

1 ENGROSSED HOUSE AMENDMENT

2 TO

2 ENGROSSED SENATE BILL NO. 891

By: Murdock and Prieto of the
Senate

3 and

4 Pae of the House

5

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7 An Act relating to kratom products; amending 63 O.S.
2021, Section 1-1432.2, as amended by Section 1,
8 Chapter 278, O.S.L. 2024 (63 O.S. Supp. 2024, Section
9 1-1432.2), which relates to definitions used in the
Oklahoma Kratom Consumer Protection Act; modifying
definitions; updating statutory reference; and
providing an effective date.

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13 AMENDMENT NO. 1. Strike the title, enacting clause, and entire bill
and insert:

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16 "An Act relating to kratom products; amending 63 O.S.
2021, Sections 1-1432.2 and 1-1432.4, as amended by
17 Sections 1 and 2, Chapter 278, O.S.L. 2024 (63 O.S.
Supp. 2024, Sections 1-1432.2 and 1-1432.4), which
18 relate to the Oklahoma Kratom Consumer Protection
Act; modifying and adding definitions; removing
19 certain packaging and labeling requirements;
requiring inclusion of certain statement on labels;
20 directing vendors to provide test results from
independent testing laboratories upon request; and
21 providing an effective date.

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24 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

1 SECTION 1. AMENDATORY 63 O.S. 2021, Section 1-1432.2, as
2 amended by Section 1, Chapter 278, O.S.L. 2024 (63 O.S. Supp. 2024,
3 Section 1-1432.2), is amended to read as follows:

4 Section 1-1432.2. As used in ~~this act~~ the Oklahoma Kratom
5 Consumer Protection Act:

6 1. "Food" means a food, food product, food ingredient, dietary
7 ingredient, dietary supplement or beverage for human consumption;

8 2. "Independent testing laboratory" means a laboratory that:

9 a. does not have a direct or indirect interest in the
10 entity whose product is being tested,

11 b. does not have a direct or indirect interest in a
12 facility that processes, distributes, dispenses, or
13 sells kratom products in this state or in another
14 jurisdiction, and

15 c. is nationally accredited by an accrediting body as
16 defined by Section 150.37 of Title 74 of the Oklahoma
17 Statutes;

18 2. 3. "Kratom leaf" means the leaf of the kratom plant,

19 Mitragyna speciosa, in fresh or dehydrated or dried form that
20 undergoes no post-harvest processing other than drying or size
21 reduction by cutting, milling, or similar procedure, and may be
22 cleaned or sterilized using standard treatments applied to food
23 ingredients, such as heat, steam, pressurization, or irradiation or
24 other standard treatments applied to food ingredients. The total

1 alkaloid content of kratom leaf material used in the kratom product
2 shall not exceed three and one-half percent (3.5%) measured on a
3 dried weight-to-weight basis;

4 3. 4. "Kratom leaf extract" means the material obtained by
5 extracting kratom using a solvent consisting of:

- 6 a. water, ethanol, or food-grade carbon dioxide (CO₂), or
 - 7 b. any other solvent allowed by federal or state
- 8 regulation for use in manufacturing a food ingredient.

9 The extracted material shall contain mitragynine as the most
10 abundant alkaloid, measured on a weight-to-weight basis, ~~and at a~~
11 ~~level that is equal to or exceeds twice that of any other alkaloid~~
12 ~~present. The ratio of mitragynine to other alkaloids in the extract~~
13 ~~shall be equal to or greater than the ratio found in the starting~~
14 ~~material;~~

15 4. 5. "Kratom product" means a food or dietary supplement that
16 consists of or contains kratom leaf or kratom leaf extract that does
17 not contain any synthesized kratom alkaloids, other synthesized
18 kratom constituents, or synthesized metabolites of any kratom
19 constituent in which the level of 7-hydroxymitragynine, on a percent
20 weight basis, is not greater than one percent (1%) of the amount of
21 total kratom alkaloids, as confirmed with a high-performance liquid
22 chromatography testing method. For purposes of this paragraph,
23 "synthesized" refers to substances produced using directed synthetic

