FROM THE KELLER ARMY COMMUNITY HOSPITAL PHARMACY: The FDA announced on April 30, 2021, that Acella Pharmaceuticals, LLC is voluntarily recalling certain lots of NP Thyroid® (Thyroid Tablets, USP) to the consumer level. On May 24, 2021, FDA classified this recall as a Class I drug recall.

The products are being recalled because routine testing has found these lots to be sub potent. The product contains less than 90% of the labeled amount of liothyronine (T3) and/or levothyroxine (T4).

The Keller Pharmacy has identified and notified patients who receive NP thyroid (from Keller) to inform them of the recall.

The pharmacy is still able to obtain non-recalled NP Thyroid, and does not anticipate any break in the ability to provide this medication to patients.

If any patients have questions or concerns, please contact the pharmacy at (845) 938-2271.

FDA Drug Recall Classes	
Class I	The most serious of recalls; ongoing product use may result in serious health threat or death.
Class II	Moderate severity concern; ongoing product use may pose serious adverse events or irreversible consequences.
Class III	Lowest severity concern; ongoing product use unlikely to cause adverse health threat; however, a marginal chance of injury may exist, so the product is being recalled.