CHAPTER 39—FAIR PACKAGING AND LABELING PROGRAM

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§1451. Congressional declaration of policy

Informed consumers are essential to the fair and efficient functioning of a free market economy. Packages and their labels should enable consumers to obtain accurate information as to the quantity of the contents and should facilitate value comparisons. Therefore, it is hereby declared to be the policy of the Congress to assist consumers and manufacturers in reaching these goals in the marketing of consumer goods.


Effective Date

Section 13 of Pub. L. 89–755 provided that: “This Act [enacting this chapter] shall take effect on July 1, 1967: Provided, That the Secretary (with respect to any consumer commodity which is a food, drug, device, or cosmetic, as those terms are defined by the Federal Food, Drug, and Cosmetic Act) [section 301 et seq. of Title 21, Food and Drugs], and the Commission (with respect to any other consumer commodity) may by regulation postpone, for an additional twelve-month period, the effective date of this Act [this chapter] with respect to any class or type of consumer commodity on the basis of a finding that such a postponement would be in the public interest.”

Short Title

Section 1 of Pub. L. 89–755 provided: “That this Act [enacting this chapter] may be cited as the ‘Fair Packaging and Labeling Act’.”
§1452. Unfair and deceptive packaging and labeling; scope of prohibition

(a) Nonconforming labels

It shall be unlawful for any person engaged in the packaging or labeling of any consumer commodity (as defined in this chapter) for distribution in commerce, or for any person (other than a common carrier for hire, a contract carrier for hire, or a freight forwarder for hire) engaged in the distribution in commerce of any packaged or labeled consumer commodity, to distribute or to cause to be distributed in commerce any such commodity if such commodity is contained in a package, or if there is affixed to that commodity a label, which does not conform to the provisions of this chapter and of regulations promulgated under the authority of this chapter.

(b) Exemptions

The prohibition contained in subsection (a) of this section shall not apply to persons engaged in business as wholesale or retail distributors of consumer commodities except to the extent that such persons (1) are engaged in the packaging or labeling of such commodities, or (2) prescribe or specify by any means the manner in which such commodities are packaged or labeled.


§1453. Requirements of labeling; placement, form, and contents of statement of quantity; supplemental statement of quantity

(a) Contents of label

No person subject to the prohibition contained in section 1452 of this title shall distribute or cause to be distributed in commerce any packaged consumer commodity unless in conformity with regulations which shall be established by the promulgating authority pursuant to section 1455 of this title which shall provide that—

(1) The commodity shall bear a label specifying the identity of the commodity and the name and place of business of the manufacturer, packer, or distributor;

(2) The net quantity of contents (in terms of weight or mass, measure, or numerical count) shall be separately and accurately stated in a uniform location upon the principal display panel of that label, using the most appropriate units of both the customary inch/pound system of measure, as provided in paragraph (3) of this subsection, and, except as provided in paragraph (3)(A)(ii) or paragraph (6) of this subsection, the SI metric system;

(3) The separate label statement of net quantity of contents appearing upon or affixed to any package—

(A)(i) if on a package labeled in terms of weight, shall be expressed in pounds, with any remainder in terms of ounces or common or decimal fractions of the pound; or in the case of liquid measure, in the largest whole unit (quarts, quarts and pints, or pints, as appropriate) with any remainder in terms of fluid ounces or common or decimal fractions of the pint or quart;

(ii) if on a random package, may be expressed in terms of pounds and decimal fractions of the pound carried out to not more than three decimal places and is not required to, but may, include a statement in terms of the SI metric system carried out to not more than three decimal places;

(iii) if on a package labeled in terms of linear measure, shall be expressed in terms of the largest whole unit (yards, yards and feet, or feet, as appropriate) with any remainder in terms
(iv) if on a package labeled in terms of measure of area, shall be expressed in terms of the largest whole square unit (square yards, square yards and square feet, or square feet, as appropriate) with any remainder in terms of square inches or common or decimal fractions of the square foot or square yard;

