



General Assembly

January Session, 2013

***Raised Bill No. 6519***

LCO No. 3817



Referred to Committee on PUBLIC HEALTH

Introduced by:  
(PH)

***AN ACT CONCERNING THE LABELING OF GENETICALLY ENGINEERED FOOD.***

Be it enacted by the Senate and House of Representatives in General Assembly convened:

1 Section 1. Section 21a-92 of the general statutes is repealed and the  
2 following is substituted in lieu thereof (*Effective October 1, 2013*):

3 For the purposes of this chapter, [and] section 21a-65 and sections 2  
4 and 3 of this act, the following terms shall have the meanings  
5 hereinafter specified:

6 (1) "Advertisement" means all representations disseminated in any  
7 manner or by any means, other than by labeling, for the purpose of  
8 inducing, or which are likely to induce, directly or indirectly, the  
9 purchase of food, drugs, devices or cosmetics;

10 (2) (A) "Color additive" means a material which (i) is a dye, pigment  
11 or other substance made by a process of synthesis or similar artifice, or  
12 extracted, isolated or otherwise derived, with or without intermediate  
13 or final change of identity, from a vegetable, animal, mineral or other

14 source, and (ii) when added or applied to a food, drug or cosmetic, or  
15 to the human body or any of its parts, is capable, alone or through  
16 reaction with other substance, of imparting color thereto, except that  
17 the term "color additive" does not include any material exempted by  
18 regulation under the federal act, or which the commissioner, by  
19 regulation, determines is used, or intended to be used, solely for a  
20 purpose or purposes other than coloring; (B) the term "color" includes  
21 black, white and intermediate grays, as well as all other colors; (C)  
22 nothing in subparagraph (A) of this subdivision shall be construed to  
23 apply to any pesticide chemical, soil or plant nutrient, or other  
24 agricultural chemical used, or intended to be used, solely because of its  
25 effect in aiding, retarding or otherwise affecting, directly or indirectly,  
26 the growth or other natural physiological processes of produce of the  
27 soil which thereby affects its color, whether before or after harvest;

28 (3) "Commissioner" means the Commissioner of Consumer  
29 Protection;

30 (4) "Contaminated with filth" applies to any food, drug, device or  
31 cosmetic not securely protected from dust or dirt, and as far as may be  
32 necessary, by all reasonable means, from all foreign or injurious  
33 contaminations;

34 (5) "Cosmetic" means (A) articles intended to be rubbed, poured,  
35 sprinkled or sprayed on, introduced into, or otherwise applied to the  
36 human body or any of its parts for cleansing, beautifying, promoting  
37 attractiveness or altering the appearance and (B) articles intended for  
38 use as a component of any such articles; except that such term shall not  
39 include soap;

40 (6) "Device", except when used in subdivision (15) of this section  
41 and in subsection (i) of section 21a-93, [subsection (f)] subdivision 10 of  
42 section 21a-102, as amended by this act, subsection (c) of section 21a-  
43 106 and subsection (c) of section 21a-112, means instruments,  
44 apparatus and contrivances, including their components, parts and

45 accessories, intended (A) for use in the diagnosis, cure, mitigation,  
46 treatment or prevention of disease in man or other animals or (B) to  
47 affect the structure or any function of the body of man or other  
48 animals;

49 (7) "Director" means the director of the agricultural experiment  
50 station;

51 (8) "Drug" means (A) articles recognized in the official United States  
52 Pharmacopoeia, official Homeopathic Pharmacopoeia of the United  
53 States or official National Formulary, or any supplement to any of  
54 them; (B) articles intended for use in the diagnosis, cure, mitigation,  
55 treatment or prevention of disease in man or other animals; (C)  
56 articles, other than food, intended to affect the structure or any  
57 function of the body of man or any other animal; and (D) articles  
58 intended for use as a component of any articles specified in this  
59 subdivision; but shall not include devices or their components, parts or  
60 accessories;

61 (9) "Federal act" means the federal Food, Drug and Cosmetic Act, as  
62 amended, Title 21 USC 301 et seq.: 52 Stat. 1040 et seq.;

