



Classification of Mitragynine as a Schedule I Controlled Substance

Section 1: Summary

The Ohio Board of Pharmacy (Board), pursuant to section 3719.44 of the Ohio Revised Code, proposes the placement of mitragynine, the primary psychoactive alkaloid found in the *Mitragyna speciosa* plant, commonly known as kratom, into Schedule I.

Section 2: Background

Pursuant to section 3719.44 of the Ohio Revised Code, the Board may add or transfer a compound, mixture, preparation, or substance to Schedule I when it appears that there is a high potential for abuse, that it has no accepted medical use in treatment in this state, or that it lacks accepted safety for use in treatment under medical supervision.

In making a determination to add an unscheduled compound, the Board is required to consider the following 8 criteria:

- (1) The actual or relative potential for abuse;
- (2) The scientific evidence of the pharmacological effect of the substance;
- (3) The state of current scientific knowledge regarding the substance;
- (4) The history and current pattern of abuse;
- (5) The scope, duration, and significance of abuse;
- (6) The risk to the public health;
- (7) The potential of the substance to produce psychic or physiological dependence liability; and
- (8) Whether the substance is an immediate precursor.

Section 3: Evaluating Mitragynine Under the 8-Factor Criteria

(1) The actual or relative potential for abuse.

Mitragynine is the primary psychoactive alkaloid found in the *Mitragyna speciosa* plant, commonly known as kratom. Mitragynine accounts for up to 66 percent of the alkaloid content in kratom.ⁱ Manufacturers of natural kratom products profess that minor alkaloids in natural kratom are undetectable.ⁱⁱ Kratom has a long history of traditional use in Southeast Asia with the use of its leaf or extracts typically ingested orally for the treatment of pain and to aid in the performance of agricultural labor.ⁱⁱⁱ These effects have been attributed to mitragynine, which acts as a partial agonist at the human μ -opioid receptor.^{iv} The effects of kratom in humans are dose-dependent, with small doses producing a ‘cocaine-like’ stimulation while larger amounts cause ‘morphine-like’ sedative-narcotic effects.^v

Although mitragynine is the most abundant alkaloid in kratom and credited for its effects, the plant contains over 40 naturally occurring known alkaloids. One study examined the likely effects of 25 of those compounds. The study concluded that all the compounds share the most structural similarities with controlled opioid analgesics, such as morphine derivatives. Further analysis determined that 22 (including mitragynine) of the 25 compounds in kratom bind to μ -opioid receptors. The authors further noted:

The data from the PHASE model shows us that kratom compounds are predicted to affect the body just like opioids. Based on the scientific information in the literature and further supported by our computational modeling and the reports of its adverse effects in humans, we feel confident in calling compounds found in kratom, opioids.^{vi}

The FDA confirms that no marketer has sought to develop a drug that includes kratom in the US.^{vii} Further, the World Health Organization verifies that kratom and its alkaloids have never been licensed for therapeutic use.^{viii} Mitragynine has not been approved for medical use as a drug, nor is it generally recognized as safe by the FDA.^{ix}

Kratom is illegal in 33 countries, including Malaysia where it is found naturally.^x It is also banned in several states, including Alabama, Arkansas, Indiana, Vermont, Rhode Island, Louisiana, and Wisconsin.^{xi xii} A new law in Connecticut, effective October 1, 2025, requires the state’s Department of Consumer Protection (DCP) to designate kratom and 7-hydroxymitragynine as Schedule I controlled substances.^{xiii} On December 10, 2025, Connecticut’s DCP issued a notice of decision to move forward with the scheduling action.^{xiv}

Kratom has also been identified as a Drug of Concern by the US Drug Enforcement Administration (DEA).^{xv}

Despite its lack of approved use, kratom and its primary alkaloid, mitragynine, have been used to self-treat conditions such as chronic pain, fatigue, and psychiatric disorders (including substance use disorder and attention deficit hyperactivity disorder). For example, the Ohio Substance Abuse Monitoring Network reported about the availability of kratom in Ohio and its marketing towards individuals who are experiencing opioid use disorder. One respondent noted: “Advertisement for kratom [are] on Facebook...advertised as a way to get off of opiates, certain kinds of opiates; [Kratom] has similar withdrawal [symptoms as heroin]; Kratom mimics the effects of heroin...but the POs (parole officers) are aware of it now.” A treatment provider also reported that people are using kratom for its purported health benefits such as the treatment of pain, noting, “you can go to the store and buy [kratom] now...because people look up positive things about it (health benefits). It’s a pain reliever...it does give the effect of opiates or heroin...If someone has a little pain and they think it’s healthy and natural [then they may try it]....”^{xvi}

Still others begin using kratom for relaxation, euphoria, or to enhance performance.^{xvii} The end result for many consumers of these products has been an inability to control their use—actual abuse. Below are comments submitted to the Ohio Board of Pharmacy by consumers of products whose manufacturers profess are natural, herbal, or kratom.

