

FLOOR AMENDMENT
HOUSE OF REPRESENTATIVES
State of Oklahoma

SPEAKER:

CHAIR:

I move to amend SB891 _____
Of the printed Bill
Page _____ Section _____ Lines _____
Of the Engrossed Bill

By deleting the content of the entire measure, and by inserting in lieu thereof the following language:

AMEND TITLE TO CONFORM TO AMENDMENTS

Adopted: _____

Amendment submitted by: Daniel Pae _____

Reading Clerk

1 STATE OF OKLAHOMA

2 1st Session of the 60th Legislature (2025)

3 FLOOR SUBSTITUTE
4 FOR ENGROSSED

5 SENATE BILL NO. 891

By: Murdock and Prieto of the
Senate

6 and

7 Pae of the House

8
9
10 FLOOR SUBSTITUTE

11 An Act relating to kratom products; amending 63 O.S.
12 2021, Sections 1-1432.2 and 1-1432.4, as amended by
13 Section 1, Chapter 278, O.S.L. 2024 (63 O.S. Supp.
14 2024, Sections 1-1432.2 and 1-1432.4), which relates
15 to the Oklahoma Kratom Consumer Protection Act;
16 modifying definitions; adding a definition; removing
17 certain packaging requirements; removing a certain
18 labeling requirement; updating statutory reference;
19 and providing an effective date.

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

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

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

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

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

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

6 b. does not have a direct or indirect interest in a
7 facility that processes, distributes, dispenses, or
8 sells kratom products in this state or in another
9 jurisdiction, and

10 c. is nationally accredited by an accrediting body as
11 defined by Section 150.37 of Title 74 of the Oklahoma
12 Statutes;

13 ~~2.~~ 3. "Kratom leaf" means the leaf of the kratom plant,
14 *Mitragyna speciosa*, in fresh or dehydrated or dried form that
15 undergoes no post-harvest processing other than drying or size
16 reduction by cutting, milling, or similar procedure, and may be
17 cleaned or sterilized using standard treatments applied to food
18 ingredients, such as heat, steam, pressurization, or irradiation or
19 other standard treatments applied to food ingredients. The total
20 alkaloid content of kratom leaf material used in the kratom product
21 shall not exceed three and one-half percent (3.5%) measured on a
22 dried weight-to-weight basis;

23 ~~3.~~ 4. "Kratom leaf extract" means the material obtained by
24 extracting kratom using a solvent consisting of:

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

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

10 ~~4.~~ 5. "Kratom product" means a food or dietary supplement that
11 consists of or contains kratom leaf or kratom leaf extract that does
12 not contain any synthesized kratom alkaloids, other synthesized
13 kratom constituents, or synthesized metabolites of any kratom
14 constituent in which the level of 7-hydroxymitragynine, on a percent
15 weight basis, is not greater than one percent (1%) of the amount of
16 total kratom alkaloids, as confirmed with a high-performance liquid
17 chromatography testing method. For purposes of this paragraph,
18 "synthesized" refers to substances produced using directed synthetic
19 or biosynthetic chemistry, as opposed to traditional food
20 preparation techniques such as heating or extracting;

21 ~~5.~~ 6. "Total kratom alkaloids" means the sum of mitragynine,
22 speciociliatine, speciogynine, paynantheine, and 7-
23 hydroxymitragynine; and
24

1 ~~6.~~ 7. "Vendor" means a person or entity that sells, prepares or
2 maintains kratom products or that advertises, represents, or holds
3 himself, herself, or itself out as selling, preparing or maintaining
4 kratom products and includes a manufacturer, wholesaler, store,
5 restaurant, hotel, catering facility, camp, bakery, delicatessen,
6 supermarket, grocery store, convenience store, nursing home, or food
7 or drink company.

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

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

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

15 2. A kratom product that is contaminated with a dangerous
16 nonkratom substance. A kratom product is contaminated with a
17 dangerous nonkratom substance if the kratom product contains a
18 substance that is not safe for human consumption;

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

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

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

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

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

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

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

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

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

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

1 6. Any federal food allergen labeling requirements, if
2 applicable, and clear and adequate directions for the consumption
3 and safe and effective use of such product, including the
4 recommended serving size, the number of servings in the container,
5 and the number of servings that can be safely consumed in a day.
6 Provided, liquid kratom products shall be packaged in a retail
7 container that has clear serving size markings and be subject to the
8 following requirements:

- 9 a. products of less than eight (8) fluid ounces which
10 contain more than three servings shall be accompanied
11 by a calibrated measuring device, and
- 12 b. if such a product contains more than the eight (8)
13 fluid ounces, the requirements specified in
14 subparagraph a of this paragraph do not apply.

15 ~~Provided further, packaging for powdered kratom products not in~~
16 ~~capsule form shall have a calibrated measuring device included in~~
17 ~~the container;~~

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

24

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

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

9 D. Upon request by the State Department of Health, the vendor
10 shall provide test results from a United States-based testing
11 facility, that is an independent testing laboratory as defined in
12 this act, to confirm the items listed on the product label.

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

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