Losartan Tablets, USP 25mg, 50mg, 100mg – Camber Pharmaceuticals

On February 28, 2019 the U.S. Food and Drug Administration (FDA) announced that Camber Pharmaceuticals is voluntarily recalling 87 lots of Losartan Tablets, USP 25mg, 50mg and 100mg. This product is being recalled 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 Camber's products are as follows: Losartan 25mg 31722-700-90, 31722-70-05, 31722-700-10; Losartan 50mg 31722-701-30, 31722-701-90, 31722-701-10; Losartan 100mg 31722-702-30, 31722-702-90, and 31722-702-10.

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.