(B) shall appear in conspicuous and easily legible type in distinct contrast (by topography, layout, color, embossing, or molding) with other matter on the package;

(C) shall contain letters or numerals in a type size which shall be (i) established in relationship to the area of the principal display panel of the package, and (ii) uniform for all packages of substantially the same size; and

(D) shall be so placed that the lines of printed matter included in that statement are generally parallel to the base on which the package rests as it is designed to be displayed; and

(4) The label of any package of a consumer commodity which bears a representation as to the number of servings of such commodity contained in such package shall bear a statement of the net quantity (in terms of weight or mass, measure, or numerical count) of each such serving.

(5) For purposes of paragraph (3)(A)(ii) of this subsection the term “random package” means a package which is one of a lot, shipment, or delivery of packages of the same consumer commodity with varying weights or masses, that is, packages with no fixed weight or mass pattern.

(6) The requirement of paragraph (2) that the statement of net quantity of contents include a statement in terms of the SI metric system shall not apply to foods that are packaged at the retail store level.

(b) Supplemental statements

No person subject to the prohibition contained in section 1452 of this title shall distribute or cause to be distributed in commerce any packaged consumer commodity if any qualifying words or phrases appear in conjunction with the separate statement of the net quantity of contents required by subsection (a) of this section, but nothing in this subsection or in paragraph (2) of subsection (a) of this section shall prohibit supplemental statements, at other places on the package, describing in nondeceptive terms the net quantity of contents: Provided, That such supplemental statements of net quantity of contents shall not include any term qualifying a unit of weight or mass, measure, or count that tends to exaggerate the amount of the commodity contained in the package.


AMENDMENTS

1992—Pub. L. 102–245, §107, which directed amendment of section, effective two years after Feb. 14, 1992, by substituting “weight or mass” for “weight” in subssecs. (a)(2), (4), (5) and (b) and “weights or masses” for “weights” in subsec. (a)(5), by inserting “, using the most appropriate units of the SI metric system as the primary system for measuring quantity” after “panel of that label” in subsec. (a)(2), by substituting “that also displays the avoirdupois system of measure, and that contains” for “containing” in subsec. (a)(3)(A)(i), by inserting “that also displays the avoirdupois system of measure” after “random package” in subsec. (a)(3)(A)(ii), by inserting “that also displays the avoirdupois system of measure” after “linear measure” in subsec. (a)(3)(A)(iii), and by inserting “that also displays the avoirdupois system of measure” in subsec. (a)(3)(A)(iv), was repealed by Pub. L. 102–329, §3.

Subsec. (a)(2). Pub. L. 102–329, §1(1), (3), substituted “weight or mass” for “weight” and inserted before semicolon at end “, using the most appropriate units of both the customary inch/pound system of measure, as provided in paragraph (3) of this subsection, and, except as provided in paragraph (3)(A)(iii) or paragraph
(6) of this subsection, the SI metric system”.

Subsec. (a)(3)(A)(i). Pub. L. 102–329, §1(4)(A), substituted “labeled in terms of weight, shall be expressed in pounds” for “containing less than four pounds or one gallon and labeled in terms of weight or fluid measure, shall, unless subparagraph (ii) applies and such statement is set forth in accordance with such subparagraph, be expressed both in ounces (with identification as to avoirdupois or fluid ounces) and, if applicable, in pounds for weight units”.

Subsec. (a)(3)(A)(ii). Pub. L. 102–329, §1(4)(B), (C), substituted “three” for “two” and inserted before semicolon at end “and is not required to, but may, include a statement in terms of the SI metric system carried out to not more than three decimal places”.


Subsec. (a)(4). Pub. L. 102–329, §1(1), substituted “weight or mass” for “weight”.

Subsec. (a)(5). Pub. L. 102–329, §1(1), (2), substituted “weight or mass” for “weight” and “weights or masses” for “weights”.