63 (10) "Food" means (A) articles used for food or drink for man or  
64 other animals, and (B) chewing gum, and (C) articles used for  
65 components of any such article;

66 (11) "Food additive" means any substance the intended use of which  
67 results or reasonably may be expected to result, directly or indirectly,  
68 in its becoming a component or otherwise affecting the characteristics  
69 of any food, including any substance intended for use in producing,  
70 manufacturing, packing, processing, preparing, treating, packaging,  
71 transporting or holding food; and including any source of radiation  
72 intended for any such use, if such substance is not generally  
73 recognized, among experts qualified by scientific training and  
74 experience to evaluate its safety, as having been adequately shown  
75 through scientific procedures or, in the case of a substance used in

76 food prior to January 1, 1958, through either scientific procedures or  
77 experience based on common use in food, to be safe under the  
78 conditions of its intended use; except that such term does not include  
79 (A) a pesticide chemical in or on a raw agricultural commodity; or (B) a  
80 pesticide chemical to the extent that it is intended for use or is used in  
81 the production, storage or transportation of any raw agricultural  
82 commodity; or (C) a color additive; or (D) any substance used in  
83 accordance with a sanction or approval granted prior to June 12, 1963,  
84 or the federal Food, Drug and Cosmetic Act, the Poultry Products  
85 Inspection Act (21 USC 451 et seq.) or the Meat Inspection Act of  
86 March 4, 1907, as amended;

87 (12) "Immediate container" shall not include package liners;

88 (13) "Intrastate commerce" means any and all commerce within the  
89 state of Connecticut and subject to its jurisdiction, and shall include the  
90 operation of any business or service establishment;

91 (14) "Label" means a display of written, printed or graphic matter  
92 upon the immediate container of any article, provided a requirement  
93 made by or under authority of this chapter that any information or  
94 other word or statement appear on the label shall not be considered to  
95 be complied with unless such information or other word or statement  
96 also appears on the outside container or wrapper, if any, of the retail  
97 package of such article, or is easily legible through the outside  
98 container or wrapper;

99 (15) "Labeling" means all labels and other written, printed or  
100 graphic matter (A) upon any article or any of its containers or  
101 wrappers, or (B) accompanying such article; provided, if an article is  
102 alleged to be misbranded because the labeling is misleading, or if an  
103 advertisement is alleged to be false because it is misleading, then, in  
104 determining whether the labeling or advertisement is misleading, there  
105 shall be taken into account, among other things, not only  
106 representations made or suggested by statement, word, design, device

107 or sound, or any combination thereof, but also the extent to which the  
108 labeling or advertisement fails to reveal facts material in the light of  
109 such representations or material with respect to consequences which  
110 may result from the use of the article to which the labeling or  
111 advertisement relates under the conditions of use prescribed in the  
112 labeling or advertisement thereof or under such conditions of use as  
113 are customary or usual, and provided the representation of a drug, in  
114 its labeling or advertisement, as an antiseptic shall be considered to be  
115 a representation that it is a germicide, except in the case of a drug  
116 purporting to be, or represented as, an antiseptic for inhibitory use as a  
117 wet dressing, ointment or dusting powder or for such other use as  
118 involves prolonged contact with the body;

119 (16) "Natural food" means food (A) which has not been treated with  
120 preservatives, antibiotics, synthetic additives, artificial flavoring or  
121 artificial coloring; [and] (B) which has not been processed in a manner  
122 that makes such food significantly less nutritive; and (C) which has not  
123 been genetically engineered, as defined in section 2 of this act.  
124 Processing of food by extracting, purifying, heating, fermenting,  
125 concentrating, dehydrating, cooling or freezing shall not, of itself,  
126 prevent the designation of such food as "natural food";

