2.4.2022 Email to Dispensaries

Hello,

The Department conducted a statewide review of all vaporization products containing added ingredients and has determined that certain vaporization products containing added ingredients, such as externally sourced flavorings or terpenes, have not been approved for inhalation by the United States Food and Drug Administration. 35 P.S. § 10231.702(a)(5), 28 Pa. Code § 1151.27(f).

The affected grower/processors have just been notified that these products meet the conditions for recall under 28 Pa. Code § 1151.42(c)(1); accordingly, mandatory recall procedures must be implemented. 28 Pa. Code § 1151.42(c). Certain vaporization products will no longer be available for dispensing to patients or caregivers. The list of affected products is posted on our website at www.medicalmarijuana.pa.gov.

You must return all recalled products to the grower/processor for proper disposal of these products in accordance with 28 Pa. Code § 1161.38(c). **You must provide proof of the return of all recalled products.** A manifest is acceptable as proof of return and should be emailed to RA-DHMMRCompliance@pa.gov. Failure to comply will result in the Department acting to impose sanctions against you under 28 Pa. Code § 1141.47.

You may appeal this action to the Secretary of Health in writing **within 30 days of the date of emailing** in accordance with 28 Pa. Code Chapter 1230 (relating to practice and procedure – temporary regulations).

If you have questions about specific products, please contact the grower/processor.

Thank you for your commitment to keeping patients safe.

Sunny