

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 <u>resolution</u> by the Medical Marihuana Licensing Board passed on January 16, 2019.

The products were sold between 12/12/2018 and 12/29/2018 at Compassionate Care By Design in Kalamazoo, Michigan. 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 Compassionate Care By Design License PC-000142 located at 401 N. Sage St, Kalamazoo, MI 49006.

All products failed for chemical residue:

Blueberry

1A4040100000515000000019

Critical Cali Skunk #1

1A4040100000515000000014 1A4040100000515000000020

IO Chunk D

1A4040100000515000000015 1A4040100000515000000021

Silver HazeAmnesia Lemon Kush
1A4040100000515000000016
1A40401000005150000000022

GSC Special Kush

GSCSpecial Kush
1A4040100000515000000017
1A4040100000515000000023

 Critical Kush
 Purple Punch

 1A4040100000515000000018
 1A4040100000515000000024

1A4040100000515000000018 1A4040100000515000000024

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



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