LICENSING AND REGULATORY AFFAIRS
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PUBLIC HEALTH AND SAFETY BULLETIN

February 26, 2019

Notification of Multiple Marijuana Products Recall

The Bureau of Marijuana Regulation (BMR) is issuing a health and safety advisory recall due to the sale of marijuana which failed laboratory testing.

All marijuana product subject to recall in this notice was purchased by the licensed facility from a registered primary caregiver under a resolution by the Medical Marihuana Licensing Board passed on January 16, 2019.

The products were sold between 12/28/2018 and 1/23/2019 at Utopia Gardens. All affected medical marijuana has a label affixed to the container that, at a minimum, indicates the license number of the marijuana facility that obtained the marijuana product, as well as the production batch number assigned to the marijuana product.

This recall affects the following marijuana flower products sold from Utopia Gardens- License PC-000079 located at 6541 E. Lafayette, Detroit, MI:

Utopia Concentrates
1A405010000641000000075
1A405010000641000000041
Failed for chemical residue

Utopia Cartridges
1A405010000641000000066
Failed for chemical residue

Patients or caregivers who have these affected medical marijuana products in their possession should return them to Utopia Gardens for proper disposal. Utopia Gardens must notify patients or caregivers that purchased these medical marijuana products of the recall.

For more information about BMR, please visit www.michigan.gov/bmr
For more information about LARA, please visit www.michigan.gov/lara

This advisory bulletin does not constitute legal advice and is subject to change. Licensees are encouraged to seek legal counsel to ensure their operations comply with the Medical Marihuana Facilities Licensing Act and associated Administrative Rules.