127 (17) "New drug" means (A) any drug the composition of which is  
128 such that such drug is not generally recognized, among experts  
129 qualified by scientific training and experience to evaluate the safety  
130 and effectiveness of drugs, as safe and effective for use under the  
131 conditions prescribed, recommended or suggested in its labeling or (B)  
132 any drug the composition of which is such that such drug, as a result  
133 of investigation to determine its safety and effectiveness for use under  
134 such conditions, has become so recognized, but which has not,  
135 otherwise than in such investigations, been used to a material extent or  
136 for a material time under such conditions, except that the provisions of  
137 this subsection pertaining to "effectiveness" shall not apply to any drug  
138 which (i) was commercially sold or used in the United States on  
139 October 9, 1962, (ii) was not a new drug as defined by this subsection

140 prior to the enactment of these provisions, and (iii) was not covered by  
141 an effective application under section 21a-110 or under Section 355 of  
142 the federal act, when such drug is intended solely for use under  
143 conditions prescribed, recommended, or suggested in labeling with  
144 respect to such drug on whichever of the above dates is applicable;

145 (18) "Official compendium" means the official United States  
146 Pharmacopoeia, official Homeopathic Pharmacopoeia of the United  
147 States, official National Formulary, or any supplement to any of them;

148 (19) "Organically grown" means produced through organic farming  
149 methods, which involve a system of ecological soil management and  
150 mechanical or biological methods to control insects, weeds, pathogens  
151 and other pests and which rely on crop rotation, crop residues,  
152 composted animal manures, legumes, green manures, composted  
153 organic waste or mineral-bearing rocks and not genetically engineered,  
154 as defined in section 2 of this act;

155 (20) "Person" includes any individual, partnership, corporation,  
156 limited liability company or association;

157 (21) "Pesticide chemical" means any substance which, alone, in  
158 chemical combination or in formulation with one or more other  
159 substances is an "economic poison" within the meaning of the federal  
160 Insecticide, Fungicide and Rodenticide Act, 7 USC 135-135k, and  
161 which is used in the production, storage or transportation of raw  
162 agricultural commodities;

163 (22) "Raw agricultural commodity" means any food in its raw or  
164 natural state, including all fruits that are washed, colored or otherwise  
165 treated in their unpeeled natural form prior to marketing;

166 (23) The term "safe" has reference to the health of man or animal;

167 (24) "Sale" means any and every sale and includes (A) manufacture,  
168 processing, packing, canning, bottling or any other production,

169 preparation or putting up; (B) exposure, offer or any other proffer; (C)  
170 holding, storing or any other possessing; (D) dispensing, giving,  
171 delivering, serving or any other supplying; and (E) applying,  
172 administering or any other using.

173 Sec. 2. (NEW) (*Effective October 1, 2013*) For purposes of this section  
174 and section 3 of this act:

175 (1) "Cultivated commercially" means grown or raised by a person in  
176 the course of his or her business or trade and sold within the state;

177 (2) "Enzyme" means a protein that catalyzes chemical reactions of  
178 other substances without being destroyed or altered upon completion  
179 of such reactions;

180 (3) "Genetically engineered" or "genetic engineering" means a  
181 process whereby any food intended for human consumption (A) is  
182 produced from an organism or organisms in which the genetics are  
183 materially altered through the application of: (i) In vitro nucleic acid  
184 techniques, including recombinant DNA (deoxyribonucleic acid)  
185 techniques, the direct injection of nucleic acid into cells or organelles,  
186 encapsulation, gene deletion and doubling, or (ii) methods of fusing  
187 cells that do not fall within the same taxonomic family, that overcome  
188 natural physiological reproductive or recombinant barriers and that  
189 are not techniques used in traditional breeding and selection such as  
190 conjugation, transduction and hybridization; (B) is treated with a  
191 material described in subparagraph (A) of this subdivision, except  
192 manure that is used as a fertilizer for a raw agricultural commodity; or  
193 (C) contains an ingredient, component or substance described in  
194 subparagraph (A) of this subdivision.

