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 # / Exp Date	NDC	UPC
METFORM ER TAB 500MG APX 100	ND1973 02/28/21; NE5800 04/30/21; MZ8147 10/31/20; MZ8148 10/31/20; MZ8149 10/31/20; NC3603 04/30/21; NC3604 11/30/20; NC3605 04/30/21; MZ8145 11/30/20; NC3607 11/30/20; MZ8144 10/31/20; ND1974 02/28/21; ND7849 04/30/21; ND7850 02/28/21; ND7851 02/28/21; ND7852 02/28/21; ND7853 04/30/21; NE5799 04/30/21; MN6854 06/30/20; NC3606 04/30/21; MW2800 07/31/20; MP8932 06/30/20; MP8933 06/30/20; MP8934 06/30/20; MR6717 06/30/20; MR6718 06/30/20; MV6913 07/31/20; MV6914 07/31/20; MZ8146 02/28/21; MW2799 07/31/20; NE5801 04/30/21; MW2801 07/31/20; MZ1936 10/31/20; MZ1937 09/30/20; MZ1938 10/31/20; MZ1939 10/31/20; MZ1940 10/31/20; MZ1941 11/30/20; MZ8143 11/30/20; MV6915 07/31/20	60505026001	36050502601
METFORM ER TB 500MG APXDOD 100	ND1973 02/28/21; MN6854 06/30/20; MZ8147 10/31/20; MZ8148 10/31/20; MZ8149 10/31/20; NC3603 04/30/21; NC3604 11/30/20; NC3605 04/30/21; MZ8145 11/30/20; NC3607 11/30/20; MZ8144 10/31/20; ND1974 02/28/21; ND7849 04/30/21; ND7850 02/28/21; ND7851 02/28/21; ND7852 02/28/21; ND7853 04/30/21; NE5799 04/30/21; NE5800 04/30/21; NC3606 04/30/21; MW2800 07/31/20; MP8932 06/30/20; MP8933 06/30/20; MP8934 06/30/20; MR6717 06/30/20; MR6718 06/30/20;	60505026001	36050502601

Description	Lot # / Exp Date	NDC	UPC
	MV6913 07/31/20; MV6914 07/31/20; MZ8146 02/28/21; MW2799 07/31/20; NE5801 04/30/21; MW2801 07/31/20; MZ1936 10/31/20; MZ1937 09/30/20; MZ1938 10/31/20; MZ1939 10/31/20; MZ1940 10/31/20; MZ1941 11/30/20; MZ8143 11/30/20; MV6915 07/31/20		

Apotex is recalling the above items/lots due to showed results for NDMA levels in excess of the daily acceptable intake limit. This recall is to the retail level. Affected product started shipping June 29, 2016.