SUBSTITUTE FOR HOUSE BILL NO. 4969

A bill to regulate certain conduct related to kratom and kratom products; to prohibit the manufacturing, processing, distribution, and sale of certain kratom products; to require labeling of certain kratom products; and to prescribe fines.

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

- Sec. 1. This act may be cited as the "kratom consumer protection act".
- 3 Sec. 3. As used in this act:
- 4 (a) "Certified independent third-party laboratory" means a
- 5 laboratory that is accredited to the International Organization for
- 6 Standardization, ISO/IEC 17025 standard, by an accreditation body
- 7 recognized by the International Laboratory Accreditation
- 8 Cooperation.





- 1 (b) "Kratom extract" means a substance or compound that meets 2 all of the following criteria:
- 3 (i) Is obtained by extraction of the Mitragyna speciosa leaf.
- 4 (ii) Contains more than trace amounts of Mitragyna speciosa and 5 other alkaloids of the Mitragyna speciosa plant.
- 6 (iii) Is intended for ingestion.
- 7 (iv) Does not contain any controlled substances or levels of 8 residual solvents higher than is allowed under the United States 9 Pharmacopeia 467.
- 10 (c) "Kratom leaf" means the leaf of the *Mitragyna speciosa*11 plant in fresh or dehydrated form.
- 12 (d) "Kratom product" means a consumable product that contains 13 any part of the kratom leaf or kratom extract. A kratom product is 14 not considered a food under the food law, 2000 PA 92, MCL 289.1101 15 to 289.8111.
- Sec. 5. A person that sells a kratom product must label the kratom product, and the label must clearly and conspicuously provide the following information:
- 19 (a) The following statements:
- (i) A statement against the use of the kratom product by individuals who are under 21 years, who are pregnant, or who are breastfeeding.
- (ii) A statement that kratom may be habit forming.
- (iii) A statement that reads as follows: "These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease."
- 28 (b) Directions for use of the kratom product that include, but 29 are not limited to, all of the following:



- 1 (i) A recommended serving size that is clearly described on the
- 2 label for each form of kratom product, such as capsules, gummies,
- 3 prepacked single-serving units, or similar product forms. If the
- 4 kratom product is a beverage, liquid, or loose powder, the label
- 5 must include clear instructions or a mark on the package or
- 6 container that clearly designates the recommended serving size.
- 7 (ii) The recommended number of servings that can be safely 8 consumed in a 24-hour period.
- 9 (iii) The recommended number of servings per container.
- 10 (iv) A list of the kratom alkaloid content and other
- 11 ingredients in the kratom product that includes quantitative, not
- 12 to exceed declarations of the amount per serving of each of the
- 13 following:
- 14 (A) Mitragynine.
- 15 (B) 7-hydroxymitragynine.
- Sec. 7. A person shall not manufacture, process, distribute,
- 17 sell, or offer for sale or distribution, in person or through an
- 18 online website, any of the following:
- 19 (a) A kratom product that contains a level of 7-
- 20 hydroxymitragynine in the alkaloid fraction that is greater than 2%
- 21 of the alkaloid composition of the kratom product.
- 22 (b) A kratom product that contains a synthesized material,
- 23 such as an alkaloid, metabolite, or alkaloid derivative that is
- 24 created by chemical synthesis or biosynthetic means, which
- 25 includes, but is not limited to, fermentation, recombinant
- 26 techniques, yeast-derived, or enzymatic techniques, rather than
- 27 traditional food preparation techniques, such as heating or
- 28 extracting, that synthetically alters the composition of kratom
- 29 alkaloid or constituent.



- 1 (c) A kratom product that does not include on its package or 2 label the amount of mitragynine and 7-hydroxymitragynine contained 3 in the kratom product.
 - (d) A kratom product that contains a kratom extract that was manufactured using extraction solvents that contain levels of residual solvents higher than is allowed under the United States Pharmacopeia 467 and not manufactured using food-grade solvents approved by the United States Food and Drug Administration.
 - Sec. 9. (1) Before a kratom product can be distributed, sold, or offered for distribution or sale, a person that manufactures or processes a kratom product shall have the kratom product tested by a certified independent third-party laboratory to ensure that the kratom product complies with the requirements of this act. If the kratom product complies with the requirements of this act, the certified independent third-party laboratory shall issue the person a certificate of analysis.
 - (2) The person shall maintain a certification of analysis issued under this section for 10 years and make the certificate of analysis available to a law enforcement officer on request. If a law enforcement officer makes a request to review a certificate of analysis issued under this section, the person shall produce the certificate of analysis within 30 days after the request is made.
 - Sec. 11. A person that violates section 5, 7, or 9 may be ordered to pay a civil fine of not more than \$1,000.00 for the first offense and not more than \$5,000.00 for a second or subsequent offense. The prosecutor of the county in which the violation occurred or the attorney general may bring an action to collect the fine. A fine collected under this section must be exclusively applied to the support of public libraries and county

- 1 law libraries in the same manner as is provided by law for penal
- 2 fines assessed and collected for violation of a penal law of this
- 3 state.