195 (4) "Organism" means any biological entity capable of replication,  
196 reproduction or transferring genetic material;

197 (5) "Processed food" means any food other than a raw agricultural  
198 commodity and includes any food produced from a raw agricultural

199 commodity that has been processed through canning, smoking,  
200 pressing, cooking, freezing, dehydration, fermentation or milling;

201 (6) "Processing aid" means: (A) Any substance that is added to a  
202 food during the processing of such food but that is removed in some  
203 manner from the food before the food is packaged in a finished form;  
204 (B) any substance that is added to a food during processing, that is  
205 converted into constituents normally present in the food, and that does  
206 not significantly increase the amount of the constituents naturally  
207 found in the food; or (C) any substance that is added to a food for its  
208 technical or functional effect in the processing but that is present in the  
209 finished food at insignificant levels and that does not have any  
210 technical or functional effect in the finished food;

211 (7) "Raw agricultural commodity" means any food in its raw or  
212 natural state, including a fruit that is washed, colored or treated in its  
213 unpeeled, natural form;

214 (8) "Retailer" means a person or entity that engages in the sale of  
215 food to a consumer;

216 (9) "Distributor" means a person or entity that sells, supplies,  
217 furnishes or transports food in this state that such person or entity  
218 does not produce; and

219 (10) "Manufacturer" means a person who produces seed, seed stock  
220 or food and sells such item to a retailer or distributor.

221 Sec. 3. (NEW) (*Effective October 1, 2013*) (a) On and after October 1,  
222 2014, any food, seed or seed stock offered or intended for retail sale in  
223 this state that is, or may have been, entirely or partially genetically  
224 engineered, except a processed food in which one or more processing  
225 aids or enzymes were produced or derived from genetic engineering,  
226 shall be labeled as follows: (1) In the case of food for retail sale  
227 contained in a package, by the manufacturer, distributor or retailer of  
228 the food, with the clear and conspicuous words: "Produced with

229 Genetic Engineering"; (2) in the case of food that is a raw agricultural  
230 commodity, on the package offered for retail sale or, in the case of any  
231 such commodity that is not separately packaged or labeled, on the  
232 retail store shelf or bin that holds such commodity displayed for sale,  
233 by the retailer, with the clear and conspicuous words: "Produced with  
234 Genetic Engineering"; and (3) in the case of any seed or seed stock, on  
235 the container holding the seed or seed stock displayed for sale, the  
236 sales receipt, or any label identifying ownership or possession of the  
237 commodity, by the manufacturer or distributor, with the clear and  
238 conspicuous words: "Produced with Genetic Engineering".

239 (b) Notwithstanding the provisions of chapter 418 of the general  
240 statutes, the Commissioner of Consumer Protection, in consultation  
241 with the Commissioners of Agriculture, Energy and Environmental  
242 Protection and Public Health, may adopt regulations, pursuant to  
243 chapter 54 of the general statutes, to implement and enforce the  
244 provisions of this section.

245 Sec. 4. Section 21a-102 of the general statutes is repealed and the  
246 following is substituted in lieu thereof (*Effective October 1, 2013*):

247 A food shall be deemed to be misbranded: [(a)] (1) If its labeling is  
248 false or misleading in any particular. A statement on the label or  
249 labeling either directly or indirectly implying that the product is  
250 recommended or endorsed by any agency of the federal or state  
251 government shall be considered misleading, unless the agency  
252 concerned has approved the statement prior to its use; [(b)] (2) if it is  
253 offered for sale under the name of another food; [(c)] (3) if it is an  
254 imitation of another food, unless its label bears, in type of uniform size  
255 and prominence, the word "imitation" and, immediately thereafter, the  
256 name of the food imitated; [(d)] (4) if its container is so made, formed  
257 or filled as to be misleading; [(e)] (5) if in package form, unless it bears  
258 a label containing [(1)] (A) the name and place of business of the  
259 manufacturer, packer or distributor; and [(2)] (B) an accurate statement  
260 of the quantity of the contents in terms of weight, measure or