Subsec. (b). Pub. L. 102–329, §1(1), substituted “weight or mass” for “weight”.

Effective Date of 1992 Amendments

Section 2 of Pub. L. 102–329 provided that: “The amendments made by section 1 [amending this section and section 1454 of this title] shall take effect on February 14, 1994. The amendments made by section 1 shall have no effect on the sale or distribution of products whose labels have been printed before such effective date. Nothing in the amendments made by section 1 shall apply to unit pricing, advertising, recipe programs, nutrition labeling, or other general pricing information. Nothing in the amendments made by section 1 shall be construed to require changes in package size or to affect in any way the size of packages.”

Section 107(b) of Pub. L. 102–245, which provided that section 107 of Pub. L. 102–245 which amended this section and section 1454 of this title was to take effect 2 years after Feb. 14, 1992, was repealed by Pub. L. 102–329, §3, Aug. 3, 1992, 106 Stat. 848.

§1454. Rules and regulations

(a) Promulgating authority

The authority to promulgate regulations under this chapter is vested in (A) the Secretary of Health and Human Services (referred to hereinafter as the “Secretary”) with respect to any consumer commodity which is a food, drug, device, or cosmetic, as each such term is defined by section 321 of title 21; and (B) the Federal Trade Commission (referred to hereinafter as the “Commission”) with respect to any other consumer commodity.

(b) Exemption of commodities from regulations

If the promulgating authority specified in this section finds that, because of the nature, form, or quantity of a particular consumer commodity, or for other good and sufficient reasons, full compliance with all the requirements otherwise applicable under section 1453 of this title is impracticable or is not necessary for the adequate protection of consumers, the Secretary or the Commission (whichever the case may be) shall promulgate regulations exempting such commodity from those requirements to the extent and under such conditions as the promulgating authority determines to be consistent with section 1451 of this title.

(c) Scope of additional regulations

Whenever the promulgating authority determines that regulations containing prohibitions or requirements other than those prescribed by section 1453 of this title are necessary to prevent the
deception of consumers or to facilitate value comparisons as to any consumer commodity, such
authority shall promulgate with respect to that commodity regulations effective to—

(1) establish and define standards for characterization of the size of a package enclosing any
consumer commodity, which may be used to supplement the label statement of net quantity of
contents of packages containing such commodity, but this paragraph shall not be construed as
authorizing any limitation on the size, shape, weight or mass, dimensions, or number of packages
which may be used to enclose any commodity;

(2) regulate the placement upon any package containing any commodity, or upon any label
affixed to such commodity, of any printed matter stating or representing by implication that such
commodity is offered for retail sale at a price lower than the ordinary and customary retail sale
price or that a retail sale price advantage is accorded to purchasers thereof by reason of the size
of that package or the quantity of its contents;

(3) require that the label on each package of a consumer commodity (other than one which is
a food within the meaning of section 321(f) of title 21) bear (A) the common or usual name of
such consumer commodity, if any, and (B) in case such consumer commodity consists of two or
more ingredients, the common or usual name of each such ingredient listed in order of
decreasing predominance, but nothing in this paragraph shall be deemed to require that any
trade secret be divulged; or

(4) prevent the nonfunctional-slack-fill of packages containing consumer commodities.

For purposes of paragraph (4) of this subsection, a package shall be deemed to be
nonfunctionally slack-filled if it is filled to substantially less than its capacity for reasons other than
(A) protection of the contents of such package or (B) the requirements of machines used for
enclosing the contents in such package.

(d) Development by manufacturers, packers, and distributors of voluntary product standards

Whenever the Secretary of Commerce determines that there is undue proliferation of the
weights or masses, measures, or quantities in which any consumer commodity or reasonably
comparable consumer commodities are being distributed in packages for sale at retail and such
undue proliferation impairs the reasonable ability of consumers to make value comparisons with
respect to such consumer commodity or commodities, he shall request manufacturers, packers,
and distributors of the commodity or commodities to participate in the development of a voluntary
product standard for such commodity or commodities under the procedures for the development
of voluntary products standards established by the Secretary pursuant to section 272 of this title.
Such procedures shall provide adequate manufacturer, packer, distributor, and consumer
representation.

