



Drug Recall

DATE OF RECALL: April 15, 2020

DRUG NAME: Losartan Potassium 50mg & 100mg Tablets

RECALLING FIRM: Avet Pharmaceuticals, Inc.

REASON FOR RECALL: This recall is being issued due to CGMP Deviations. The FDA lab confirmed the presence of an impurity, N-Methylnitrosobutyric acid (NMBA) in the finished product above the interim acceptable daily intake level.

FDA LINK: N/A

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Affected Lots/Expire Date
Losartan Potassium 50 mg Tablets, 30 Count	23155-0645-03	CL018008A exp. 03/2020 CL018008B exp. 03/2020 CL018009A exp. 03/2020
Losartan Potassium 50 mg Tablets, 90 Count	23155-0645-09	CL018008A exp. 03/2020 CL018008B exp. 03/2020 CL018009A exp. 03/2020
Losartan Potassium 50 mg Tablets, 1000 Count	23155-0645-10	CL018008A exp. 03/2020 CL018008B exp. 03/2020 CL018009A exp. 03/2020
Losartan Potassium 100 mg Tablets, 30 Count	23155-0646-03	CL018022B exp. 04/2020
Losartan Potassium 100 mg Tablets, 90 Count	23155-0646-09	CL018010A exp. 03/2020 CL018017B exp. 04/2020 CL018018A exp. 04/2020
Losartan Potassium 100 mg Tablets, 1000 Count	23155-0646-10	CL018011A exp. 03/2020 CL018012A exp. 03/2020 CL018013A exp. 04/2020 CL018015A exp. 04/2020 CL018016A exp. 04/2020 CL018017A exp. 04/2020