear in conspicuous and legible type.