Losartan Potassium – Legacy Pharmaceutical Packaging, LLC

On March 25, 2019 the U.S. Food and Drug Administration (FDA) announced that Legacy Pharmaceutical Packaging, LLC is recalling 40 repackaged lots of Losartan Tablets, USP 25mg, 50mg and 100mg. This recall was prompted due to Camber Pharmaceuticals, Inc. issuing a Voluntary Nationwide recall due to the detection of trace amounts of N-Nitroso N-Methyl 4-amino butyric acid (NMBA) a possible process impurity or contaminant in an active pharmaceutical ingredient. NMBA is a potential human carcinogen.

Losartan Potassium is a prescription medication used to treat high blood pressure and congestive heart failure. Patients should contact their doctor for further guidance and potential change of treatment before they stop taking the product.

The identifying NDC #s associated with Legacy's products are as follows: Losartan 25mg 68645-0577-54; Losartan 50mg 68645-0578-54; Losartan 100mg 68645-0579-54.

Patients with questions regarding this recall can contact Camber Pharmaceutical's Med line at 1-866-495-1995 Monday-Friday 9:00 AM – 5:00 PM (EST). Patients should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this drug product.