PUBLIC HEALTH AND SAFETY BULLETIN

November 17, 2021

Notification of Marijuana Product Recall

The Marijuana Regulatory Agency (MRA) has identified inaccurate and/or unreliable results of products tested by safety compliance facilities Viridis North, LLC and Viridis Laboratories, LLC.

In the interest of public health and safety, the MRA is issuing this health and safety advisory bulletin for **all** marijuana products tested by Viridis Laboratories, LLC (license numbers SC-000009 and AU-SC-000113) and Viridis North, LLC (license numbers SC-000014 and AU-SC-000103) **except** for inhalable marijuana concentrate products such as:

- Vape carts.
- Live resin.
- Distillate.
- Any other cannabis concentrate created through residual solvent extractions.

The marijuana products impacted have a test date between August 10, 2021 and November 16, 2021. All marijuana product labels are required to list the name and license number of the safety compliance facility that conducted the testing and date the product was tested.

Note: An MRA investigation is still on-going.

Consumers who have marijuana products in their possession that meet the recall criteria may return the products to the marijuana sales location where they were purchased for proper disposal. Consumers with weakened immune systems or lung disease are at the highest risk for health-related incidents such as aspergillosis, which can impact lung function, if these potentially harmful products are consumed.

Consumers who have experienced adverse reactions after using these products should report their symptoms and product use to their physician. Consumers are requested to report any adverse product reactions to the MRA via email: MRA-Enforcement@michigan.gov or via phone: 517-284-8599.

Marijuana sales locations that sold product covered by this bulletin must display this recall notice on the sales floor, visible to all customers, for 30 days from the date of this

notice. Marijuana sales locations that receive adverse product reactions from consumers should report the adverse product reactions to the agency at MRA-Enforcement@michigan.gov and document these reports in METRC.

Licensees with products remaining in their inventory that meet the recall criteria have the following options:

- Destroy the product and provide proof of destruction: MRA-Compliance@michigan.gov.
- Have the product retested for the microbials compliance panel.
- Send the product back to the original licensee source so they can destroy or have the product retested as a larger batch.

Licensees that opt to have product sent back or retested will need to create new METRC packages with new METRC identification numbers prior to transferring or submitting the products for testing. Additional guidance can be provided to licensees who need assistance in creating these packages by reaching out to MRA-Compliance@michigan.gov.

Additional questions can be sent to the MRA's Operations Support Section: MRA-Compliance@michigan.gov.