

103RD GENERAL ASSEMBLY State of Illinois 2023 and 2024 HB2812

Introduced 2/16/2023, by Rep. Bob Morgan - Kelly M. Cassidy

SYNOPSIS AS INTRODUCED:

35 ILCS 105/3-10 410 ILCS 130/105 410 ILCS 705/55-21

Amends the Compassionate Use of Medical Cannabis Program Act. Provides that a medical cannabis container shall be compliant with standards established by the Consumer Product Safety Commission, unless the medical cannabis container carries a warning that it is not recommended for use in households with children. Amends the Use Tax Act and the Cannabis Regulation and Tax Act to make corresponding changes.

LRB103 29816 CPF 56224 b

1 AN ACT concerning safety.

Be it enacted by the People of the State of Illinois, represented in the General Assembly:

- Section 5. The Use Tax Act is amended by changing Section 3-10 as follows:
- 6 (35 ILCS 105/3-10)

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Sec. 3-10. Rate of tax. Unless otherwise provided in this Section, the tax imposed by this Act is at the rate of 6.25% of either the selling price or the fair market value, if any, of the tangible personal property. In all cases where property functionally used or consumed is the same as the property that was purchased at retail, then the tax is imposed on the selling price of the property. In all cases where property functionally used or consumed is a by-product or waste product that has been refined, manufactured, or produced from property purchased at retail, then the tax is imposed on the lower of the fair market value, if any, of the specific property so used in this State or on the selling price of the property purchased at retail. For purposes of this Section "fair market value" means the price at which property would change hands between a willing buyer and a willing seller, neither being under any compulsion to buy or sell and both having reasonable knowledge of the relevant facts. The fair market value shall be

established by Illinois sales by the taxpayer of the same property as that functionally used or consumed, or if there are no such sales by the taxpayer, then comparable sales or purchases of property of like kind and character in Illinois.

Beginning on July 1, 2000 and through December 31, 2000, with respect to motor fuel, as defined in Section 1.1 of the Motor Fuel Tax Law, and gasohol, as defined in Section 3-40 of the Use Tax Act, the tax is imposed at the rate of 1.25%.

Beginning on August 6, 2010 through August 15, 2010, and beginning again on August 5, 2022 through August 14, 2022, with respect to sales tax holiday items as defined in Section 3-6 of this Act, the tax is imposed at the rate of 1.25%.

With respect to gasohol, the tax imposed by this Act applies to (i) 70% of the proceeds of sales made on or after January 1, 1990, and before July 1, 2003, (ii) 80% of the proceeds of sales made on or after July 1, 2003 and on or before July 1, 2017, and (iii) 100% of the proceeds of sales made thereafter. If, at any time, however, the tax under this Act on sales of gasohol is imposed at the rate of 1.25%, then the tax imposed by this Act applies to 100% of the proceeds of sales of gasohol made during that time.

With respect to majority blended ethanol fuel, the tax imposed by this Act does not apply to the proceeds of sales made on or after July 1, 2003 and on or before December 31, 2023 but applies to 100% of the proceeds of sales made thereafter.

With respect to biodiesel blends with no less than 1% and no more than 10% biodiesel, the tax imposed by this Act applies to (i) 80% of the proceeds of sales made on or after July 1, 2003 and on or before December 31, 2018 and (ii) 100% of the proceeds of sales made after December 31, 2018 and before January 1, 2024. On and after January 1, 2024 and on or before December 31, 2030, the taxation of biodiesel, renewable diesel, and biodiesel blends shall be as provided in Section 3-5.1. If, at any time, however, the tax under this Act on sales of biodiesel blends with no less than 1% and no more than 10% biodiesel is imposed at the rate of 1.25%, then the tax imposed by this Act applies to 100% of the proceeds of sales of biodiesel blends with no less than 1% and no more than 10% biodiesel made during that time.

