

Kratom-Induced Liver Injury: A Case Series and Clinical Implications

Mahesh Botejue ¹, Gurjot Walia ², Omar Shahin ³, Jyotsna Sharma ⁴, Rasiq Zackria ³

¹. Graduate Medical Education, Riverside Community Hospital, Riverside, USA ². Internal Medicine, University of California (UC) Riverside, Riverside, USA ³. Internal Medicine, Riverside Community Hospital/University of California (UC) Riverside, Riverside, USA ⁴. Internal Medicine, Riverside Community Hospital, Riverside, USA

Corresponding author: Mahesh Botejue, mahesh.botejue@hcahealthcare.com

Abstract

Drug-induced liver injury (DILI) is among the most common causes of acute liver injury and acute liver failure in the United States. Kratom is an herbal supplement made from the leaves of a tropical evergreen tree (*Mitragyna speciosa*) that is native to Southeast Asia. Due to its psychotropic and opioid-like activity, there has been an increase in its use as a recreational drug. Despite this increase, little is known regarding the toxicities and adverse effects though it is known to cause DILI. We present two cases of DILI associated with Kratom use.

Categories: Internal Medicine, Gastroenterology

Keywords: dili, kratom, acute liver injury

Introduction

Drug-induced liver injury (DILI) is one of the most common etiologies of acute liver failure (ALF) in the United States. Patients may present with a range of clinical manifestations from an asymptomatic transaminase elevation to ALF [1-3]. Clinically, DILI has been described as the rise of alanine aminotransferase (ALT) or aspartate aminotransferase (AST) greater than five times the upper limit of normal (ULN), alkaline phosphatase (ALP) greater than two times the ULN, or bilirubin greater than two times the ULN with an associated rise of AST and ALT [4-5]. While certain over-the-counter and prescription medications have been known to cause liver injury, the incidence of herbal and dietary supplements as a precursor has been rising.

Case Presentation

Case 1

A 27-year-old man with no reported medical history presented with sharp, non-radiating, epigastric abdominal pain, which started one day earlier. He denied any inciting event and reported the pain had been constant since onset. Associated symptoms included fevers, chills, and nausea; he denied vomiting or changes in bowel habits. He denied any alcohol or drug abuse. He did, however, endorse a two-week history of taking Kratom for foot osteoarthritis. Physical examination was unremarkable on presentation except for abdominal pain on palpation. Initial lab work demonstrated elevated liver chemistries of AST 345 U/L, ALT 404 U/L, ALP 115 U/L, and total bilirubin of 1.3 mg/dL. His liver synthetic function tests were normal. A urine drug screen was positive for opiates and tetrahydrocannabinol. Infectious workup, autoimmune panel, salicylate, and acetaminophen toxicity were negative. Imaging studies, including a right upper quadrant ultrasound (US) and magnetic resonance cholangiopancreatography (MRCP), were unremarkable. He was treated symptomatically for DILI from Kratom use. His liver chemistries down-trended with a resolution of his symptoms, and he was discharged home in stable condition.

Case 2

A 36-year-old woman, with a history of hepatic steatosis, ventral hernia repair complicated by two episodes of dehiscence, opioid dependence, and recent choledocholithiasis status post biliary stent placement five months prior, with subsequent removal one month ago and cholecystectomy, presented for worsening jaundice of one-month duration. She reported the onset of symptoms after biliary stent removal and also endorsed lethargy, poor appetite, non-bilious vomiting, and watery diarrhea. She denied any medication, alcohol, or drug use. On presentation, physical examination demonstrated jaundice, scleral icterus, moderate ascites with a positive fluid wave, and pitting edema. Lab work revealed leukocytosis 16.8 K/mm³, sodium 132 mmol/L, hyperbilirubinemia 39.5 mg/dL, AST 138 U/L, ALT 28 U/L, ALP 156 U/L, international normalized ratio (INR) of 2.7, and albumin of 2.3 g/dL. Other lab studies included an unremarkable autoimmune and hepatitis panel. The US demonstrated fatty liver with no bile duct dilatation. MRCP revealed hepatomegaly with diffuse fatty infiltration without biliary tree dilatation. Given no identifiable biliary or hepatocellular etiology, a core liver biopsy demonstrated cholestatic hepatitis consistent with DILI, and steatohepatitis with marked pericellular fibrosis and scarring. On further questioning, she