(e) Report and recommendations to Congress upon industry failure to develop or abide by
voluntary product standards

If (1) after one year after the date on which the Secretary of Commerce first makes the request
of manufacturers, packers, and distributors to participate in the development of a voluntary
product standard as provided in subsection (d) of this section, he determines that such a standard
will not be published pursuant to the provisions of such subsection (d), or (2) if such a standard is
published and the Secretary of Commerce determines that it has not been observed, he shall
promptly report such determination to the Congress with a statement of the efforts that have been
made under the voluntary standards program and his recommendation as to whether Congress
should enact legislation providing regulatory authority to deal with the situation in question.


AMENDMENTS

1992—Pub. L. 102–245, §107(a)(1), (2), (b), which directed amendment of section, effective two years after Feb. 14, 1992, by substituting “weight or mass” for “weight” in subsec. (c)(1) and “weights or masses” for “weights” in subsec. (d), was repealed by Pub. L. 102–329, §3.

Subsec. (c)(1). Pub. L. 102–329, §1(1), substituted “weight or mass” for “weight”.

Subsec. (d). Pub. L. 102–329, §1(2), substituted “weights or masses” for “weights”.

CHANGE OF NAME

“Secretary of Health and Human Services” substituted for “Secretary of Health, Education, and Welfare” in subsec. (a) pursuant to section 509(b) of Pub. L. 96–88, which is classified to section 3508(b) of Title 20, Education.

EFFECTIVE DATE OF 1992 AMENDMENT

Amendment by Pub. L. 102–329 effective Feb. 14, 1994, but with such amendment to have no effect on the sale or distribution of products whose labels have been printed before such date, no application to unit pricing, advertising, recipe programs, nutrition labeling, or other general pricing information, and no construction requiring changes in package size or affecting in any way the size of packages, see section 2 of Pub. L. 102–329, set out as a note under section 1453 of this title.

§1455. Procedure for promulgation of regulations

(a) Hearings by Secretary of Health and Human Services

Regulations promulgated by the Secretary under section 1453 or 1454 of this title shall be promulgated, and shall be subject to judicial review, pursuant to the provisions of subsections (e), (f), and (g) of section 371 of title 21. Hearings authorized or required for the promulgation of any such regulations by the Secretary shall be conducted by the Secretary or by such officer or employees of the Department of Health and Human Services as he may designate for that purpose.

(b) Judicial review; hearings by Federal Trade Commission

Regulations promulgated by the Commission under section 1453 or 1454 of this title shall be promulgated, and shall be subject to judicial review, by proceedings taken in conformity with the provisions of subsections (e), (f), and (g) of section 371 of title 21 in the same manner, and with the same effect, as if such proceedings were taken by the Secretary pursuant to subsection (a) of this section. Hearings authorized or required for the promulgation of any such regulations by the Commission shall be conducted by the Commission or by such officer or employee of the Commission as the Commission may designate for that purpose.

(c) Cooperation with other departments and agencies

In carrying into effect the provisions of this chapter, the Secretary and the Commission are authorized to cooperate with any department or agency of the United States, with any State, Commonwealth, or possession of the United States, and with any department, agency, or political subdivision of any such State, Commonwealth, or possession.

