

spect to the United States [Jan. 1, 1995], see section 352 of Pub. L. 103-465, set out as a note under section 2531 of this title.

### § 2572. Exemptions

This subchapter does not apply to—

(1) any standards activity engaged in by any Federal agency or State agency for the use (including, but not limited to, use with respect to research and development, production, or consumption) of that agency or the use of another such agency; or

(2) any standards activity engaged in by any private person solely for use in the production or consumption of products by that person.

(Pub. L. 96-39, title IV, §452, July 26, 1979, 93 Stat. 250.)

### § 2573. Reports to Congress on operation of agreement

As soon as practicable after the close of the 3-year period beginning on the date on which this subchapter takes effect, and as soon as practicable after the close of each succeeding 3-year period through 2001, the Trade Representative shall prepare and submit to Congress a report containing an evaluation of the operation of the Agreement, both domestically and internationally, during the period.

(Pub. L. 96-39, title IV, §453, July 26, 1979, 93 Stat. 250; Pub. L. 103-182, title III, §351(b)(2)(A), Dec. 8, 1993, 107 Stat. 2122; Pub. L. 103-465, title III, §351(f), Dec. 8, 1994, 108 Stat. 4957; Pub. L. 104-295, §21(b)(1), Oct. 11, 1996, 110 Stat. 3529.)

#### AMENDMENTS

1996—Pub. L. 104-295 amended directory language of Pub. L. 103-182. See 1993 Amendment note below.

1994—Pub. L. 103-465 inserted “through 2001” after “succeeding 3-year period”.

1993—Pub. L. 103-182, as amended by Pub. L. 104-295, substituted “Trade Representative” for “Special Representative”.

#### EFFECTIVE DATE OF 1994 AMENDMENT

Amendment by Pub. L. 103-465 effective on the date on which the WTO Agreement enters into force with respect to the United States [Jan. 1, 1995], see section 352 of Pub. L. 103-465, set out as a note under section 2531 of this title.

## PART E—STANDARDS AND MEASURES UNDER THE NORTH AMERICAN FREE TRADE AGREEMENT

### SUBPART 1—SANITARY AND PHYTOSANITARY MEASURES

#### § 2575. General

Nothing in this subpart may be construed—

(1) to prohibit a Federal agency or State agency from engaging in activity related to sanitary or phytosanitary measures to protect human, animal, or plant life or health; or

(2) to limit the authority of a Federal agency or State agency to determine the level of protection of human, animal, or plant life or health the agency considers appropriate.

(Pub. L. 96-39, title IV, §461, as added Pub. L. 103-182, title III, §351(a), Dec. 8, 1993, 107 Stat. 2118.)

#### § 2575a. Inquiry point

The standards information center maintained under section 2544 of this title shall, in addition to the functions specified therein, make available to the public relevant documents, at such reasonable fees as the Secretary of Commerce may prescribe, and information regarding—

(1) any sanitary or phytosanitary measure of general application, including any control or inspection procedure or approval procedure proposed, adopted, or maintained by a Federal or State agency;

(2) the procedures of a Federal or State agency for risk assessment, and factors the agency considers in conducting the assessment and in establishing the levels of protection that the agency considers appropriate;

(3) the membership and participation of the Federal Government and State governments in international and regional sanitary and phytosanitary organizations and systems, and in bilateral and multilateral arrangements regarding sanitary and phytosanitary measures, and the provisions of those systems and arrangements; and

(4) the location of notices of the type required under article 719 of the NAFTA, or where the information contained in such notices can be obtained.

(Pub. L. 96-39, title IV, §462, as added Pub. L. 103-182, title III, §351(a), Dec. 8, 1993, 107 Stat. 2118.)

#### § 2575b. Subpart definitions

Notwithstanding section 2571 of this title, for purposes of this subpart—

##### (1) Animal

The term “animal” includes fish, bees, and wild fauna.

##### (2) Approval procedure

The term “approval procedure” means any registration, notification, or other mandatory administrative procedure for—

(A) approving the use of an additive for a stated purpose or under stated conditions, or

(B) establishing a tolerance for a stated purpose or under stated conditions for a contaminant,

in a food, beverage, or feedstuff prior to permitting the use of the additive or the marketing of a food, beverage, or feedstuff containing the additive or contaminant.

##### (3) Contaminant

The term “contaminant” includes pesticide and veterinary drug residues and extraneous matter.

