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.

Biogen is recalling:

VUMERITY® 231 mg bottles lots #101801, #101799, #102826, and #102362

Lot Number	NDC Codes	Expiry
101801 (106 ct)	64406-020-01	31-May-21
101799 (120 ct)	64406-020-03	31-May-21
102826 (106 ct)	64406-020-01	30-Jun-21
102362 (120 ct)	64406-020-03	30-Jun-21

This recall has been initiated due to out of specification results observed during the buffer stage dissolution testing on Vumerity® 231mg drug product for all stability conditions at 25°C/60% (long term conditions) 30°C/65% (intermediate conditions) and 40°C/75% (accelerated conditions) on three (3) manufactured bulk product lots. All other key product quality indicators are within specification and demonstrate no atypical results.

This recall is to the retail level. Biogen began shipping this product on 21 November 2019.