1 A bill to be entitled

An act relating to labeling of genetically engineered foods; creating s. 500.92, F.S.; providing definitions; providing mandatory labeling requirements for genetically engineered raw foods and processed foods made with or derived from genetically engineered ingredients by a specified date; exempting specified foods, commodities, ingredients, and other substances from the labeling requirements; directing the Department of Health to adopt rules; providing for enforcement of the labeling requirements; providing administrative and civil remedies and penalties; providing legislative intent with regard to such penalties; providing for injunctive relief actions; requiring the court to award costs and fees under certain circumstances; specifying that injunctive relief actions do not preclude civil actions for damages or personal injury; providing an effective date.

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WHEREAS, Florida has the right to protect the liberty of its citizens to be free to make the most fundamental of life choices of what to eat and put on their tables to feed their families, and

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WHEREAS, the Legislature finds that consumers should have the right to know whether the foods they purchase contain

Page 1 of 11

genetically engineered material, and

WHEREAS, without mandatory labeling of genetically engineered foods, consumers may unknowingly violate their own dietary or religious principles, and

WHEREAS, the lack of labeling denies health professionals the ability to trace potential toxic or allergic reactions to, and other adverse health effects from, genetically engineered food, and

WHEREAS, labeling requirements for genetically engineered foods are needed to facilitate both the withdrawal of products where unforeseen adverse effects on human health, animal health, or the environment, including ecosystems, are established, and the targeting of monitoring to examine potential effects on health and the environment, and

WHEREAS, many medical and public health groups still have questions regarding the potential long-term impact of genetically engineered foods on human health and the environment, and

WHEREAS, many medical and public health groups, including, but not limited to, the American College of Physicians, American Public Health Association, American Nurses Association, British Medical Association, Australian Medical Association, Irish Medical Organization, and German Medical Association, have passed resolutions or otherwise supported the mandatory labeling of genetically engineered foods to facilitate further health research, and

Page 2 of 11

WHEREAS, sixty-four developed or developing nations have banned, restricted, or required labeling of products that are genetically engineered, and

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WHEREAS, Floridians should have the same freedom to make informed choices about the food they eat as consumers or grow and offer to market as farmers, and

WHEREAS, no international agreement prohibits the mandatory labeling of genetically engineered foods, and

WHEREAS, the cultivation of genetically engineered crops can negatively impact the environment, in some cases necessitating the use of increasingly toxic herbicides that can damage agricultural areas, impair drinking water, and pose health risks to consumers and farmworkers, and

WHEREAS, consumers should have the choice to avoid purchasing foods that they believe cause adverse health and environmental effects, and

WHEREAS, currently, there is no federal requirement mandating disclosure of genetically engineered foods on food labels, NOW, THEREFORE,

Be It Enacted by the Legislature of the State of Florida:

Section 1. Section 500.92, Florida Statutes, is created to read:

- 500.92 Genetically engineered foods.-
- (1) As used in this section, the term:

Page 3 of 11

(a) "Department" means the Department of Health.

- (b) "Food facility" means an operation that stores, prepares, packages, serves, vends, or otherwise provides food for human consumption at the retail level, including an operation where food is consumed on or off the premises, regardless of whether there is a charge for the food.
- (c) "Genetically engineered" means any food that consists of, is composed of, contains, or is produced from an organism or organisms in which the genetic material has been changed, commonly referred to as a "genetically modified organism" or "GMO," through the application of:
- 1. In vitro nucleic acid techniques, including recombinant deoxyribonucleic acid techniques and the direct injection of nucleic acid into cells or organelles. Such techniques include, but are not limited to, recombinant deoxyribonucleic acid or ribonucleic acid techniques that use vector systems and techniques involving the direct introduction into the organisms of hereditary material prepared outside the organisms, such as microinjection, macroinjection, chemoporation, electroporation, microencapsulation, and liposome fusion; or
- 2. Fusion of cells, including protoplast fusion, or hybridization techniques that overcome natural physiological, reproductive, or recombination barriers, where the donor cells or protoplasts do not fall within the same taxonomic family, in a way that does not occur by natural multiplication or natural recombination.

The term does not include the centuries-old hybridization technique used by farmers and breeders relying on nature or similar plant-to-plant or similar animal-to-animal selective breeding.

- 110 (d) "Ingredient" means any substance that is used in the
 111 manufacture, or contained in the final form, of a processed
 112 food.
 - (e) "Processed food" means any food other than a raw agricultural commodity and includes any food produced from a raw agricultural commodity that has been subject to processing, such as canning, smoking, pressing, cooking, freezing, dehydration, fermentation, or milling.
 - (2) Beginning January 1, 2018:
 - (a) Any genetically engineered raw food that is offered for retail sale must include a clear and conspicuous statement with the words "genetically engineered" on the front package or label of any such commodity. For such a commodity that is not separately packaged or labeled, the statement must appear on a label on the retail store shelf or bin where the commodity is displayed for sale.
 - (b) Any package offered for retail sale containing processed food that is made with or derived from any genetically engineered ingredient or is produced from a source that contains recombinant bovine growth hormone must include a clear and conspicuous statement on the front or back of the package with

Page 5 of 11

the words "contains genetically engineered ingredients,"

followed by the name of the genetically engineered ingredient or

ingredients. If an ingredients list appears on the package, the

statement must appear underneath the ingredients list. For a

processed food containing more than one genetically engineered

ingredient or recombinant bovine growth hormone, the genetically

engineered ingredients listed after the statement must be listed

in the same order in which they appear in the full ingredients

list.