1 or biosynthetic chemistry, as opposed to traditional food
2 preparation techniques such as heating or extracting;

3 5. 6. "Total kratom alkaloids" means the sum of mitragynine,
4 speciociliatine, speciogynine, paynantheine, and 7-
5 hydroxymitragynine; and

6 6. 7. "Vendor" means a person or entity that sells, prepares or
7 maintains kratom products or that advertises, represents, or holds
8 himself, herself, or itself out as selling, preparing or maintaining
9 kratom products and includes a manufacturer, wholesaler, store,
10 restaurant, hotel, catering facility, camp, bakery, delicatessen,
11 supermarket, grocery store, convenience store, nursing home, or food
12 or drink company.

13 SECTION 2. AMENDATORY 63 O.S. 2021, Section 1-1432.4, as
14 amended by Section 2, Chapter 278, O.S.L. 2024 (63 O.S. Supp. 2024,
15 Section 1-1432.4), is amended to read as follows:

16 Section 1-1432.4. A. A vendor shall not prepare, distribute,
17 sell, or expose for sale any of the following:

18 1. A kratom product that does not meet the definition for a
19 kratom product pursuant to Section 1-1432.2 of this title;

20 2. A kratom product that is contaminated with a dangerous
21 nonkratom substance. A kratom product is contaminated with a
22 dangerous nonkratom substance if the kratom product contains a
23 substance that is not safe for human consumption;

1 3. A kratom product containing a level of 7-hydroxymitragynine
2 in the alkaloid fraction that is greater than one percent (1%) of
3 the alkaloid composition of the product;

4 4. A kratom product containing any synthesized alkaloid
5 including synthesized mitragynine, synthesized 7-hydroxymitragynine
6 or any other synthesized compounds of the kratom plant;

7 5. A kratom product containing any controlled substance listed
8 in the Uniform Controlled Dangerous Substances Act, unless the
9 product is compounded by a licensed pharmacist with the controlled
10 substance dispensed in accordance with a valid prescription; or

11 6. A kratom product containing a level of any residual solvent
12 that was used in the manufacturing of the extract that exceeds the
13 residual level specified for pharmaceutical products in the document
14 "Q3C - Tables and List, Guidance for Industry, [June 2017] ICH
15 Revision 3" issued by the United States Department of Health and
16 Human Services, Food and Drug Administration.

17 B. Kratom products shall be accompanied by a label bearing the
18 following information prior to its sale in this state:

19 1. A list of the ingredients, which shall include the common or
20 usual name of each ingredient used in the manufacture of the
21 product, listed in descending order of predominance;

22 2. That the sale or transfer of kratom to a person under
23 eighteen (18) years of age is prohibited;

1 3. The amount of total kratom alkaloids, mitragynine, and 7-
2 hydroxymitragynine contained in the product;

3 4. The amount of total kratom alkaloids, mitragynine, and 7-
4 hydroxymitragynine contained in packaging for the product;

5 5. The name and the principal street address of the vendor or
6 the person responsible for distributing the product;

7 6. Any federal food allergen labeling requirements, if
8 applicable, and clear and adequate directions for the consumption
9 and safe and effective use of such product, including the
10 recommended serving size, the number of servings in the container,
11 and the number of servings that can be safely consumed in a day.

12 Provided, liquid kratom products shall be packaged in a retail
13 container that has clear serving size markings and be subject to the
14 following requirements:

15 a. products of less than eight (8) fluid ounces which
16 contain more than three servings shall be accompanied
17 by a calibrated measuring device, and

18 b. if such a product contains more than the eight (8)
19 fluid ounces, the requirements specified in
20 subparagraph a of this paragraph do not apply-

21 ~~Provided further, packaging for powdered kratom products not in~~
22 ~~capsule form shall have a calibrated measuring device included in~~
23 ~~the container;~~

1 7. Any precautionary statements as to the safety and
2 effectiveness of the product, including a warning that a consumer
3 should consult a health care professional on questions about the use
4 of kratom, ~~and~~ that the product may be habit-forming, ~~and a~~
5 ~~statement that the kratom product is not intended to "diagnose,~~
6 ~~treat, cure, or prevent any disease"; and~~

7 8. A statement that ~~a kratom product label is prohibited from~~
8 ~~making any therapeutic claims unless approved by the United States~~
9 ~~Food and Drug Administration.~~ states, "These statements have not
10 been evaluated by the United States Food and Drug Administration.
11 This product is not intended to diagnose, treat, cure, or prevent
12 any disease."

