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.

Dr. Reddy's has initiated a recall of 11 following lots of various unit-dose products which were packaged for institutional use but may have been dispensed to patients through non-institutional pharmacies. Dr. Reddy's sold the recalled medications to wholesalers starting in 2018. Ultimately, these medications may have been sold to consumers at retail pharmacies in the United States. The packaging of the products is not child resistant and can pose a risk of poisoning if the contents are swallowed by young children.

NDC	**Product**	**Lot Number(s)**
43598-344-31	Imatinib Mesylate 100 mg (3x10s) Blisters	H2000138
43598-345-31	Imatinib Mesylate 400 mg (3x10s) Blisters	H2000127
43598-292-66	Pregabalin Capsules 50 mg (5x10s) Blisters	T900876
43598-293-66	Pregabalin Capsules 75 mg (5x10s) Blisters	T901021
43598-294-66	Pregabalin Capsules 100 mg (5x10s) Blisters	T901022
43598-295-66	Pregabalin Capsules 150 mg (5x10s) Blisters	T901023
55111-789-11	Sevelamer Carbonate Tablets 800 mg (25x4s) Blisters	T000009, T801003, T900221
43598-575-31	Tadalafil Tablets, USP 5 mg (3x10s) Blisters	T000376
43598-573-31	Tadalafil Tablets, USP 20 mg (3x10s) Blisters	T000425

Distribution Dates: 2018 – 2020