With respect to biodiesel and biodiesel blends with more than 10% but no more than 99% biodiesel, the tax imposed by this Act does not apply to the proceeds of sales made on or after July 1, 2003 and on or before December 31, 2023. On and after January 1, 2024 and on or before December 31, 2030, the taxation of biodiesel, renewable diesel, and biodiesel blends shall be as provided in Section 3-5.1.

Until July 1, 2022 and beginning again on July 1, 2023, with respect to food for human consumption that is to be consumed off the premises where it is sold (other than alcoholic beverages, food consisting of or infused with adult use cannabis, soft drinks, and food that has been prepared for

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Beginning on July 1, 2022 and until July 1, 2023, with respect to food for human consumption that is to be consumed off the premises where it is sold (other than alcoholic beverages,

immediate consumption), the tax is imposed at the rate of 1%.

food consisting of or infused with adult use cannabis, soft

drinks, and food that has been prepared for immediate

consumption), the tax is imposed at the rate of 0%.

respect to prescription With and nonprescription medicines, drugs, medical appliances, products classified as Class III medical devices by the United States Food and Drug Administration that are used for cancer treatment pursuant to a prescription, as well as any accessories and components related to those devices, modifications to a motor vehicle for the purpose of rendering it usable by a person with a disability, and insulin, blood sugar testing materials, syringes, and needles used by human diabetics, the tax is imposed at the rate of 1%. For the purposes of this Section, until September 1, 2009: the term "soft drinks" means any complete, finished, ready-to-use, non-alcoholic drink, whether carbonated or not, including, but not limited to, soda water, cola, fruit juice, vegetable juice, carbonated water, and all other preparations commonly known as soft drinks of whatever kind or description that are contained in any closed or sealed bottle, can, carton, or container, regardless of size; but "soft drinks" does not include coffee, tea, non-carbonated water, infant formula, milk or milk products as defined in the

Grade A Pasteurized Milk and Milk Products Act, or drinks containing 50% or more natural fruit or vegetable juice.

Notwithstanding any other provisions of this Act, beginning September 1, 2009, "soft drinks" means non-alcoholic beverages that contain natural or artificial sweeteners. "Soft drinks" does do not include beverages that contain milk or milk products, soy, rice or similar milk substitutes, or greater than 50% of vegetable or fruit juice by volume.

Until August 1, 2009, and notwithstanding any other provisions of this Act, "food for human consumption that is to be consumed off the premises where it is sold" includes all food sold through a vending machine, except soft drinks and food products that are dispensed hot from a vending machine, regardless of the location of the vending machine. Beginning August 1, 2009, and notwithstanding any other provisions of this Act, "food for human consumption that is to be consumed off the premises where it is sold" includes all food sold through a vending machine, except soft drinks, candy, and food products that are dispensed hot from a vending machine, regardless of the location of the vending machine.

Notwithstanding any other provisions of this Act, beginning September 1, 2009, "food for human consumption that is to be consumed off the premises where it is sold" does not include candy. For purposes of this Section, "candy" means a preparation of sugar, honey, or other natural or artificial sweeteners in combination with chocolate, fruits, nuts or

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other ingredients or flavorings in the form of bars, drops, or pieces. "Candy" does not include any preparation that contains flour or requires refrigeration.

Notwithstanding any other provisions of this Act, beginning September 1, 2009, "nonprescription medicines and drugs" does not include grooming and hygiene products. For purposes of this Section, "grooming and hygiene products" includes, but is not limited to, soaps and cleaning solutions, shampoo, toothpaste, mouthwash, antiperspirants, and sun tan lotions and screens, unless those products are available by prescription only, regardless of whether the products meet the definition of "over-the-counter-drugs". For the purposes of this paragraph, "over-the-counter-drug" means a drug for human use that contains a label that identifies the product as a drug C.F.R. S required by 21 CFR 201.66. "over-the-counter-drug" label includes:

- (A) a A "Drug Facts" panel; or
- 18 (B) <u>a</u> A statement of the "active ingredient(s)" with a

 19 list of those ingredients contained in the compound,

 20 substance or preparation.

