



# PUBLIC HEALTH AND SAFETY BULLETIN

February 7, 2020

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## Notification of Medical Marijuana Product Recall

The Marijuana Regulatory Agency (MRA) is issuing this health and safety advisory bulletin due to the presence of Vitamin E Acetate in several vape cartridges which failed safety compliance testing. These cartridges were manufactured prior to the Emergency Rules for marijuana products intended for inhalation which were filed on November 22, 2019.

All affected vape cartridges will have a label that indicates the license number of the marijuana facility that sold the marijuana product as well as tag number assigned to the product in the statewide monitoring system.

### **Fuel 420**

This recall affects the following marijuana products sold from Fuel 420 – License PC-000077 – located at 1255 Falahee RD Jackson, MI 49203:

**METRC # 1A4050100001AF5000000026** (Cart 510 .6G Wedding Cake - FUEL 420)  
Sold between April 3, 2019 and November 11, 2019

**METRC # 1A4050100001AF5000000032** (Carts 510 .6G Lime - FUEL 420)  
Sold between March 31, 2019 and November 22, 2019

**METRC # 1A4050100001AF5000000022** (Cart 510 .6G Skittlez - FUEL 420)  
Sold between March 31, 2019 and November 22, 2019

**METRC # 1A4050100001AF5000000031** (Carts 510 .6G Lemonade - FUEL 420)  
Sold between March 31, 2019 and November 22, 2019

### **Green House of Walled Lake**

This recall affects the following marijuana products sold from Green House of Walled Lake – License PC-000129 – located at 103 E. Walled Lake DR Walled Lake, MI 48390:

**METRC # 1A40401000006A5000001961** (True-Green Crack)  
Sold between July 10, 2019 and November 22, 2019

**METRC # 1A4050100000BB9000001101** (Motor City High | Green Crack | 1g Cart)  
Sold between January 28, 2020 and February 5, 2019



# PUBLIC HEALTH AND SAFETY BULLETIN

February 7, 2020

---

**METRC # 1A40401000006A5000001963** (True-Sandferando)  
Sold between July 20, 2019 and November 22, 2019

**METRC # 1A4050100000BB9000001103** (Motor City High | San Fernando Valley | 1g Cart)  
Sold between January 28, 2020 and February 5, 2019

**METRC # 1A40401000006A5000001962** (True-North Lights)  
Sold between July 18, 2019 and November 22, 2019

**METRC # 1A4050100000BB9000001102** (Motor City High | Northern Lights | 1g Cart)  
Sold between January 28, 2020 and February 5, 2019

## **Liv Wellness Center, LLC**

This recall affects the following marijuana products sold from Liv Wellness Center, LLC – License PC-000298 – located at 2625 Hilton Rd Suite 100 Ferndale, MI 48220:

**METRC # 1A40401000006A5000002323** (True-Green Crack)  
Sold between September 18, 2019 and November 06, 2019

**METRC # 1A40401000006A5000002325** (True-Sandferando)  
Sold between September 18, 2019 and November 21, 2019

**METRC # 1A40401000006A5000002326** (True-North Lights)  
Sold between September 13, 2019 and November 06, 2019

**METRC # 1A40401000006A5000002205** (True-CBD/THC)  
Sold between September 12, 2019 and November 19, 2019

## **The Green Mile Detroit**

This recall affects the following marijuana products sold from The Green Mile Detroit – License PC-000144 – located at 6650 Eight Mile RD Detroit, MI 48234:

**METRC # 1A40401000006A5000001527** (True-CBD/THC)  
Sold between June 8, 2019 and July 1, 2019



# PUBLIC HEALTH AND SAFETY BULLETIN

February 7, 2020

---

## **664 Vassar, LLC**

This recall affects the following marijuana products sold from 664 Vassar, LLC – License PC-000035 – located at 664 State RD Vassar, MI 48768:

**METRC # 1A40501000045ED000002619** (True-Green Crack)

Sold between January 10, 2020 and February 4, 2020

**METRC # 1A40501000045ED000002609** (True-Sandferando)

Sold between January 10, 2020 and February 5, 2020

**METRC # 1A40501000045ED000002620** (True-North Lights)

Sold between January 10, 2020 and February 5, 2020

**METRC # 1A40501000045ED000002618** (True-CBD/THC)

Sold between January 10, 2020 and February 5, 2020

Patients or caregivers who have these affected vape cartridges in their possession should return them to the provisioning center where they purchased for proper disposal. The provisioning centers must notify patients or caregivers that purchased these vape cartridges 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](mailto:MRA-Enforcement@michigan.gov) or via phone: 517-284-8599.