(d) Returnable or reusable glass containers for beverages

No regulation adopted under this chapter shall preclude the continued use of returnable or reusable glass containers for beverages in inventory or with the trade as of the effective date of this Act, nor shall any regulation under this chapter preclude the orderly disposal of packages in inventory or with the trade as of the effective date of such regulation.
§1456. Enforcement

(a) Misbranded consumer commodities

Any consumer commodity which is a food, drug, device, or cosmetic, as each such term is defined by section 201 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321), and which is introduced or delivered for introduction into commerce in violation of any of the provisions of this chapter, or the regulations issued pursuant to this chapter, shall be deemed to be misbranded within the meaning of chapter III of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 331 et seq.], but the provisions of section 303 of that Act (21 U.S.C. 333) shall have no application to any violation of section 1452 of this title.

(b) Unfair or deceptive acts or practices in commerce

Any violation of any of the provisions of this chapter, or the regulations issued pursuant to this chapter, with respect to any consumer commodity which is not a food, drug, device, or cosmetic, shall constitute an unfair or deceptive act or practice in commerce in violation of section 45(a) of this title and shall be subject to enforcement under section 45(b) of this title.

(c) Imports

In the case of any imports into the United States of any consumer commodity covered by this chapter, the provisions of sections 1453 and 1454 of this title shall be enforced by the Secretary of the Treasury pursuant to section 801(a) and (b) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 381).


References in Text

The Federal Food, Drug, and Cosmetic Act, referred to in subsecs. (a) and (c), is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended. Chapter III of the Act is classified generally to subchapter III (§331 et seq.) of chapter 9 of Title 21, Food and Drugs. For complete classification of this Act to the Code, see section 301 of Title 21 and Tables.

§1457. Omitted

Codification

Congress describing activities carried out for the administration and enforcement of this chapter, terminated, effective May 15, 2000, pursuant to section 3003 of Pub. L. 104–66, as amended, set out as a note under section 1113 of Title 31, Money and Finance. See, also, pages 54, 92, and 172 of House Document No. 103–7.

§1458. Cooperation with State authorities; transmittal of regulations to States; noninterference with existing programs

(a) A copy of each regulation promulgated under this chapter shall be transmitted promptly to the Secretary of Commerce, who shall (1) transmit copies thereof to all appropriate State officers and agencies, and (2) furnish to such State officers and agencies information and assistance to promote to the greatest practicable extent uniformity in State and Federal regulation of the labeling of consumer commodities.

(b) Nothing contained in this section shall be construed to impair or otherwise interfere with any program carried into effect by the Secretary of Health and Human Services under other provisions of law in cooperation with State governments or agencies, instrumentalities, or political subdivisions thereof.


CHANGE OF NAME

“Secretary of Health and Human Services” substituted for “Secretary of Health, Education, and Welfare” in subsec. (b) pursuant to section 509(b) of Pub. L. 96–88, which is classified to section 3508(b) of Title 20, Education.

§1459. Definitions

For the purpose of this chapter—

(a) The term “consumer commodity”, except as otherwise specifically provided by this subsection, means any food, drug, device, or cosmetic (as those terms are defined by the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.]), and any other article, product, or commodity of any kind or class which is customarily produced or distributed for sale through retail sales agencies or instrumentalities for consumption by individuals, or use by individuals for purposes of personal care or in the performance of services ordinarily rendered within the household, and which usually is consumed or expended in the course of such consumption or use. Such term does not include—

(1) any meat or meat product, poultry or poultry product, or tobacco or tobacco product;
(2) any commodity subject to packaging or labeling requirements imposed by the Secretary of Agriculture pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act [7 U.S.C. 136 et seq.], or the provisions of the eighth paragraph under the heading “Bureau of Animal Industry” of the Act of March 4, 1913 [21 U.S.C. 151 et seq.], commonly known as the Virus-Serum-Toxin Act;
(3) any drug subject to the provisions of section 503(b)(1) or 506 of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 353(b)(1) and 356];
(4) any beverage subject to or complying with packaging or labeling requirements imposed under the Federal Alcohol Administration Act [27 U.S.C. 201 et seq.]; or
(5) any commodity subject to the provisions of the Federal Seed Act [7 U.S.C. 1551 et seq.].