261 numerical count; provided, under [subdivision (2) of this subsection]  
262 subparagraph (B) of this subdivision, reasonable variations shall be  
263 permitted, and exemptions as to small packages shall be established by  
264 regulations promulgated by the commissioner and director, acting  
265 jointly; [(f)] (6) if any information or other word or statement, required  
266 by or under authority of this chapter to appear on the label or labeling,  
267 is not prominently placed thereon with such conspicuousness, as  
268 compared with other words, statements, designs or devices, in the  
269 labeling, and in such terms, as to render it likely to be read and  
270 understood by the ordinary individual under customary conditions of  
271 purchase and use; [(g)] (7) if it purports to be or simulates or is  
272 represented as a food for which a definition and standard of identity  
273 has been prescribed by regulations as provided by section 21a-100,  
274 unless [(1)] (A) it conforms to such definition and standard, and [(2)]  
275 (B) its label bears the name of the food specified in the definition and  
276 standard, and, so far as may be required by such regulations, the  
277 common names of optional ingredients, other than spices, flavoring  
278 and coloring, present in such food; [(h)] (8) if it purports to be or is  
279 represented as [(1)] (A) a food for which a standard of quality has been  
280 prescribed by regulations as provided by section 21a-100 and its  
281 quality falls below such standard, unless its label bears, in such  
282 manner and form as such regulations specify, a statement that it falls  
283 below such standard; [or (2)] (B) a food for which a standard or  
284 standards of fill of container have been prescribed by regulations as  
285 provided by section 21a-100, and it falls below the standard of fill of  
286 container applicable thereto, unless its label bears, in such manner and  
287 form as such regulations specify, a statement that it falls below such  
288 standard; [(3)] or (C) a food for which no definition and standard of  
289 identity and no standard of quality has been prescribed by regulations  
290 as provided by section 21a-100, and it falls below the standard of  
291 purity, quality or strength which it purports or is represented to  
292 possess; [(i)] (9) if it is not subject to the provisions of [subsection (g)]  
293 subdivision (7) of this section, unless its label bears [(1)] (A) the  
294 common or usual name of the food, if any, and [(2)] (B) if it is

295 fabricated from two or more ingredients, the common or usual name  
296 of each such ingredient; except that spices, flavorings and colorings,  
297 other than those sold as such, may be designated as spices, flavorings  
298 and colorings without naming each; provided, to the extent that  
299 compliance with the requirements of [subdivision (2) of this  
300 subsection] subparagraph (B) of this subdivision is impracticable, or  
301 results in deception or unfair competition, exemptions shall be  
302 established by regulations promulgated by the commissioner and  
303 director, acting jointly; [(j)] (10) if it purports to be or is represented to  
304 be for special dietary uses, unless its label bears such information  
305 concerning its vitamin, mineral and other dietary properties as is  
306 necessary in order fully to inform purchasers as to its value for such  
307 uses, as provided by regulations promulgated by the commissioner  
308 and director, acting jointly; [(k)] (11) if it bears or contains any artificial  
309 flavoring, artificial coloring, artificial sweetening or chemical  
310 preservative, unless it bears labeling stating that fact; provided, to the  
311 extent that compliance with the requirements of this subsection is  
312 impracticable, exemptions shall be established by regulations  
313 promulgated by the commissioner and director, acting jointly; (12) if it  
314 is genetically engineered, as defined in section 2, of this act, and does  
315 not bear labeling as required in accordance with section 3 of this act,  
316 unless (A) it is a food produced without the producer's knowledge that  
317 a seed or other component of the food was genetically engineered, or  
318 (B) on or before July 1, 2019, it is a processed food, as defined in section  
319 2 of this act, that is subject to the provisions of section 3 of this act,  
320 solely because it contains one or more materials that are genetically  
321 engineered, as defined in section 2 of this act, provided such  
322 genetically engineered materials do not, in the aggregate, account for  
323 more than nine-tenths of one per cent of the total weight of the  
324 processed food.

This act shall take effect as follows and shall amend the following sections:		
Section 1	October 1, 2013	21a-92

Sec. 2	<i>October 1, 2013</i>	New section
Sec. 3	<i>October 1, 2013</i>	New section
Sec. 4	<i>October 1, 2013</i>	21a-102

**Statement of Purpose:**

To require the labeling of genetically engineered food.

*[Proposed deletions are enclosed in brackets. Proposed additions are indicated by underline, except that when the entire text of a bill or resolution or a section of a bill or resolution is new, it is not underlined.]*