##### (4) Control or inspection procedure

The term “control or inspection procedure” means any procedure used, directly or indirectly, to determine that a sanitary or phytosanitary measure is fulfilled, including sampling, testing, inspection, evaluation, verification, monitoring, auditing, assurance of conformity, accreditation, registration, certification, or other procedure involving the physical examination of a good, of the packaging of a good, or of the equipment or facilities di-

rectly related to production, marketing, or use of a good, but does not mean an approval procedure.

**(5) Plant**

The term “plant” includes wild flora.

**(6) Risk assessment**

The term “risk assessment” means an evaluation of—

(A) the potential for the introduction, establishment or spread of a pest or disease and associated biological and economic consequences; or

(B) the potential for adverse effects on human or animal life or health arising from the presence of an additive, contaminant, toxin or disease-causing organism in a food, beverage, or feedstuff.

**(7) Sanitary or phytosanitary measure**

**(A) In general**

The term “sanitary or phytosanitary measure” means a measure to—

(i) protect animal or plant life or health in the United States from risks arising from the introduction, establishment, or spread of a pest or disease;

(ii) protect human or animal life or health in the United States from risks arising from the presence of an additive, contaminant, toxin, or disease-causing organism in a food, beverage, or feedstuff;

(iii) protect human life or health in the United States from risks arising from a disease-causing organism or pest carried by an animal or plant, or a product thereof; or

(iv) prevent or limit other damage in the United States arising from the introduction, establishment, or spread of a pest.

**(B) Form**

The form of a sanitary or phytosanitary measure includes—

- (i) end product criteria;
- (ii) a product-related processing or production method;
- (iii) a testing, inspection, certification, or approval procedure;
- (iv) a relevant statistical method;
- (v) a sampling procedure;
- (vi) a method of risk assessment;
- (vii) a packaging and labeling requirement directly related to food safety; and
- (viii) a quarantine treatment, such as a relevant requirement associated with the transportation of animals or plants or with material necessary for their survival during transportation.

(Pub. L. 96-39, title IV, §463, as added Pub. L. 103-182, title III, §351(a), Dec. 8, 1993, 107 Stat. 2119.)

SUBPART 2—STANDARDS-RELATED MEASURES

**§ 2576. General**

**(a) No bar to engaging in standards activity**

Nothing in this subpart shall be construed—

(1) to prohibit a Federal agency from engaging in activity related to standards-related

measures, including any such measure relating to safety, the protection of human, animal, or plant life or health, the environment or consumers; or

(2) to limit the authority of a Federal agency to determine the level it considers appropriate of safety or of protection of human, animal, or plant life or health, the environment or consumers.

**(b) Exclusion**

This subpart does not apply to—

(1) technical specifications prepared by a Federal agency for production or consumption requirements of the agency; or

(2) sanitary or phytosanitary measures under subpart 1.

(Pub. L. 96-39, title IV, §471, as added Pub. L. 103-182, title III, §351(a), Dec. 8, 1993, 107 Stat. 2120.)

**§ 2576a. Inquiry point**

The standards information center maintained under section 2544 of this title shall, in addition to the functions specified therein, make available to the public relevant documents, at such reasonable fees as the Secretary of Commerce may prescribe, and information regarding—

(1) the membership and participation of the Federal Government, State governments, and relevant nongovernmental bodies in the United States in international and regional standardizing bodies and conformity assessment systems, and in bilateral and multilateral arrangements regarding standards-related measures, and the provisions of those systems and arrangements;

(2) the location of notices of the type required under article 909 of the NAFTA, or where the information contained in such notice can be obtained; and

(3) the Federal agency procedures for assessment of risk, and factors the agency considers in conducting the assessment and establishing the levels of protection that the agency considers appropriate.

(Pub. L. 96-39, title IV, §472, as added Pub. L. 103-182, title III, §351(a), Dec. 8, 1993, 107 Stat. 2120.)

**§ 2576b. Subpart definitions**

Notwithstanding section 2571 of this title, for purposes of this subpart—

**(1) Approval procedure**

The term “approval procedure” means any registration, notification, or other mandatory administrative procedure for granting permission for a good or service to be produced, marketed, or used for a stated purpose or under stated conditions.

**(2) Conformity assessment procedure**

The term “conformity assessment procedure” means any procedure used, directly or indirectly, to determine that a technical regulation or standard is fulfilled, including sampling, testing, inspection, evaluation, verification, monitoring, auditing, assurance of conformity, accreditation, registration, or ap-