How to cite this article

Botejue M, Walia G, Shahin O, et al. (April 25, 2021) Kratom-Induced Liver Injury: A Case Series and Clinical Implications. Cureus 13(4): e14679. DOI 10.7759/cureus.14679

endorsed using an herbal supplement, Kratom, for several years to aid with her opioid abuse and withdrawal. Due to a persistently poor model for end-stage liver disease (MELD)-Na score, she was transferred to a liver transplant center for appropriate management.

Discussion

Kratom, known by many street names, including “thang,” is available in several forms, including powder, tea, extract, tablets, or capsules. It behaves as a stimulant at low doses, opioid-like at moderate doses, and sedative at high doses [6-9]. With the recent promulgation of over-the-counter supplements, physicians must be familiar with their various forms, uses, and toxicities.

In the US, poison centers have reported an increasing number of calls regarding exposure to Kratom [10]. A literature review studied the cases of DILI in the Drug-Induced Liver Injury Network and found eight out of 404 cases to be associated with Kratom [11]. This review determined a causal association in seven out of eight of those cases, with the median Kratom use of 22 days before the onset of injury and reported symptoms of jaundice, itching, abdominal pain, or fever. Of these, all patients recovered, with six hospitalized. This herbal substance is currently considered a “drug of concern” by the Drug Enforcement Administration (DEA), which initially was to classify the extract as a Schedule I drug, but the classification was postponed [12].

Although the hepatotoxicity of Kratom has been suggested, it is unclear to what extent any byproducts or contaminants may have a role in liver injury. A recent comprehensive review discussed the mechanisms of associated hepatotoxicity and suggested that physiologic metabolism is likely hepatic [13]. However, case reports have demonstrated a cholestatic or mixed pattern as well [14].

One of the more likely explanations for the increasing number of Kratom-associated DILI is the opioid epidemic and policymaking shift, including Centers for Disease Control and Prevention (CDC) guidelines, which discourage physicians from prescribing opioids for non-cancer pain [15]. A majority of Kratom users reported using the drug as a treatment for acute or chronic pain, endorsed fewer side effects, and less addiction potential than with opioids [16]. As chronic pain and opioid prescription patterns evolve, this has the potential to encourage the use of less regulated compounds as both an adjunct and replacement for traditional pain control methods [17].

Conclusions

These cases highlight a rare but serious complication of DILI associated with Kratom use. This contributes to the growing body of evidence supporting Kratom as a cause of liver injury. A thorough history of medication and supplement use is critical for patients with hepatic dysfunction or possible suspicion of DILI.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Suh JI: Drug-induced liver injury. *Yeungnam Univ J Med.* 2020, 37:2-12. [10.12701/yujm.2019.00297](https://doi.org/10.12701/yujm.2019.00297)
2. Larson AM, Polson J, Fontana RJ, et al.: Acetaminophen-induced acute liver failure: results of a United States multicenter, prospective study. *Hepatology.* 2005, 42:1364-72. [10.1002/hep.20948](https://doi.org/10.1002/hep.20948)
3. Ostapowicz G, Fontana RJ, Schiødt FV, et al.: Results of a prospective study of acute liver failure at 17 tertiary care centers in the United States. *Ann Intern Med.* 2002, 137:947-54. [10.7326/0003-4819-137-12-200212170-00007](https://doi.org/10.7326/0003-4819-137-12-200212170-00007)
4. Fontana RJ, Seeff LB, Andrade RJ, Björnsson E, Day CP, Serrano J, Hoofnagle JH: Standardization of nomenclature and causality assessment in drug-induced liver injury: summary of a clinical research workshop. *Hepatology.* 2010, 52:730-42. [10.1002/hep.23696](https://doi.org/10.1002/hep.23696)
5. Devarbhavi H: An update on drug-induced liver injury. *J Clin Exp Hepatol.* 2012, 2:247-59. [10.1016/j.jceh.2012.05.002](https://doi.org/10.1016/j.jceh.2012.05.002)
6. Kratom. LiverTox: Clinical and Research Information on Drug-Induced Liver Injury [Internet]. National Institute of Diabetes and Digestive and Kidney Diseases, Bethesda (MD); 2012.
7. Prozialeck WC, Jivan JK, Andurkar SV: Pharmacology of Kratom: an emerging botanical agent with stimulant, analgesic and opioid-like effects. *J Am Osteopath Assoc.* 2012, 112:792-9.
8. Hassan Z, Muzaimi M, Navaratnam V, et al.: From Kratom to mitragynine and its derivatives: physiological and behavioural effects related to use, abuse, and addiction. *Neurosci Biobehav Rev.* 2013, 37:138-51. [10.1016/j.neubiorev.2012.11.012](https://doi.org/10.1016/j.neubiorev.2012.11.012)