- *Good morning. I understand the Ohio Board of Pharmacy is considering a ban on Kratom. I am in support of this ban. I have a family member that has become addicted to Feel Free made by Botanic Tonics. It was originally marketed as a kava drink and alcohol alternative. Come to find out the company is facing many lawsuits due to its false marketing. While they now say it includes kratom, it's too late for my family. The drink is extremely addictive and very strong. I don't believe that it is pure leaf kratom, there has to be more in this drink. It has created many issues for my family and severely negatively impacted my family's life. Please consider this ban. This drink is legal opioid and available at most gas stations. This drink is very addictive and dangerous. Please seriously consider the ban on kratom and more specifically Feel Free. Also, please reference the Reddit thread quitting feelfree, you will see all the people suffering from the dangerous and addictive qualities of this drink.*^{xviii}

- *Hello, I am a married wife of a kratom addict. I married my husband 7 years ago after he was clean off drugs (pills, cocaine, weed) for 8 years. He relapsed after 4 years into our marriage and has been addicted to kratom since. He was offered a sample by a gas station attendant ... stated "are you looking for energy, try this". Since then he takes up to 60 capsules a day lost 25 pounds and is in a constant state of irritability, speedy, highs, and super lows withdrawling ect. He tried detox once and relapse 2 days later. He says it's so hard to quit bc it's so easy to get and cheap. This garbage is horrible and I cannot believe kids can get a hold of this junk. I am pleading for your office to understand the severity of this "supplement". My soon to be ex-husband has now transitioned to the "7oh tabs" that are more potent and also easy to get apparently sold at smoke shops and cbd stores. I know it is the choice of the individual to pick up a substance and it's their responsibility to quit, but I worry about my kids (16 and 5) in their future if they stumble upon this gas station heroine. To whatever it's worth I just wanted to share my story.^{xix}*
- *My husband is a recovering opioid addict. Recently, he was introduced to kratom tablets and began taking them without telling me. When I first noticed the physical changes—his pupils shrinking and his energy levels spiking—I honestly believed he had relapsed on opioids. It was terrifying and brought back painful memories of our past struggles. He has now been taking kratom daily for months. He repeatedly tells me he plans to stop, but I can see that he cannot. He is worried about withdrawal and continues to use it despite the harm it is causing to his health and to our family. To me, this feels like we are back at square one, only with a substance that is currently unregulated and readily available. This is not the first time we have faced this danger. About a year ago, my husband was using the “blue bottle Feel Free” drinks (also containing kratom). After months of daily use, he ended up hospitalized with dangerously high liver enzyme levels and signs of serious liver damage. It took a long recovery, and now we are living through a repeat situation with kratom tablets. I know kratom is sometimes marketed as a “safe alternative” or a supplement, but my family’s experience shows otherwise. For people in recovery, it is not a harmless plant—it is another addictive drug that threatens to undo years of hard work and healing.^{xx}*
- *Someone very close to me began using kratom thinking it was a safe, natural way to manage stress and pain with out opiates. Over time, I watched it take over their life. They became anxious, detached, angry, and sick when they tried to stop. It broke my*

heart to see how something so easy to buy could cause such deep harm – not only to them, but to everyone who loves them. He would pass out while alone with our 22month old, he has been hospitalized with heart problems from use. We both have missed work, it has damaged our relationship, careers, children, and every aspect of our lives.^{xxi}

According to a recent news report, “many people who descended into kratom addiction say gas station products sucked them in. The ‘Quitting Kratom’ subreddit has 52,000 members and several posts a day from people documenting their journeys trying to quit kratom and 7-OH.”^{xxii} Medications approved by the FDA to treat opioid-use disorder have been used to treat individuals experiencing substance-use disorder associated with kratom. For example, a case report of a male in his 40s with a history of kratom use was successfully treated with buprenorphine/naloxone, which helped alleviate his withdrawal symptoms and allowed him to abstain from kratom.^{xxiii} Other case examples have been cited throughout medical literature to support the use of buprenorphine/naloxone to treat kratom-use disorder.^{xxiv xxv}

Another study conducted interviews with 395 kratom users and classified users into four distinct categories. Two classes included people using kratom for the self-treatment of chronic pain or substance use disorder and comprised approximately 52 percent of respondents. The other two classes, about 48 percent of respondents, reported using kratom to “get high.” The study also notes that individuals who reported using kratom to “get high” and who had the highest probability of kratom-use disorder were more likely to characterize kratom as “habit-forming, addictive, problematic, and inconsistent in its effects.”^{xxvi}

(2) The scientific evidence of the pharmacological effect of the substance.

The European Monitoring Centre for Drugs and Drug Addiction (EMCDDA) notes that kratom and its related compounds are selective and full agonists of the μ -opioid receptor. The effects of kratom in humans are dose-dependent, with small doses producing a ‘cocaine-like’ stimulation while larger amounts cause ‘morphine-like’ sedative-narcotic effects.^{xxvii}

A review published in 2024 notes that “mitragynine intake may be beneficial according to previous scientific data that presents its characteristics in antinociception, anti-inflammation, antidepressant, stimulant activity, sedative activity, anxiolytic activity, cognitive effects, and management of opioid withdrawal, which might be used as natural

remedies for various illnesses.” However, the authors noted that uncontrolled use of mitragynine may promote tolerance development and produce withdrawal symptoms upon abstinence.”^{xxviii}

One study conducted of kratom users in Malaysia found high rates of severe kratom dependence among regular users. More than half of regular users (those using for more than 6 months) reported severe kratom dependence, while 45% showed moderate kratom dependence. Physical withdrawal symptoms commonly experienced include muscle spasms and pain, sleeping difficulty, watery eyes/nose, hot flashes, fever, decreased appetite, and diarrhea. Psychological withdrawal symptoms commonly reported were restlessness, tension, anger, sadness, and nervousness. The study authors also noted that regular users who consumed 3 or more glasses of kratom per day had “higher odds of developing severe Kratom dependence, withdrawal symptoms, and inability to control Kratom craving.”^{xxix}

Medications approved by the FDA to treat opioid-use disorder have been used to treat individuals experiencing kratom-use disorder. Case examples have been cited throughout medical literature to support the use of buprenorphine/naloxone to treat kratom-use disorder.^{xxx} ^{xxxi} There are also examples of treating kratom withdrawal symptoms with medications such as clonidine and hydroxyzine that are often used off-label to manage opioid withdrawal.^{xxxii} ^{xxxiii} ^{xxxiv}

A case report of a male in his 40s with a history of kratom use was successfully treated with buprenorphine/naloxone, which helped alleviate his withdrawal symptoms and allowed him to abstain from kratom:

