Ohio Medical Marijuana Control Program

Vaporizing Devices Submission Guidance Document

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I. Document Purpose, Audience and Topics Covered

A. Document Purpose

This reference document is designed to provide a step by step guide for the medical marijuana industry to submit vaporizing devices to be reviewed by the MMCP to ensure the devices are in accordance with the relevant legal authority.

B. Intended Audience

The intended audience for this document includes individuals employed by or acting on behalf of dispensaries, processors, cultivators, or any person that is seeking to have their vaporizing device reviewed by the MMCP.

C. Topics Covered

The topics covered in this guidance document include:

- Submission and Status Process
- Minimum Submission Requirements
- Basic Non-Compliance Issues

II. Applicable Rules and Definitions

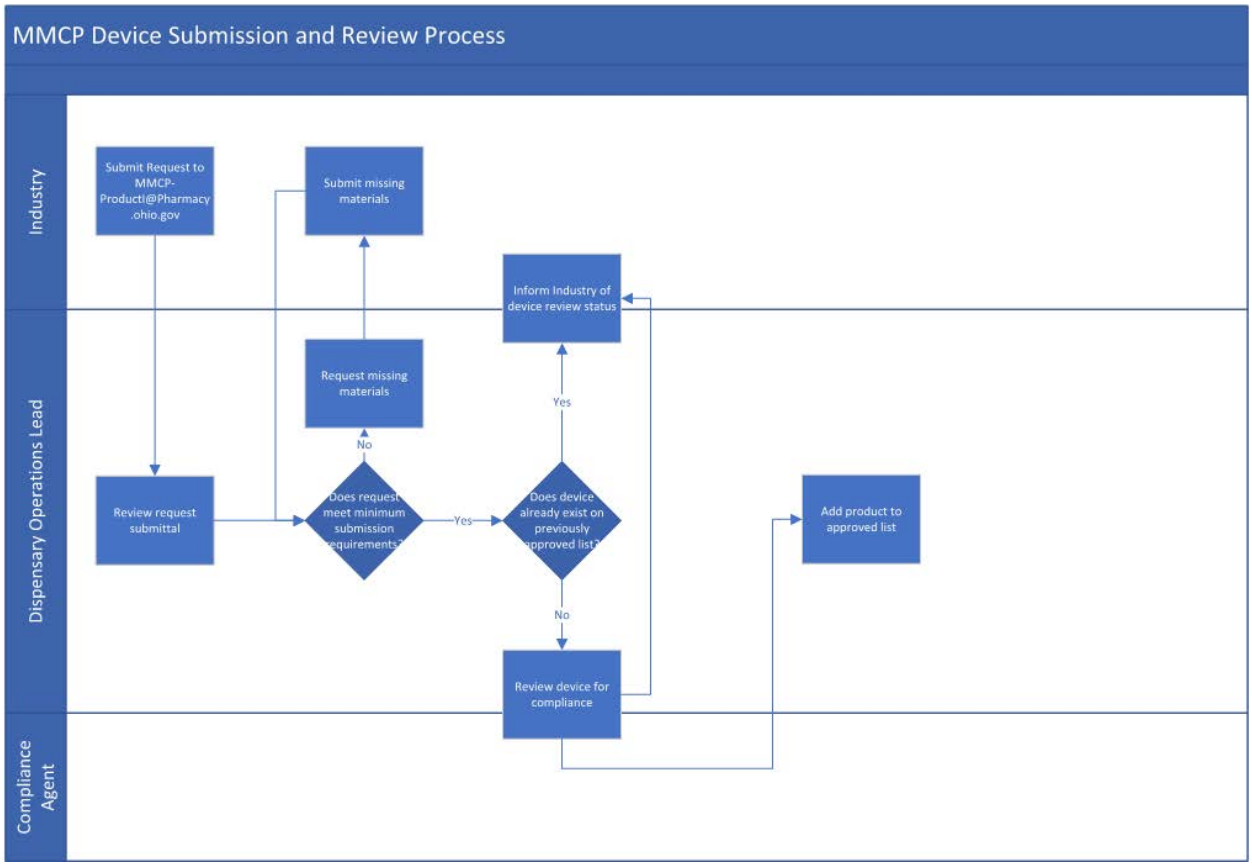
A. Relevant Legal Authority

For further information refer to the links below:

- Chapter 3796 of the Ohio Revised Code
- Ohio Administrative Code Rule 3796:8-2-01 - Authorized medical marijuana forms and methods of administration.

III. Submission Process and Status Process

- Send email to MMCP-ProductID@Pharmacy.ohio.gov with the Subject titled, "Vaporizing Device Review Request" with all of the required information for review (Appendix 1, A).
- After vaporizing device materials are submitted, the Board will review and decide as to whether the submission is compliant, non-compliant or more information is needed.
- After the determination has been made, a response including the approval status will be sent to the email containing the materials related to the vaporizing device.


MMCP Device Submission and Review Process
Industry
Dispensary Operations Lead
Compliance Agent
Submit Request to MMCP-Producti@Pharmacy.ohio.gov
Submit missing materials
Inform Industry of device review status
Request missing materials
Yes
No
Review request submittal
Does request meet minimum submission requirements?
Yes
Does device already exist on previously approved list?
Add product to approved list
No
Review device for compliance

IV. Appendix – Vaporizing Checklist

A. Minimum Submission Requirements

Includes all required materials submitted for review:

- ☐ The manufacturer name of the vaporizing device submitted for review
- ☐ The model name of the vaporizing device submitted for review
- ☐ A description of the device that includes the following:
  - Mechanics and functionality of the device
  - How the vaporizing device meets the all the regulatory requirements
  - Specify that the vaporizing device will or will not be sold with medical marijuana products
  - Specify the vaporizing device is intended for plant material or metered oil or solid oil.
- ☐ Diagrams and /or videos of the vaporizing device
- ☐ Any other useful information pertaining to the review of the vaporizing device

B. Basic Non-Compliance

- ☐ The vaporizing device must not place medical marijuana in direct contact with the device's heating element. This includes an extract of medical marijuana.
- ☐ The device must be capable of meter dosing the product - If the device does not include this functionality, then the medical marijuana product will need to be sold pre-dosed without the device.