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.

This drug product was manufactured by Ultra Tab Laboratories Inc. (Hightstown, N.Y.) for Par Pharmaceuticals Inc. This recall by ANI Pharmaceuticals has been initiated due to potential cross-contamination with other drug substances during the manufacturing process. ANI Pharmaceuticals is the current application holder for the affected drug product. This recall does not involve product manufactured or distributed by ANI Pharmaceuticals or its subsidiary Novitium Pharma.

The affected products were distributed within the following dates:

- Alprazolam Tab lets, USP 02/21/2019 to 10/03/2019
- Pyrazinamide Tablets, USP 04/04/2019 to 10/01/2019

Product & NDC Number	Lot Number
Alprazolam Tablets, USP 0.25 mg	
(NDC Numbers)	19C003A, 03 /2022; 19C004B, 03/2022;
67253-900-10, 100 TABLET in 1 BOTTLE,	
67253-900-50, 500 TABLET in 1 BOTTLE,	19C048C, 03/2022; 19G002A , 07/2022
67253-900-11, 1000 TABLET in 1 BOTTLE	
	19A087B, 02/2022; 19A088B, 02/2022;
Alprazolam Tablets, USP 0.5 mg	19A089B, 02/2022; 19A090B, 02/2022;
(NDC Numbers)	19B020C, 02/2022; 19B021C, 02/2022;
67253-901-10, 100 TABLET in 1 BOTTLE,	19B027C, 02/2022; 19B028C , 02/2022;
67253-901-50, 500 TABLET in 1 BOTTLE ,	19B029A, 02/2022; 19E056C, 05/2022;
67253-901-11, 1000 TABLET in 1 BOTTLE	19A086B, 02/2022; 19A091B, 02/2022; 19B019B, 02/2022; 19D021A, 04/2022; 19E057C, 05/2022; 19E059C, 06/2022; 19G072C, 07/2022

Product & NDC Number	Lot Number
Alprazolam Tablets, USP 1.0 mg	19A102B, 02/2022; 19B081A, 02/2022;
(NDC Numbers)	19B082C, 03/2022; 19B083C, 03/2022;
67253-902-10, 100 TABLET in 1 BOTTLE,	19D067B, 04/2022; 19D068B, 04/2022;
67253-902-50, 500 TABLET in 1 BOTTLE,	19D069C, 05/2022; 19D070C, 05/2022; 19E088A, 05/2022; 19E089A, 05/2022;
67253-902-11, 1000 TABLET in 1 BOTTLE	19045C, 06/2022; 19F046C, 06/2022
Alprazolam Tablets, USP 2.0 mg	
(NDC Numbers)	19C002A, 03/2022; 19C100B, 04/2022; 19E001B, 05/2022; 19E002B, 05/2022;
67253-903-50, 500 TABLET in 1 BOTTLE,	1520015, 05, 2022, 1520025, 05, 2022,
	19E012A, 05/2022; 19E013A, 05/2022
67253-903-10, 100 TABLET in 1 BOTTLE	
Pyrazinamide Tablets, USP 500mg	
(NDC Numbers)	19B064A, 03/2022
67253-660-10, 100 TABLET in 1 BOTTLE	