The patient was introduced to kratom 12 years ago by his brother for relaxation, stress relief, and as a substitute for oxycodone. He gradually increased his dose up to 40g of powder per day from an initial 6g/daily. He purchased kratom online and at a local gas station, and his weekly expenditure on the substance ranged from \$100 to \$600. Over time, he spent less time with family and friends, engaged in fewer recreational activities, and spent more time using kratom to relax at home. He realized that he was dependent on the substance when he experienced withdrawal symptoms such as diarrhea and anxiety when he skipped or reduced his use of the substance. He attempted to self-treat his addiction by decreasing his dose but experienced severe withdrawal symptoms, including stress, headaches, diarrhea, nausea, and dizziness. His inability to quit kratom

prompted him to seek medical help from his primary care physician. This was his first time seeking help from a physician. The patient was provided with the opportunity to connect with substance abuse clinics and other self-help groups, but he preferred pharmacological treatment.^{xxxv}

(3) The state of current scientific knowledge regarding the substance.

Little clinical dosing research exists for kratom and its primary alkaloid, mitragynine. Neither kratom nor mitragynine have progressed past Phase 1 clinical trials.^{xxxvi} ^{xxxvii} “Despite the increasing availability of human in vitro and in vivo animal data, rigorous fundamental information about the pharmacokinetics of kratom alkaloids in humans remains limited.”^{xxxviii}

Of the sparse pharmacokinetic research available, one study of twelve enrolled subjects monitored plasma levels of mitragynine and one of its metabolites, 7-hydroxymitragynine (7-OH), at various doses. The study found that 7-OH in plasma increased in a dose proportional manner after subjects consumed mitragynine contained in botanical kratom. The study also concluded that 7-OH steady state is reached after seven days of continuous dosing.^{xxxix}

An increasing amount of 7-OH in plasma that is proportional with an individual’s dose of mitragynine is important for two reasons. First, as discussed under Factor 4 below, mitragynine found in consumer products today is often several times more potent than traditional leaf products. Second, 7-OH is approximately 10 times more potent than mitragynine.^{xl} At just 10 times the potency of mitragynine, the effects and actual abuse of 7-OH prompted Food and Drug Administration (FDA) Commissioner Dr. Martin Makary to warn that it could be the next wave of the opioid crisis.^{xli}

Mitragynine has not undergone randomized, placebo-controlled studies necessary to demonstrate efficacy for any condition while minimizing the significant risks associated with it.^{xlii} “FDA has warned consumers not to use kratom because of the risk of serious adverse events, including liver toxicity, seizures, and substance use disorder (SUD). In rare cases, deaths have been associated with kratom use, as confirmed by a medical examiner or toxicology reports.”

Significantly, Phase I clinical studies explore the safety and tolerability of substances on 100 or fewer healthy adults.^{xliv} Most drugs—nearly 70 percent—that undergo Phase 1 clinical trials succeed to Phase 2 clinical trials. It is not until Phase 2 that research on a substance’s ability to treat a condition and its side effects are studied. This usually occurs with several hundred subjects who have the disease or condition in question. Approximately 33 percent of substances proceed to Phase 3 where efficacy is continued to be studied, and adverse effects are monitored on somewhere between several hundred to several thousand individuals with the disease or condition in question.^{xlv}

(4) The history and current pattern of abuse.

Kratom has been used for centuries in Southeast Asia by laborers for its stimulant and analgesic effects.^{xlvi} As previously discussed, kratom and its primary alkaloid, mitragynine, have been used to self-treat conditions such as chronic pain, fatigue, and psychiatric disorders (including substance use disorder and attention deficit hyperactivity disorder). Still others begin using kratom for relaxation, euphoria, or to enhance performance.^{xlvii}

Despite a long history of use originating from Southeast Asia, the widespread commercialization of kratom and its primary alkaloid, mitragynine, is a recent development. Both products sold as vegetation and those labeled as ground vegetation have been found to contain several times the amount of mitragynine than actual kratom leaf material.^{xlviii} Indeed, manufacturers of products marketed as “all-



Figure 1. Example of a manufacturer claim that its "all-natural and unadulterated" products are four times the strength of the kratom from which it originated.

“natural and unadulterated” acknowledge that extracts are several times greater than the plant from which they start (see Figure 1).^{xliv} Postmortem toxicology findings from patients who used kratom had concentrations “higher than those reported in Thai individuals consuming traditional kratom tea without adverse effects.”^l

As kratom commercialization has grown, so too have the number of adverse events reported to poison control centers. From January 1 to July 31, 2025, poison control centers in the United States (US) received 1,690 reports of exposure cases involving kratom, a number that already surpasses the total for all of 2024.^{li} Notably, adverse events associated with items that can be categorized as supplements are underreported. At least one source estimates the reporting rate for adverse events related to supplements to be approximately 2 percent.^{lii} Reports of kratom exposure have been associated with opioid toxicities including seizures, agitation, and death.^{liii}

The Ohio Substance Abuse Monitoring (OSAM) Network report from January 2025 found kratom use associated with opioid use, as the substance reportedly has similar effects, and kratom is reportedly advertised to help mitigate opioid withdrawal symptoms and stop opioid use. While its notoriety appears to be causing drug screening to expand, there is evidence that some individuals consumed kratom because it would not appear on drug screens. One testimonial provided, “[a]dvertisements for kratom [are] on Facebook ... advertised as a way to get off of opiates, certain kinds of opiates; [Kratom] has similar withdrawal [symptoms as heroin]; Kratom mimics the effects of heroin ... but the POs (parole officers) are aware of it now.”^{liv}

Access to mitragynine can be further abused when it is isolated from kratom leaves. It can then be synthetically transformed into 7-OH and other semi-synthetic opioids—some with significantly enhanced μ -opioid receptor activity. One such compound, MGM-15 (dihydro-7-hydroxy mitragynine), first described in 2014, is a highly potent semi-synthetic opioid that can be synthesized from 7-hydroxymitragynine in a single step. In September 2025, the presence of MGM-15 was confirmed in commercially available tablets.^{lv} These compounds were legally available in Ohio until rule 4729:9-1-01.1 of the Ohio Administrative Code became effective on December 12, 2025.^{lvii}

(5) The scope, duration, and significance of abuse.