"Prescription Beginning on the effective date of this amendatory Act of the 98th General Assembly, "prescription and nonprescription medicines and drugs" includes medical cannabis purchased by a qualifying patient or a designated caregiver from a registered dispensing organization under the Compassionate Use of Medical Cannabis Program Act or a

- 1 <u>secondary site dispensary or dispensary under the Cannabis</u>
- 2 Regulation and Tax Act.
- 3 As used in this Section, "adult use cannabis" means
- 4 cannabis subject to tax under the Cannabis Cultivation
- 5 Privilege Tax Law and the Cannabis Purchaser Excise Tax Law
- 6 and does not include cannabis subject to tax under the
- 7 Compassionate Use of Medical Cannabis Program Act.
- 8 If the property that is purchased at retail from a
- 9 retailer is acquired outside Illinois and used outside
- 10 Illinois before being brought to Illinois for use here and is
- 11 taxable under this Act, the "selling price" on which the tax is
- 12 computed shall be reduced by an amount that represents a
- 13 reasonable allowance for depreciation for the period of prior
- 14 out-of-state use.
- 15 (Source: P.A. 101-363, eff. 8-9-19; 101-593, eff. 12-4-19;
- 16 102-4, eff. 4-27-21; 102-700, Article 20, Section 20-5, eff.
- 17 4-19-22; 102-700, Article 60, Section 60-15, eff. 4-19-22;
- 18 102-700, Article 65, Section 65-5, eff. 4-19-22; revised
- 19 5-27-22.)
- 20 Section 10. The Compassionate Use of Medical Cannabis
- 21 Program Act is amended by changing Section 105 as follows:
- 22 (410 ILCS 130/105)
- Sec. 105. Requirements; prohibitions; penalties for
- 24 cultivation centers.

- (a) The operating documents of a registered cultivation center shall include procedures for the oversight of the cultivation center, a cannabis plant monitoring system including a physical inventory recorded weekly, a cannabis container system including a physical inventory recorded weekly, accurate record keeping, and a staffing plan.
- (b) A registered cultivation center shall implement a security plan reviewed by the Illinois State Police and including but not limited to: facility access controls, perimeter intrusion detection systems, personnel identification systems, 24-hour surveillance system to monitor the interior and exterior of the registered cultivation center facility and accessible to authorized law enforcement and the Department of Agriculture in real-time.
- (c) A registered cultivation center may not be located within 2,500 feet of the property line of a pre-existing public or private preschool or elementary or secondary school or day care center, day care home, group day care home, part day child care facility, or an area zoned for residential use.
- (d) All cultivation of cannabis for distribution to a registered dispensing organization must take place in an enclosed, locked facility as it applies to cultivation centers at the physical address provided to the Department of Agriculture during the registration process. The cultivation center location shall only be accessed by the cultivation center agents working for the registered cultivation center,

- 1 Department of Agriculture staff performing inspections,
- 2 Department of Public Health staff performing inspections, law
- 3 enforcement or other emergency personnel, and contractors
- 4 working on jobs unrelated to medical cannabis, such as
- 5 installing or maintaining security devices or performing
- 6 electrical wiring.
- 7 (e) A cultivation center may not sell or distribute any
- 8 cannabis to any individual or entity other than another
- 9 cultivation center, a dispensing organization registered under
- 10 this Act, or a laboratory licensed by the Department of
- 11 Agriculture.
- 12 (f) All harvested cannabis intended for distribution to a
- dispensing organization must be packaged in a labeled medical
- 14 cannabis container and entered into a data collection system.
- 15 A medical cannabis container shall be compliant with standards
- 16 established by the Consumer Product Safety Commission, unless
- 17 the medical cannabis container carries a warning that it is
- 18 not recommended for use in households with children.
- 19 (g) No person who has been convicted of an excluded
- offense may be a cultivation center agent.
- 21 (h) Registered cultivation centers are subject to random
- inspection by the Illinois State Police.
- 23 (i) Registered cultivation centers are subject to random
- 24 inspections by the Department of Agriculture and the
- 25 Department of Public Health.
- 26 (j) A cultivation center agent shall notify local law

