The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Description	Lot # and Exp Date	NDC	UPC
	Lot # and Exp Date H102287 03/31/23; H000556 11/30/22; H903582 10/31/22; H102286 03/31/23; H101821 03/31/23; H101496 03/31/23; H101479 03/31/23; H101078 03/31/23; H100901 03/31/23; H100220 02/28/23; H003198 02/28/23; H003045 12/31/22; H001062 11/30/22; H003044 02/28/23; H000557 11/30/22; H001431 11/30/22; H001706 12/31/22; H001707 12/31/22;	NDC 68180037809	
	H002312 01/31/23; H002341 11/30/22; H002391 12/31/22; H002517 01/31/23; H002841 01/31/23; H001061 11/30/22		
LOSARTAN POT TAB 25MG LUP 90	H001940 12/31/22; H003080 01/31/23; H101989 03/31/23; H101789 03/31/23; H101285 02/28/23; H101282 02/28/23; H100642 02/28/23; H100109 02/28/23; H002955 01/31/23; H002786 01/31/23; H000523 01/31/23; H002388 01/31/23; H001714 12/31/22; H001333 11/30/22; H001189 11/30/22; H002389 01/31/23	68180037609	36818037609

Description	Lot # and Exp Date	NDC	UPC
LOSARTAN POT TAB 50MG LUP 90	H001599 12/31/22; H000605 01/31/23; H001063 11/30/22; H001188 11/30/22; H001401 11/30/22; H001875 12/31/22; H001455 11/30/22; H002126 12/31/22; H002642 01/31/23; H002838 01/31/23; H100148 02/28/23; H101495 03/31/23;	68180037709	36818037709
LOSARTAN POT TB 25MG LUPI1000	H102043 03/31/23 H100110 02/28/23; H003122 02/28/23; H003121 02/28/23; H003079 01/31/23; H002958 01/31/23; H100111 02/28/23; H002787 01/31/23; H100644 03/31/23; H002957 01/31/23; H100643 02/28/23; H002489 01/31/23; H100643 02/28/23; H002003 12/31/22; H100869 03/31/23; H101283 02/28/23; H101284 02/28/23; H101990 03/31/23; H101991 03/31/23; H100147 02/28/23; H001715 12/31/22; H000847 11/30/22; H000848 11/30/22; H001059 11/30/22; H001058 11/30/22; H001059 11/30/22; H00190 11/30/22; H001191 11/30/22; H001486 01/31/23; H001275 11/30/22; H001717 12/31/22; H001716 12/31/22; H001941 12/31/22; H001718 12/31/22; H001941 12/31/22; H002002 12/31/22; H001941 12/31/22; H002390 01/31/23; H002487 01/31/23; H001192 11/30/22	68180037603	36818037603
Losartan Potassium Tablets USP, 50mg	See vendor notice for affected lots and expiration dates.	68180037703	
Losartan Potassium Tablets USP, 100mg	See vendor notice for affected lots and expiration dates.	68180037803	

Lupin is voluntarily recalling the above items/lots due to results observed above acceptable limits in Losartan Azido Tetrazole impurity in nine analysed lots of Losartan Potassium Tablets USP. This recall is to the retail level. Affected product started shipping March 17, 2020.