Although there is apparent consensus that the number of Americans who use kratom is in the millions, the precise number of those who use it is more nebulous. The Substance Abuse and Mental Health Services Administration estimated that in 2021, 1.7 million Americans aged 12 and older used kratom.^{lvii} More recently, Harvard Medical School estimated that between 2 and 15 million Americans may be using kratom.^{lviii}

The Ohio Substance Abuse Monitoring (OSAM) Network report from January 2025 found kratom use associated with opioid use, as the substance reportedly has similar effects, and kratom is reportedly advertised to help mitigate opioid withdrawal symptoms and stop opioid use. They stated: “[a]dvertisements for kratom [are] on Facebook ... advertised as a way to get off of opiates, certain kinds of opiates; [Kratom] has similar withdrawal [symptoms as heroin]; Kratom mimics the effects of heroin ... but the POs (parole officers) are aware of it now.”^{lix}

One research group prioritized its study in part because, “[t]eas and powders made from the kratom ... plant are gaining in popularity due to their opioid-like effects.”^{lx} A treatment provider in the Cleveland region spoke about the promoted health benefits of kratom, saying, “[y]ou can go to the store and buy [kratom] now ... because people look up positive things about it (health benefits). It's a pain reliever ... it does give the effect of opiates or heroin.... If someone has a little pain and they think it's healthy and natural [then they may try it]...”^{lxii}

Reportedly, kratom is desirable to some because it is not always included on drug screens. Treatment providers observed, “Kratom is still one of those things that's out there that's being abused, that's not being caught [through drug screening]; I spoke to our nurse practitioner, and in small doses [kratom] doesn't show up on the [drug] screen. It's because people are taking way high doses that are detectable [on drug screens] that are not good for you, you know, addiction wise; A lot of places don't test for it [on drug screens]. So, until we started testing for it, I didn't hear that much about it, but now our lab tests routinely include kratom.”^{lxii}

Public comments serve as further evidence of the significance of kratom abuse in Ohio. As discussed in the analysis of Factor 1, the Board of Pharmacy received comments from several individuals regarding the impact of kratom use. Reported impacts include relapses by those

in recovery, significant weight loss, inability to cease use despite adverse impacts on personal relationships and finances, liver damage, and heart issues.

Finally, a reality of modern drug misuse is that an increasing number of individuals engage in polydrug use with little to no knowledge of contraindications. This is true of both illicit drugs and prescription medication. From 2019 to 2023, the amount of unintentional drug overdose deaths in Ohio that involved multiple drugs ranged from 58 to 69 percent.^{lxiii} Given this context, it would be irresponsible to discount the risk of polydrug use.

Kratom is no different. “In parallel with the increase in popularity of kratom is an increase in kratom-associated overdose deaths when concomitantly used with other drugs.”^{lxiv} The impact of consuming kratom with other substances is described in two Reddit posts from those impacted by kratom misuse found in Appendix A. These posts warn that kratom consumption with common medications can cause heart issues and even death.

(6) The risk to the public health.

“FDA has warned consumers not to use kratom because of the risk of serious adverse events, including liver toxicity, seizures, and substance use disorder (SUD). In rare cases, deaths have been associated with kratom use, as confirmed by a medical examiner or toxicology reports.”^{lxv}

In Ohio, a growing number of deaths associated with kratom have been reported, indicating that it is a risk to public health. From 2019 to 2024, the Ohio Department of Health reported at least 202 deaths in which kratom is listed as a cause of death (see Table 1).

Table 1. Number of Unintentional Drug Overdose Deaths Involving Kratom (Mitragynine), 2019 - 2024^{bvi}

Year	Total No. of Overdose Deaths	No. of Deaths with Kratom Detected in Postmortem Toxicology	% of Total Overdose Deaths with Kratom Detected in Postmortem Toxicology	No. of Deaths with Kratom Detected in Postmortem Listed as a Cause of Death	% of Deaths with Kratom Detected in Postmortem Toxicology where Kratom Was Listed as a Cause of Death	No. of Deaths with Kratom Detected in Postmortem Toxicology That Were Also Positive for Other Substances	% of Deaths with Kratom Detected in Postmortem Toxicology That Were Also Positive for Other Substances
2019	3,962	19	0.5%	15	78.9%	17	89.5%
2020	4,943	34	0.7%	29	85.3%	29	85.3%
2021	5,174	49	0.9%	37	75.5%	41	83.7%
2022	4,916	47	1.0%	40	85.1%	44	93.6%
2023	4,461	52	1.2%	37	71.2%	43	82.7%
2024*	3,015	54	1.8%	44	81.5%	43	79.6%
Total	26,471	255	0.9%	202	79.2%	217	85.1%

*Data for 2024 is not yet complete.