- 1 enforcement, the Illinois State Police, and the Department of
- 2 Agriculture within 24 hours of the discovery of any loss or
- 3 theft. Notification shall be made by phone or in-person, or by
- 4 written or electronic communication.
- 5 (k) A cultivation center shall comply with all State and
- 6 federal rules and regulations regarding the use of pesticides.
- 7 (Source: P.A. 101-363, eff. 8-9-19; 102-538, eff. 8-20-21.)
- 8 Section 15. The Cannabis Regulation and Tax Act is amended
- 9 by changing Section 55-21 as follows:
- 10 (410 ILCS 705/55-21)
- 11 Sec. 55-21. Cannabis product packaging and labeling.
- 12 (a) Each cannabis product produced for sale shall be
- 13 registered with the Department of Agriculture on forms
- 14 provided by the Department of Agriculture. Each product
- 15 registration shall include a label and the required
- 16 registration fee at the rate established by the Department of
- 17 Agriculture for a comparable medical cannabis product, or as
- 18 established by rule. The registration fee is for the name of
- 19 the product offered for sale and one fee shall be sufficient
- 20 for all package sizes.
- 21 (b) All harvested cannabis intended for distribution to a
- 22 cannabis enterprise must be packaged in a sealed, labeled
- 23 container.
- 24 (c) Any product containing cannabis shall be sold in a

- sealed, odor-proof, and child-resistant cannabis container consistent with current standards, including the Consumer Product Safety Commission standards referenced by the Poison Prevention Act unless the sale is between or among a craft grower, infuser, or cultivation center or the medical cannabis container carries a warning that it is not recommended for use in households with children.
 - (d) All cannabis-infused products shall be individually wrapped or packaged at the original point of preparation. The packaging of the cannabis-infused product shall conform to the labeling requirements of the Illinois Food, Drug and Cosmetic Act, in addition to the other requirements set forth in this Section.
 - (e) Each cannabis product shall be labeled before sale and each label shall be securely affixed to the package and shall state in legible English and any languages required by the Department of Agriculture:
 - (1) the name and post office box of the registered cultivation center or craft grower where the item was manufactured;
 - (2) the common or usual name of the item and the registered name of the cannabis product that was registered with the Department of Agriculture under subsection (a);
 - (3) a unique serial number that will match the product with a cultivation center or craft grower batch and lot

1	number to facilitate any warnings or recalls the
2	Department of Agriculture, cultivation center, or craft
3	grower deems appropriate;
4	(4) the date of final testing and packaging, if
5	sampled, and the identification of the independent testing
6	laboratory;
7	(5) the date of harvest and "use by" date;
8	(6) the quantity (in ounces or grams) of cannabis
9	contained in the product;
10	(7) a pass/fail rating based on the laboratory's
11	microbiological, mycotoxins, and pesticide and solvent
12	residue analyses, if sampled;
13	(8) content list.
14	(A) A list of the following, including the minimum
15	and maximum percentage content by weight for
16	subdivisions (e)(8)(A)(i) through (iv):
17	(i) delta-9-tetrahydrocannabinol (THC);
18	(ii) tetrahydrocannabinolic acid (THCA);
19	(iii) cannabidiol (CBD);
20	(iv) cannabidiolic acid (CBDA); and
21	(v) all other ingredients of the item,
22	including any colors, artificial flavors, and
23	preservatives, listed in descending order by
24	predominance of weight shown with common or usual
25	names.
26	(B) The acceptable tolerances for the minimum

