

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.

Avet Pharmaceuticals Inc. ("Avet") located in East Brunswick, New Jersey, is initiating a recall of 9 lots of Desipramine Hydrochloride Tablets USP, 10 mg, 25 mg, 50 mg, 75 mg, 100 mg, and 150 mg. A full list of impacted lot numbers is provided in the below table. This recall is being conducted in coordination with the FDA to the Retail level. The drug product is manufactured by USV Private Ltd. (Daman, India) and is distributed by Avet. This recall is being initiated due to N-Nitroso Desipramine impurity exceeding the permissible acceptable daily intake level.

Please note that there are no adverse events or any complaints that were reported to Avet for the identified lots of Desipramine Hydrochloride Tablets USP.

Product Name	Dosage/ Package	NDC Number	Lot Number	Expiry
Desipramine Hydrochloride Tablets USP	10mg, 100s	23155- 578-01	18036908	Sept- 26
Desipramine Hydrochloride Tablets USP	25mg, 100s	23155- 579-01	18035876	Dec- 25
Desipramine Hydrochloride Tablets USP	25mg, 100s	23155- 579-01	18036909	Sept- 26
Desipramine Hydrochloride Tablets USP	50mg, 100s	23155- 580-01	18036713	Aug- 26
Desipramine Hydrochloride Tablets USP	75mg, 100s	23155- 581-01	18036662	Jul-26
Desipramine Hydrochloride Tablets USP	75mg, 100s	23155- 581-01	18037649	Mar- 27

Desipramine Hydrochloride Tablets USP	100mg, 100s	23155- 582-01	18035574	Oct- 25
Desipramine Hydrochloride Tablets USP	100mg, 100s	23155- 582-01	18036758	Aug- 26
Desipramine Hydrochloride Tablets USP	150mg, 50s	23155- 583-25	18035739	Nov- 25