Several forensic cases of opioid-like deaths with lung congestion and high postmortem mitragynine blood concentrations with no obvious alternative causes of death have now been reported.^{lxvii} ^{lxviii} There is also a report of naloxone being successfully used to reverse kratom intoxication.^{lxix} This evidence suggests that high enough doses of kratom alkaloids may cause opioid toxicity via the usual opioid-receptor pathways implicated in conventional opioid overdoses.^{lx}

In 2019, Utah developed a regulatory scheme that allowed continued access to kratom over-the-counter. Despite age restrictions and limited alkaloid access, overdose deaths related to kratom and treatment for kratom use disorder persisted. One emergency room physician reported patients sharing that they “had no idea it was just like a narcotic,” and that his patients experience the same withdrawal symptoms as those withdrawing from narcotics.^{lxxi} The Utah State Medical Examiner reported nearly 160 kratom-related deaths over a five-year period. While most of those overdoses involved polysubstance use, the state saw almost 50% as many kratom-only deaths in the most recent 12-month reporting period as Utah had seen since 2014.^{lxxii} Consequently, the Utah legislature proposed to add all alkaloids from kratom and their analogs to the state’s list of controlled substances in November 2025. The bill sponsor originally voted to regulate kratom in 2019 but now “sees kratom in all forms as a dangerous opioid masquerading as a supplement.”^{lxxiii}

There are also documented cases of kratom use by expecting mothers resulting in neonatal abstinence syndrome for newborns.^{lxxiv lxxv lxxvi lxxvii} These cases are particularly concerning given that the promotion of kratom products as safe alternatives to opioids may well persuade expecting mothers to consume kratom products to avoid the adverse impacts of opioids on their unborn child. The concern is compounded by the tendency of consumers to believe that the legal status of a substance means that it is a safe alternative to a controlled substance.^{lxxviii}

At least one manufacturer reinforces this notion with misleading statements like, “[f]eel better with the only kratom products studied in clinical trials,” and “US Made in our FDA-registered

space in Florida” (see Figure 2).^{lxxix} The FDA is unequivocal in its warning against kratom use

due to its adverse effects.^{lxxx} “There are no FDA-approved drugs containing kratom.”^{lxxxi} According to the FDA, kratom and mitragynine are also unlawful “when added to conventional foods, as dietary supplements, [and] as ingredients in any FDA-approved drug.”^{lxxxii}



Figure 2. Claims made by a manufacturer of self-proclaimed “pure, unadulterated” kratom.

Evidence supports the position that the ingredients in kratom available to consumers are void of robust quality control. In 2019, FDA conducted laboratory testing on 30 different kratom products from a variety of sources. The analysis found “significant levels of lead and nickel at concentrations that exceed safe exposure for oral daily drug intake.”^{lxxxiii} A third-party analysis of kratom products also found samples that exceeded daily exposure limits for lead. The authors noted that “non-extract products (powders, capsules, tablets) generally contain greater concentrations of elemental impurities than extract products or the soda preparation.”^{lxxxiv}

In 2018, the CDC investigated a multistate outbreak of *Salmonella* infections linked to kratom. The agency reported that a total of 199 people were infected with outbreak strains of *Salmonella* across 41 states, including 7 individuals in Ohio. Of those 199, thirty-eight percent of those individuals were hospitalized. CDC investigators stated that “epidemiologic and laboratory evidence indicated that kratom was the likely source of this multistate outbreak.”^{lxxxv}

The presence of lead and other heavy metals may increase the risk of adverse health outcomes for regular kratom users.^{lxxxvi} For example, a case report from Iowa found a patient with suspected Fanconi syndrome from cadmium toxicity due to heavy kratom use. The report found high levels of cadmium in both the patient’s blood stream and kratom the patient consumed.^{lxxxvii}

According to the World Health Organization, chronic kratom use has been associated with at least 92 cases of liver toxicity. They note that common presenting signs and symptoms include abdominal discomfort, jaundice, pruritus, and dark-colored urine. These typically start about 3 weeks after initiation of kratom use and resolve within one month of stopping kratom use.^{lxxxviii}

Additionally, there are no age verification requirements to prevent the sale of kratom products to children. This raises serious concerns about the impact of these compounds on the developing brain. The 2021 National Survey on Drug Use estimates that at least 45,000 adolescents in the US (ages 12-17) used kratom in the past year. This presents public health risks to adolescents given studies showing “adolescent kratom exposure particularly mitragynine and LKD [lyophilised kratom decoction] may cause selective cognitive and behavioural deficits. The brain metabolite profiles further suggest that the altered metabolic pathway (i.e., arachidonic acid, pantothenate and CoA, and tryptophan) may underlie the kratom-induced cognitive and behavioural deficits. Together, these findings demonstrate that adolescents’ brain is sensitive to the impact of early kratom exposure during this critical development period.”^{lxxxix}

(7) The potential of the substance to produce psychic or physiological dependence liability.

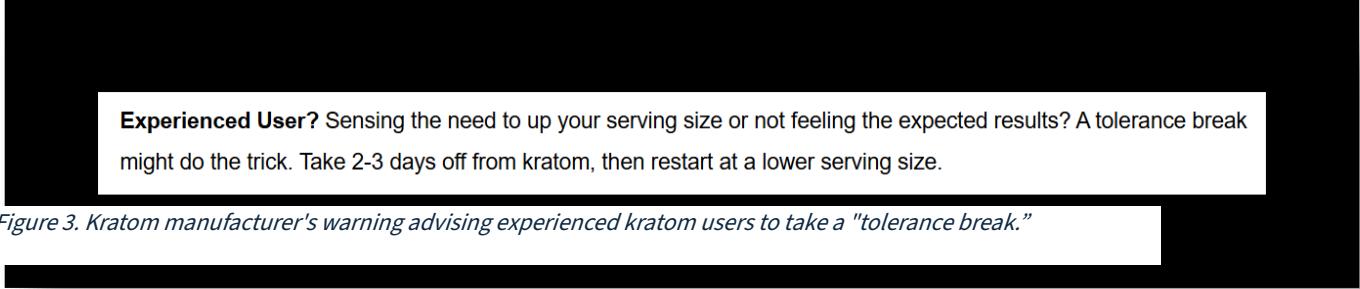
Cases of dependence to kratom have been documented for the last several decades.^{xc} Mitragynine has agonist activity at μ -opioid receptors and may lead to dependence and addiction.^{xcii} Further, the European Union Drug Agency, FDA, DEA, Mayo Clinic, and others agree that individuals can develop a dependence to kratom.^{xcii} According to the FDA, “[c]ases of kratom-related SUD have also been observed. In these cases, individuals met certain criteria for SUD, including using kratom for longer than intended, using more kratom than intended, having cravings for kratom, continuing to use kratom despite adverse consequences (either physically or in their personal life), increasing the amount of kratom used to produce the same effect (tolerance), and experiencing withdrawal symptoms when kratom use was stopped (physical dependence).”^{xciii}