9. Warner ML, Kaufman NC, Grundmann O: The pharmacology and toxicology of kratom: from traditional herb to drug of abuse. *Int J Legal Med.* 2016, 130:127-38. [10.1007/s00414-015-1279-y](https://doi.org/10.1007/s00414-015-1279-y)
10. Post S, Spiller HA, Chounthirath T, Smith GA: Kratom exposures reported to United States poison control centers: 2011-2017. *Clin Toxicol (Phila).* 2019, 57:847-54. [10.1080/15563650.2019.1569236](https://doi.org/10.1080/15563650.2019.1569236)
11. Ahmad J, Odin JA, Hayashi PH, et al.: Liver injury associated with kratom, a popular opioid-like product: experience from the U.S. drug induced liver injury network and a review of the literature. *Drug Alcohol Depend.* 2021, 218:108426. [10.1016/j.drugalcdep.2020.108426](https://doi.org/10.1016/j.drugalcdep.2020.108426)
12. Griffin OH, Webb ME: The scheduling of Kratom and selective use of data . *J Psychoactive Drugs.* 2018, 50:114-20. [10.1080/02791072.2017.1371363](https://doi.org/10.1080/02791072.2017.1371363)
13. Schimmel J, Dart RC: Kratom (*Mitragyna speciosa*) liver injury: a comprehensive review . *Drugs.* 2020, 80:263-8. [10.1007/s40265-019-01242-6](https://doi.org/10.1007/s40265-019-01242-6)
14. Osborne CS, Overstreet AN, Rockey DC, Schreiner AD: Drug-induced liver injury caused by Kratom use as an alternative pain treatment amid an ongoing opioid epidemic. *J Investig Med High Impact Case Rep.* 2019, 7:2324709619826167. [10.1177/2324709619826167](https://doi.org/10.1177/2324709619826167)
15. Dowell D, Haegerich TM, Chou R: CDC guideline for prescribing opioids for chronic pain - United States, 2016. *MMWR Recomm Rep.* 2016, 65:1-49. [10.15585/mmwr.rr6501e1](https://doi.org/10.15585/mmwr.rr6501e1)
16. Smith KE, Lawson T: Prevalence and motivations for Kratom use in a sample of substance users enrolled in a residential treatment program. *Drug Alcohol Depend.* 2017, 180:340-8. [10.1016/j.drugalcdep.2017.08.034](https://doi.org/10.1016/j.drugalcdep.2017.08.034)
17. Shahbaz O, Mahajan S, Lewis JH: Highlights of drug - and herb- induced liver injury in the literature from 2016: how best to translate new information into clinical practice?. *Expert Opin Drug Metab Toxicol.* 2017, 13:935-51. [10.1080/17425255.2017.1362391](https://doi.org/10.1080/17425255.2017.1362391)