_	percentage	printed	on	the	label	for	any	of
2	subdivisions	(e)(8)(A)) (i)	throu	gh (iv)	shall	not	be
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- (f) Packaging must not contain information that:
 - (1) is false or misleading;
 - (2) promotes excessive consumption;
- (3) depicts a person under 21 years of age consuming cannabis;
 - (4) includes the image of a cannabis leaf;
 - (5) includes any image designed or likely to appeal to minors, including cartoons, toys, animals, or children, or any other likeness to images, characters, or phrases that are popularly used to advertise to children, or any packaging or labeling that bears reasonable resemblance to any product available for consumption as a commercially available candy, or that promotes consumption of cannabis;
 - (6) contains any seal, flag, crest, coat of arms, or other insignia likely to mislead the purchaser to believe that the product has been endorsed, made, or used by the State of Illinois or any of its representatives except where authorized by this Act.
- (g) Cannabis products produced by concentrating or extracting ingredients from the cannabis plant shall contain the following information, where applicable:
 - (1) If solvents were used to create the concentrate or extract, a statement that discloses the type of extraction

- 1 method, including any solvents or gases used to create the 2 concentrate or extract; and
 - (2) Any other chemicals or compounds used to produce or were added to the concentrate or extract.
 - (h) All cannabis products must contain warning statements established for purchasers, of a size that is legible and readily visible to a consumer inspecting a package, which may not be covered or obscured in any way. The Department of Public Health shall define and update appropriate health warnings for packages including specific labeling or warning requirements for specific cannabis products.
 - (i) Unless modified by rule to strengthen or respond to new evidence and science, the following warnings shall apply to all cannabis products unless modified by rule: "This product contains cannabis and is intended for use by adults 21 and over. Its use can impair cognition and may be habit forming. This product should not be used by pregnant or breastfeeding women. It is unlawful to sell or provide this item to any individual, and it may not be transported outside the State of Illinois. It is illegal to operate a motor vehicle while under the influence of cannabis. Possession or use of this product may carry significant legal penalties in some jurisdictions and under federal law.".
 - (j) Warnings for each of the following product types must be present on labels when offered for sale to a purchaser:
 - (1) Cannabis that may be smoked must contain a

- 1 statement that "Smoking is hazardous to your health.".
 - (2) Cannabis-infused products (other than those intended for topical application) must contain a statement "CAUTION: This product contains cannabis, and intoxication following use may be delayed 2 or more hours. This product was produced in a facility that cultivates cannabis, and that may also process common food allergens.".
 - (3) Cannabis-infused products intended for topical application must contain a statement "DO NOT EAT" in bold, capital letters.
 - (k) Each cannabis-infused product intended for consumption must be individually packaged, must include the total milligram content of THC and CBD, and may not include more than a total of 100 milligrams of THC per package. A package may contain multiple servings of 10 milligrams of THC, indicated by scoring, wrapping, or by other indicators designating individual serving sizes. The Department of Agriculture may change the total amount of THC allowed for each package, or the total amount of THC allowed for each serving size, by rule.
 - (1) No individual other than the purchaser may alter or destroy any labeling affixed to the primary packaging of cannabis or cannabis-infused products.
 - (m) For each commercial weighing and measuring device used at a facility, the cultivation center or craft grower must:
 - (1) Ensure that the commercial device is licensed under the Weights and Measures Act and the associated

- 1 administrative rules (8 Ill. Adm. Code 600);
- 2 (2) Maintain documentation of the licensure of the commercial device; and
- 4 (3) Provide a copy of the license of the commercial device to the Department of Agriculture for review upon request.
- 7 (n) It is the responsibility of the Department to ensure 8 that packaging and labeling requirements, including product 9 warnings, are enforced at all times for products provided to 10 purchasers. Product registration requirements and container 11 requirements may be modified by rule by the Department of 12 Agriculture.
- 13 (o) Labeling, including warning labels, may be modified by 14 rule by the Department of Agriculture.
- 15 (Source: P.A. 101-27, eff. 6-25-19; 101-593, eff. 12-4-19;
- 16 102-98, eff. 7-15-21.)