In-person interviews with convenience samples of long-term daily oral kratom users in Southeast Asia find up to three-quarters reporting withdrawal symptoms after cessation of use. Typical withdrawal symptoms include irritability, anxiety, depression, sleep disturbance, lacrimation, rhinorrhea, muscle and bone pain, muscle spasm, diarrhea, and decreased appetite. The likelihood and severity of withdrawal are positively associated with duration, frequency, and intensity of kratom use.^{xciv}

Recent news reports also highlight the psychic or physiological dependence of kratom. For example, an Akron woman reported developing “a debilitating kratom addiction” after consuming a product containing the substance thinking it was a safe, “plant-based” drink. She ended up losing her house, marriage, and half of her body weight because of her exposure and subsequent addiction to this compound. The woman reports “her addiction would have taken her life too, had she not gone to rehab in 2024.”^{xcv}

Another news report demonstrates the danger of having these products readily accessible. An individual using a popular kratom-based drink available at local gas stations reported spending about \$2,000 per month (consuming about 10 to 12 bottles per day). According to the report, he “...soon found that after the immediate euphoria from the shot, he would be hit with a cycle of unpleasant symptoms, including a runny nose and achy body.” The individual closed down their business and had to seek in-patient treatment.^{xcvi}

Finally, it is not unusual for kratom manufacturers themselves to acknowledge the risk of developing a dependence to kratom. One such acknowledgement provides, “[s]ensing the need to up your serving size or not feeling the expected results? A tolerance break might do the trick. Take 2–3 days off from kratom, then restart at a lower serving size” (see Figure 3).^{xcvii}



Experienced User? Sensing the need to up your serving size or not feeling the expected results? A tolerance break might do the trick. Take 2-3 days off from kratom, then restart at a lower serving size.

Figure 3. Kratom manufacturer's warning advising experienced kratom users to take a "tolerance break."

Despite warnings that kratom may be habit forming and should not be consumed every day, testimonials posted by kratom manufacturers encourage daily use for relief of discomfort, stating, “this is my favorite kind of kratom. . . It is the best for relieving daily discomfort in your mind or your body.”^{xcviii} Additional examples from a variety of manufacturers warning of the risk of addiction can be found in Appendix B.

(8) Whether the substance is an immediate precursor.

Mitragynine is a known precursor to mitragynine-related compounds, which are schedule I controlled substances in Ohio. According to the European Union Drug Agency, “[t]he chemical total syntheses reported for several kratom alkaloids are too complex to be used for economic production of any [of] these compounds. However, mitragynine can serve as a chemical precursor to the more potent 7-hydroxymitragynine.”^{xcix}

Mitragynine is highly likely to follow the same pattern seen with other drug classes like bath salts. Bath salts were created by modifying cathinones, a primary active ingredient found in a plant called khat.^c As soon as one of those modifications was identified and scheduled, a new cathinone-like compound was developed to evade the law.

Looking to past illicit drug patterns, unexpected ways to synthesize mitragynine-related compounds will be established. At least one pathway would use mitragynine to make 7-OH and then MGM-15, the previously discussed potent opioid already identified in commercially

available tablets.^{ci} Should unfettered access to mitragynine remain, we can anticipate the creation of additional potent novel compounds that are marketed to the general public, including children.

Section 4: Finding of the Board

Pursuant to section 3719.44 of the Ohio Revised Code, the Board may add or transfer a compound, mixture, preparation, or substance to Schedule I when it appears that there is a high potential for abuse, that it has no accepted medical use in treatment in this state, or that it lacks accepted safety for use in treatment under medical supervision.

After a review of all available data, the Board finds mitragynine has a high potential for abuse, has no accepted medical use in treatment in this state, and that it lacks accepted safety for use in treatment under medical supervision.

Section 5: Resolution of the Board

Based on these findings, the Board hereby authorizes the filing of rule 4729:9-1-01.2 of the Administrative Code with the Common Sense Initiative and the Joint Committee on Agency Rule Review to classify as a schedule I opiate or opiate derivative any material, compound, mixture, or preparation that contains mitragynine.

Section 6: Proposed Rule

4729:9-1-01.2 – Mitragynine (NEW)

Notwithstanding any other provision of the Administrative Code, the following is classified as a schedule I controlled substance opiate or opiate derivative:

(A) Mitragynine ((α E,2S,3S,12bS)-3-ethyl-1,2,3,4,6,7,12,12b-octahydro-8-methoxy- α -(methoxymethylene)-indolo[2,3-a]quinolizine-2-acetic acid, methyl ester) and / or synthetic substances, derivatives, prodrugs, isomers, esters, ethers, salts, and salts of isomers with similar chemical structure.

Appendix A - Reddit Comments

Kratom had a toxic effect with the antidepressant and antipsychotic meds and killed my daughter

My 24 yo daughter died from mixing her regular prescribed psych meds with kratom. Her full autopsy listed the cause as poly drug toxicity and her prescribed meds were all in therapeutic levels.