13 C. A vendor may not distribute, sell, or expose for sale a
14 kratom product to an individual under eighteen (18) years of age.

15 D. Upon request by the State Department of Health, the vendor
16 shall provide test results from a United States-based testing
17 facility, that is an independent testing laboratory as defined in
18 Section 1-1432.2 of this title, to confirm the items listed on the
19 product label.

20 SECTION 3. This act shall become effective November 1, 2025."

Passed the House of Representatives the 5th day of May, 2025.

Presiding Officer of the House of
Representatives

Passed the Senate the _____ day of _____, 2025.

Presiding Officer of the Senate

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9 1-1432.2), which relates to definitions used in the
Oklahoma Kratom Consumer Protection Act; modifying
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12 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

13 SECTION 2. AMENDATORY 63 O.S. 2021, Section 1-1432.2, as
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15 Section 1-1432.2), is amended to read as follows:

16 Section 1-1432.2. As used in ~~this act~~ the Oklahoma Kratom
17 Consumer Protection Act:

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19 ingredient, dietary supplement or beverage for human consumption;

20 2. "Kratom leaf" means the leaf of the kratom plant, *Mitragyna*
speciosa, in fresh or dehydrated or dried form that undergoes no
post-harvest processing other than drying or size reduction by
cutting, milling, or similar procedure, and may be cleaned or

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2 such as heat, steam, pressurization, or irradiation or other
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5 exceed three and one-half percent (3.5%) measured on a dried weight-
6 to-weight basis;

7 3. "Kratom leaf extract" means the material obtained by
8 extracting kratom using a solvent consisting of:

- a. water, ethanol, or food-grade carbon dioxide (CO₂), or
 - b. any other solvent allowed by federal or state regulation for use in manufacturing a food ingredient.

12 The extracted material shall contain mitragynine as the most
13 abundant alkaloid, measured on a weight-to-weight basis, and at a
14 level that is equal to or exceeds twice that of any other alkaloid
15 present. The ratio of mitragynine to other alkaloids in the extract
16 shall be equal to or greater than the ratio found in the starting
17 material;

18 4. "Kratom product" means a food or dietary supplement that
19 consists of or contains kratom leaf or kratom leaf extract that does
20 not contain any synthesized kratom alkaloids, other synthesized
21 kratom constituents, or synthesized metabolites of any kratom
22 constituent in which the level of 7-hydroxymitragynine, on a percent
23 weight basis, is not greater than one percent (1%) of the amount of
24 total kratom alkaloids, as confirmed with a high-performance liquid

1 chromatography testing method. For purposes of this paragraph,
2 "synthesized" refers to substances produced using directed synthetic
3 or biosynthetic chemistry, as opposed to traditional food
4 preparation techniques such as heating or extracting;

5 5. "Total kratom alkaloids" means the sum of mitragynine,
6 speciociliatine, speciogynine, paynantheine, and 7-
7 hydroxymitragynine; and

8 6. "Vendor" means a person or entity that sells, prepares or
9 maintains kratom products or that advertises, represents or holds
10 himself, herself, or itself out as selling, preparing or maintaining
11 kratom products and includes a manufacturer, wholesaler, store,
12 restaurant, hotel, catering facility, camp, bakery, delicatessen,
13 supermarket, grocery store, convenience store, nursing home or food
14 or drink company.

15 SECTION 3. This act shall become effective November 1, 2025.
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Passed the Senate the 17th day of February, 2025.

Presiding Officer of the Senate

Passed the House of Representatives the _____ day of _____,
2025.

Presiding Officer of the House
of Representatives