The Board of Pharmacy has received notice of the following product recall:

This letter is to inform you that SterRx LLC has issued a Market Withdrawal for the following products:

Product	Batch/Lot Number	Expiration Date	NDC Number	Distribution Dates
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19355/AQK	16Jul20	70324-951-01	25 Sep 19 - 01 Oct 19
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19357/AQL	17Jul20	70324-951-01	01 Oct 19 - 21 Oct 19
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19372/AQW	19 Aug 20	70324-951-01	22 Oct 19 - 05 Nov 19
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19390/ARR	04 Sep 20	70324-951-01	13 Nov 19 - 12 Dec 19
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19414/ASO	25 Sep 20	70324-951-01	12 Dec 19 - 26 Dec 19
125mg Diltiazem HCl in 5% Dextrose Injection, 125ml in 100ml Conventional IV Bag	S19415/ASP	29 Sep 20	70324-951-01	08 Jan 20 - 21 Jan 20
125mg Diltiazem HCl in 0.7% Sodium Chloride Injection, 125ml in 100ml Conventional IV Bag	S19329/APZ	14 Aug 20	70324-976-01	02 Oct 19 -16 Oct 19
125mg Diltiazem HCl in 0.7% Sodium Chloride Injection, 125ml in 100ml Conventional IV Bag	S19333/AQB	15 Aug 20	70324-976-01	22 Oct 19 - 19 Nov 19

Product	Batch/Lot Number	Expiration Date	NDC Number	Distribution Dates
125mg Diltiazem HCl in 0.7% Sodium Chloride Injection, 125ml in 100ml Conventional IV Bag	S19351/AQI	04 Sep 20	70324-976-01	16 Oct 19 - 10 Dec 19
125mg Diltiazem HCl in 0.7% Sodium Chloride Injection, 125ml in 100ml Conventional IV Bag	S19391/ARS	29 Oct 20	70324-976-01	10 Dec 19-17 Jan 20
125mg Diltiazem HCl in 0.7% Sodium Chloride Injection, 125ml in 100ml Conventional IV Bag	S19392/ART	30 Oct 20	70324-976-01	17 Jan 20- 23 Jan 20

This market withdrawal is being made with the knowledge of the Food and Drug Administration, and has been initiated due to the trending of product potency detected during testing of stability samples for the products and lots listed above. The potency has not yet tested out-of-specification, however the potency is projected to be out-of-specification before the 309 day (125mg Diltiazem in 5% Dextrose) and 360 day (125mg Diltiazem in 0.7% Sodium Chloride) expiration date of the products. A decreased potency has the potential to decrease effectiveness of the product in patients. To date, SterRx, LLC has not received reports of any adverse events associated with this issue.

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