(b) The term “package” means any container or wrapping in which any consumer commodity is
enclosed for use in the delivery or display of that consumer commodity to retail purchasers, but

(1) shipping containers or wrappings used solely for the transportation of any consumer commodity in bulk or in quantity to manufacturers, packers, or processors, or to wholesale or retail distributors thereof;

(2) shipping containers or outer wrappings used by retailers to ship or deliver any commodity to retail customers if such containers and wrappings bear no printed matter pertaining to any particular commodity; or


(c) The term “label” means any written, printed, or graphic matter affixed to any consumer commodity or affixed to or appearing upon a package containing any consumer commodity.

(d) The term “person” includes any firm, corporation, or association.

(e) The term “commerce” means (1) commerce between any State, the District of Columbia, the Commonwealth of Puerto Rico, or any territory or possession of the United States, and any place outside thereof, and (2) commerce within the District of Columbia or within any territory or possession of the United States not organized with a legislative body, but shall not include exports to foreign countries.

(f) The term “principal display panel” means that part of a label that is most likely to be displayed, presented, shown, or examined under normal and customary conditions of display for retail sale.


REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in subsec. (a), 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.


The Virus-Serum-Toxin Act, referred to in subsec. (a)(2), is the eighth paragraph under the heading “Bureau of Animal Industry” of act Mar. 4, 1913, ch. 145, 37 Stat. 832, as amended, which is classified generally to chapter 5 (§151 et seq.) of Title 21, Food and Drugs. For complete classification of this Act to the Code, see Short Title note set out under section 151 of Title 21 and Tables.

The Federal Alcohol Administration Act, referred to in subsec. (a)(4), is act Aug. 29, 1935, ch. 814, 49 Stat. 977, as amended, which is classified generally to chapter 8 (§201 et seq.) of Title 27, Intoxicating Liquors. For complete classification of this Act to the Code, see section 201 of Title 27 and Tables.

The Federal Seed Act, referred to in subsec. (a)(5), is act Aug. 9, 1939, ch. 615, 53 Stat. 1275, as amended, which is classified generally to chapter 37 (§1551 et seq.) of Title 7, Agriculture. For complete classification of this Act to the Code, see section 1551 of Title 7 and Tables.

AMENDMENTS


EFFECTIVE DATE OF 1968 AMENDMENT

Amendment by Pub. L. 90–628 effective 60 days after Oct. 22, 1968, see section 3 of Pub. L. 90–628, set out as a note under section 251 of this title.
§1460. Savings provisions

Nothing contained in this chapter shall be construed to repeal, invalidate, or supersede—

(a) the Federal Trade Commission Act [15 U.S.C. 41 et seq.] or any statute defined therein as an antitrust Act;

(b) the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.]; or

(c) the Federal Hazardous Substances Labeling Act [15 U.S.C. 1261 et seq.].


REFERENCES IN TEXT

The Federal Trade Commission Act, referred to in text, is act Sept. 26, 1914, ch. 311, 38 Stat. 717, as amended, which is classified generally to subchapter I (§41 et seq.) of chapter 2 of this title. For complete classification of this Act to the Code, see section 58 of this title and Tables.

The Federal Food, Drug, and Cosmetic Act, referred to in text, 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.

The Federal Hazardous Substances Labeling Act, referred to in text, is Pub. L. 86–613, July 12, 1960, 74 Stat. 372, as amended, which is classified generally to chapter 30 (§1261 et seq.) of this title. For complete classification of this Act to the Code, see Short Title note set out under section 1261 of this title and Tables.

§1461. Effect upon State law

It is hereby declared that it is the express intent of Congress to supersede any and all laws of the States or political subdivisions thereof insofar as they may now or hereafter provide for the labeling of the net quantity of contents of the package of any consumer commodity covered by this chapter which are less stringent than or require information different from the requirements of section 1453 of this title or regulations promulgated pursuant thereto.