# PUBLIC HEALTH AND SAFETY BULLETIN

January 10, 2020

## Notification of Medical Marijuana Flower Recall

The Marijuana Regulatory Agency (MRA) is issuing this health and safety advisory bulletin due to marijuana flower which failed laboratory testing.

All affected medical marijuana is required to have a label affixed to the container that indicates the METRC number assigned to the marijuana product. Patients and caregivers should look for the production batch number associated with the product name or the individual package number which can be found under the name of the provisioning center at which the product was sold. This recall affects the following products sold at three provisioning centers in the state of Michigan between 10/14/2019 and 1/6/2020:

**Production batch:** 1A4050100002330000000009
**Product name:** Orange Burst – Buds
**Failed testing:** Chemical Residue (Paclobutrazol)

Pharmaco, Inc. (License # PC-000249)
3650 Patterson RD
Bay City MI 48706
**Individual Package # 1A4050100002330000000415**

Pharmaco, Inc. (License # PC-000261)
3557 Wilder RD
Bay City MI 48706
**Individual Package # 1A4050100002330000000416**

Pharmaco, Inc. (License # PC-000205)
20561 Dwyer ST
Detroit MI 48234
**Individual Package # 1A4050100002330000000041**

Patients or caregivers who have these affected medical marijuana products in their possession should return them to the provisioning center where they were purchased for proper disposal. Provisioning centers who carried the products must notify patients or caregivers that purchased these medical marijuana products of the recall.

Patients who have experienced symptoms after using these products should report their symptoms and product use to their physician.

Patients and caregivers are requested to report any adverse product reactions to the MRA via email: MRA-Enforcement@michigan.gov or via phone: 517-284-8599.