- (c) In lieu of compliance with paragraph (b), any package containing processed food that is made with or derived from any ingredient that may be genetically engineered or is produced from a source that contains recombinant bovine growth hormone must include a clear and conspicuous statement on the front or back of the package with the words "may contain genetically engineered ingredients," followed by the name of the genetically engineered ingredient or ingredients. If an ingredients list appears on the package, the statement must appear underneath the ingredients list. For a processed food containing more than one ingredient that may be genetically engineered, the genetically engineered ingredients listed after the statement must be listed in the same order in which they appear in the full ingredients list.
- (d) Except as set forth in paragraph (e), a food produced entirely or in part from genetic engineering may not be labeled on the package, in signage, or in advertising as "natural" or

Page 6 of 11

157 with any words of similar import.

- (e) This subsection does not apply to:
- 1. Food consisting entirely of, or derived entirely from, an animal that has not itself been genetically engineered and that has not been fed a feed containing more than 1.5 percent genetically engineered ingredients.
- 2. A raw agricultural commodity or ingredient that has been grown, raised, or produced without the knowing and intentional use of genetically engineered seed or food. The person responsible for complying with this section must obtain, from whoever sold the commodity or ingredient to such person, a sworn statement that the commodity or ingredient has not been knowingly or intentionally genetically engineered and has been segregated from, and not been knowingly or intentionally commingled with, goods that may have been genetically engineered at any time. The sworn statement must be notarized and include a written declaration stating that such statement is made under the penalties of perjury and fraud. In providing such a sworn statement, a person may rely on a sworn statement from his or her own supplier that contains such an affirmation.
- 3. An alcoholic beverage that is subject to regulation under chapters 561 through 568.
- 4. A processed food that would be subject to this section solely because it includes one or more genetically engineered ingredients, if a single genetically engineered ingredient does not account for more than one-half of 1 percent of the total

Page 7 of 11

183 weight of the processed food.

- 5. Any food not knowingly and intentionally produced from or commingled with genetically engineered seed or genetically engineered food, as determined by an independent organization, such as the Non-GMO Project, if such a determination has been made pursuant to a sampling and testing procedure approved for this purpose in rules adopted by the department.
- 6. Food that has been lawfully certified to be labeled, marketed, and offered for sale as organic pursuant to applicable federal organic food production laws and regulations.
 - 7. Food that is not packaged for retail sale and that is:
- <u>a. A processed food prepared and intended for immediate</u>
 human consumption;
- b. Served, sold, or otherwise provided in a restaurant or other food facility that is primarily engaged in the sale of food prepared and intended for immediate human consumption; or
 - c. Medical food, as defined in 21 U.S.C. s. 360ee(b)(3).
 - (3) (a) The department shall:
 - 1. Adopt rules to administer this section.
- 2. Select an independent nonprofit organization to approve a sampling and testing procedure consistent with sampling and testing principles recommended and developed by independent nonprofit organizations with the highest internationally recognized standards of genetically engineered labeling requirements. The organization shall be chosen on a 2-year basis by agency rule.

Page 8 of 11

3. Create an educational pamphlet regarding the requirements of this section for distribution to farmers in the state.

4. Prominently display on its website information regarding:

- a. Information regarding genetically engineered foods and crops as well as organic foods and crops.
- b. Standards for nongenetically engineered products developed by independent nonprofit organizations with the highest internationally recognized standards of genetically engineered labeling requirements.
- c. Penalties imposed under this subsection and any pending cases.
- (b) After exhausting administrative remedies under chapter 120, the department may bring an action in a court of competent jurisdiction to enjoin a person or an entity violating this section.
- (c) The department may assess a civil penalty against a person or an entity violating this section in an amount not to exceed \$5,000 per seed and \$1,000 per retail package intended to be sold by a retailer. Each day of violation is considered a separate violation. Minimum penalties per day will be based on 3 percent of the annual profit of the violating entity. It is the intent of the Legislature that such penalties are imposed to prevent violations of this section and that the cost of such penalties is not passed on to consumers as the cost of doing

Page 9 of 11

235 <u>business.</u>

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- (d) Any political subdivision or municipality of the state or a citizen of the state may maintain an action for injunctive relief against:
- The department to compel it to enforce this section or any rules adopted thereunder. As a condition precedent to the institution of an action pursuant to this subparagraph, the complaining party must first file with the department a verified complaint setting forth the facts upon which the complaint is based and the manner in which the complaining party is affected. Within 7 days after receipt of a complaint, the department must transmit, by registered or certified mail, a copy of the complaint to those parties charged with violating this section or rules adopted thereunder. The department shall have 30 days after the receipt of a complaint to take appropriate action. If such action is not taken within the time prescribed, the complaining party may institute the judicial proceedings authorized in this subparagraph. However, a complainant's failure to comply with this subparagraph does not bar an action for a temporary restraining order to prevent immediate and irreparable harm from the conduct or activity for which a complaint is made. In any action instituted pursuant to this subparagraph, the court, in the interest of justice, may add the department as a party defendant.
- 2. Any person, natural or corporate, or governmental agency or authority to enjoin such persons, agencies, or

Page 10 of 11

262	thereunder.
263	(e) In any successful action to enforce a provision of
264	this section, the court shall award the prevailing party, other
265	than the state, reasonable costs and attorney fees.
266	(f) Paragraph (d) does not preclude any person from
267	bringing civil action for damages or personal injury relating to

authorities from violating this section or rules adopted

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Section 2. This act shall take effect July 1, 2016.

Page 11 of 11

CODING: Words stricken are deletions; words underlined are additions.

violations of this section.