The toxicology and full autopsy report confirmed what was in her system:

- Aripiprazole (antipsychotic)
- Lamotrigine (mood stabilizer)
- Venlafaxine (antidepressant)
- Mitragynine (the active compound in kratom)

There was no warning on the kratom for interactions with other meds or the dangers. This stuff is sold in smoke shops and gas stations, over the counter and marketed as "natural" and "safe".

I was with her when she died. She was scared. She said she didn't want to die. She felt that something was wrong and her chest hurt. Her heart was beating so fast, then she had cardiac arrest and died right in front of me. CPR could not bring her back. The EMTs gave her narcan, it didn't help.

I've since learned that Kratom AMPLIFIES the toxicity of certain meds (psych meds specifically) and overwhelms the nervous system.

My daughter took what was prescribed to her, and she bought and took something she thought was safe, a supplement.

Kratom is NOT SAFE if you are taking other medications. Please share this information. I know she is not the only one this has happened to and if only there were warnings, it could have saved her life

stop tucking romanticizing and glamorized kratom.

Venting

i am currently in hell. i overdosed and am experiencing the wobbles. it is literally an opioid that causes addiction and ill side effects. i would've been okay had i not done this shit. we are responsible for managing our addiction but there are no proper warnings, consumers are unaware how dangerous this is. have no idea HOW it's still legal and to the kratom fans and fiends defending the hell out of it: you can and most likely will go through what im going through. i can't drive for a few days. this could easily be you



Archived post. New comments cannot be posted and votes cannot be cast.

45

96

Share

Sort by: Best

Search Comments

AutoModerator MOD • 10mo ago • Stickied comment



[deleted] • 10mo ago

One of the darkest times in my life was going through withdrawals for kratom. I'll never touch that shit again

KRATOM WARNING: Do not mix with these medications (heart problems)

Let me preface this by saying I've mixed kratom with tramadol myself and had no problems, but that doesn't mean it's safe. Always air on the side of caution.

I recently found out that kratom is something called a QTc interval prolonger. QTc intervals are an EKG reading. If you mix QTc prolongers, it can potentially cause a heart arrhythmia reaction called torsades de pointes arrhythmia. Many common medications and drugs including psych meds and tramadol are QTc interval prolongers.

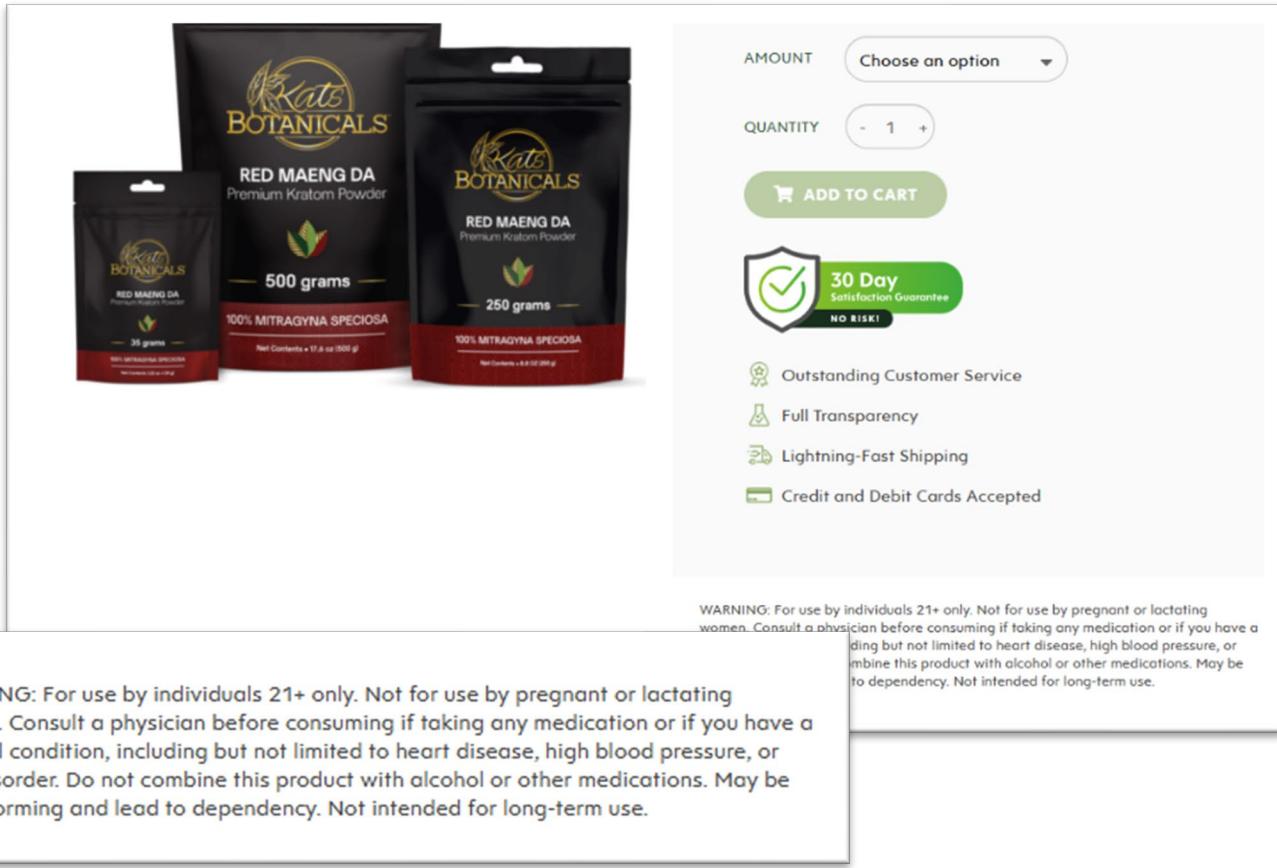
I never heard any warnings about mixing kratom with these medications. This information is important and very very unknown.

