

# Preliminary Characterization of Commercial Kratom Extract Products

NCT06640569

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**Status** RECRUITING  
**Sponsor** Johns Hopkins University  
**Enrollment** 16 participants

## Key Eligibility Criteria

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### Inclusion (5)

- \) >21 years old;
- \) Experienced kratom extract product consumer of one of the leading US brands listed in the study protocol;
- \) Reports a specific kratom extract product brand and specific dose amount (or range of typical dose amount) on the online screener;
- \) English language proficient;
- \) Willingness to provide requested samples of the kratom extract product currently being taken.

### Exclusion (11)

- \) Reports any acute adverse or unexpected or otherwise sudden health event related to their typical kratom product dose that occurred within 30 days of screening;
- \) Having ever sought medical attention for an acute adverse health event as a result of taking any kratom product;
- \) Cannot or will not provide their kratom extract product samples in the form of an unopened package that is clearly labeled with at least the product and vendor name;
- \) The kratom extract product used by the participant has any semi-synthetic or fully synthetic ingredient listed or is known by the study team to have such an ingredient;
- \) Currently breastfeeding or pregnant;
- ... and 6 more (see full listing online)

## Locations (2 total)

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Johns Hopkins University Behavioral Pharmacology Research Unit, Baltimore, Maryland, United States  
Johns Hopkins University, Baltimore, Maryland, United States