Virginia Department of Behavioral Health & Developmental Services

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Health Alert: Drug Recall

High Priority

The Food and Drug Administration has sent out an alert as follows:

"C.O. Truxton, Inc. Issues Voluntary Nationwide Recall of **Amitriptyline HCL Tablets, USP 50mg and Phenobarbital Tablets, USP 15mg, 30mg, 60mg, 100mg** Due to Potential Label Mix-Up"

"Bellmawr, New Jersey, C.O. Truxton, Inc. is expanding their 04/21/2017 voluntary recall, as a precaution to include the following C.O. Truxton, Inc. products, registered NDC numbers and corresponding lot numbers, to the consumer/user level. C.O. Truxton has not received any complaints for the products listed below. however, due to the initial recall resulting from a label mix-up error, out of an abundance of caution, we are recalling all products that were repackaged into a Truxton Incorporated label."

Please go to their website below for further information: **C.O. Truxton, Inc. Issues Voluntary Nationwide Recall of Amitriptyline HCL Tablets, USP 50mg and Phenobarbital Tablets, USP 15mg, 30mg, 60mg, 100mg Due to Potential Label Mix-Up**

If you or a person you are providing medications too is taking either **Amitriptyline** or **Phenobarbital**; please call your pharmacist or the prescriber to discuss **immediately**.

If you still have questions, please contact susan.rudolph@dbhds.virginia.gov