If you want to know what medications can cause a torsades de pointes arrhythmia, this is a good place to start (although they require registration which is free) <https://crediblemeds.org/index.php/login/dlcheck>

Click all the AVAILABLE TDP RISK CATEGORIES, set show entries to all, and sort by name or category. There are also usually pubmed articles if you look for them, that will tell you whether a medication is or isn't one.

here is my source for kratom causing this reaction <https://www.ncbi.nlm.nih.gov/pubmed/25535742>

Appendix B – Product Labels & Examples



Kats Botanicals
RED MAENG DA
Premium Kratom Powder

RED MAENG DA
Premium Kratom Powder

RED MAENG DA
Premium Kratom Powder

Choose an option

Quantity: 1

ADD TO CART

30 Day Satisfaction Guarantee
NO RISK!

Outstanding Customer Service

Full Transparency

Lightning-Fast Shipping

Credit and Debit Cards Accepted

WARNING: For use by individuals 21+ only. Not for use by pregnant or lactating women. Consult a physician before consuming if taking any medication or if you have a medical condition, including but not limited to heart disease, high blood pressure, or liver disorder. Do not combine this product with alcohol or other medications. May be habit-forming and lead to dependency. Not intended for long-term use.

Source: <https://katsbotanicals.com/product/red-maeng-da-kratom-powder> (Last Accessed Jan 3, 2026)



WARNINGS AND POTENTIAL SIDE EFFECTS:
using *Mitragyna Speciosa* a/k/a kratom or kratom-related products can be dangerous.

There have been reports of adverse health effects associated with using kratom products, including the US Food & Drug Administration who states that it exposes users to the risks of addiction, abuse and dependence and has not been approved for human consumption. Some publications have suggested kratom may be associated with serious potential side effects, including but not limited to, weight loss, dry mouth, chills, nausea, vomiting, changes in urine and constipation, liver damage, muscle pain, withdrawal, addiction, abuse, tachycardia, hepatotoxicity and as to the brain and nervous system; dizziness, drowsiness, hallucinations, delusions, depression, breathing suppression, hostile attitude, insomnia, seizures, coma and death.

You are advised to review the US Food & Drug Administration's website for further information regarding this product and its use. Furthermore, the US Drug Enforcement Administration currently does not schedule kratom as a controlled substance but has been listed as a Drug and Chemical of Concern.

Source: https://amazingbotanicals.net/product/platinum-kratom-extract-tablets-80-ultra-purified-alkaloids/?srsltid=AfmBOopW-lgl9MBHQuhAoFJ_bvVLkeKilgZoygOXYMvQH8N02PF3ZbAZ (Last Accessed Jan 3, 2026)

ADULTS 21+ ONLY
KEEP OUT OF REACH OF CHILDREN

Supplement Facts

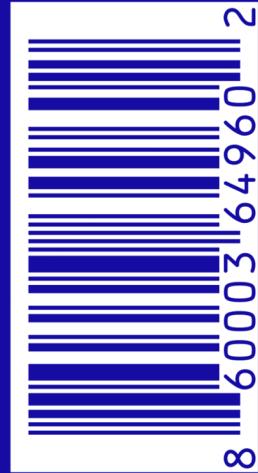
Serving Size: 1 fl oz (29.6 mL)
Servings Per Container: 2

	Amount Per Serving	%DV
Iron	0.4 mg	4%*
Kava root (extract)		
Total Kavalactones	260 mg	†
Leaf kratom (ground)		
Total Alkaloids	34 mg	†
Mitragynine	20 mg	†
7-hydroxymitragynine	<0.05 mg	†

* Percent Daily Value (DV) based on a 2,000 calorie diet.

† Daily Value (DV) not established.

Other ingredients: water, pineapple juice, natural flavors, stevia leaf, citric acid.



REV #007

**DIRECTIONS: ONLY TAKE 1 OZ (1/2 BOTTLE) BY MOUTH AT A TIME.
DO NOT EXCEED 2 OZ (1 BOTTLE) IN A 24-HOUR PERIOD.
SHAKE WELL. REFRIGERATE AFTER OPENING.**

Warning: This product contains leaf kratom which, like caffeine and alcohol, may be habit-forming and harmful if consumed irresponsibly. Avoid if you have a history of substance abuse. When consumed as recommended, *feel free* CLASSIC has not been shown to cause any serious physical or social harm. **Caution:** Not for consumption by or sale to persons under the age of 21. May interact with certain medications—consult a licensed, qualified healthcare professional before use. Do not consume with excessive alcohol. **This product is not intended for those who are sensitive to the active ingredients or women who are pregnant, nursing, or trying to become pregnant.**

These statements have not been evaluated by the FDA. This product is not intended to diagnose, treat, cure or prevent any disease or illness.

BOTANIC TONICS, LLC
13105 E. 61ST STREET, SUITE B • BROKEN ARROW, OK 74012

Source: <https://botanictonics.com/products/feel-free-classic-tonic> (Last Accessed Jan 3, 2026)

BEST SELLER



GET PERKED UP ON PLANTS.

Super SPECIOSA RAW LEAF

ALL NATURAL GREEN MAENG DA KRATOM POWDER NET WT 250g

30% OFF ON ORDERS \$100+

USE CODE: COUNTDOWN30

Tested by Cora Science on NOV 4, 2025

View results →

Description Ingredients Shipping & Returns Disclaimer

These products are not intended to diagnose, treat, cure, prevent or mitigate any disease. Not for sale to individuals under the age of 21. Product should only be used as directed on the label. Do not use if pregnant or nursing. Consult with a doctor before use and for possible interaction with drugs. Do not take kratom while operating heavy machinery. May be habit forming.

Source: https://superspeciosa.com/products/green-maeng-da-kratom-powder?_ab=0&_fd=0&_sc=1&selling_plan=5215191261 (Last Accessed Jan 3, 2026)

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