

## Drug Recall

**DATE OF RECALL:** June 28, 2022

**DRUG NAME:** Losartan potassium & Hydrochlorothiazide Tablets

**RECALLING FIRM:** Macleods Pharmaceuticals Limited

**REASON FOR RECALL:** This recall was issued due to CGMP Deviations- AZIDO Impurity levels observed to be above acceptable limits.

**FDA LINK:**

**OTHER DETAILS:** N/A

**RECALLED PRODUCT:**

Product Description	NDC Number	Affected Lots	Expiration Date
Losartan potassium & Hydrochlorothiazide Tablets 100 mg/25 mg, 30 count	33342-0052-07	BLM2114A	07/2023
Losartan potassium & Hydrochlorothiazide Tablets 100 mg/25 mg, 90 count	33342-0052-10	BLM2114A	07/2023
Losartan potassium & Hydrochlorothiazide Tablets 100 mg/25 mg, 1000 count	33342-0052-44	BLM2114A	07/2023