LICENSING AND REGULATORY AFFAIRS
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PUBLIC HEALTH AND SAFETY BULLETIN

February 11, 2019

Notification of Multiple Marijuana Products Recall (Updated)

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 12/27/2018 and 12/30/2018 at HG Lansing. 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 batches of flower sold from HG Lansing - License PC-000159 located at 1116 E. Oakland Ave., Lansing MI 48906:

Citrix
1A4050100000F3D000000009
Failed for *E.coli* and *Salmonella*

Gelato
1A4050100000F3D000000010
Failed for chemical residue

Green Crack
1A4050100000F3D000000008
Failed for chemical residue and bile-tolerant gram-negative bacteria, *E.coli*, *Salmonella* and total coliforms

Oreoz
1A4050100000F3D000000023
Failed for chemical residue and *E.coli* and S*almonella*

Patients or caregivers who have these affected medical marijuana products in their possession should return them to HG Lansing for proper disposal. HG Lansing 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.