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.

Regeneron has initiated a recall for EYLEA® (aflibercept) injection 2 mg Prefilled Syringe NDC 61755-005-01. This recall has been initiated due to reported incidents of syringe center barrel breakage that may happen upon receipt, preparation or dose delivery. There have been no increased incidence of Adverse Events reported for the subject six (6) lots.

This recall does NOT include any other forms (vial) or doses (8 mg or EYLEA® HD) of aflibercept.

Although glass breakage is an inherent risk with any glass container, this particular break occurred at a frequency level (i.e., in this case 0.035%) that we determined, in coordination with the FDA, warrants this precautionary recall action due to potential risks or hazards associated in the event breakage were to occur while handling or dosing the product. This limited recall is not anticipated to have any adverse impact to product supply.

This recall includes the following finished labeled drug product lots. Additionally, the dates of distribution (from Regeneron) are also included in the below chart.

Lot	**Distributed** Units	**First Date of** Distribution	Last **Date** of **Distribution**	**Expiration Date**
8231500321	39,240	30-Aug-23	6-Sep-23	Oct-24
8231500335	20,592	20-Sep-23	20-Sep-23	Jan-25
8231500333	50,496	25-Sep-23	28-Sep-23	Jan-25
8231500334	50,249	31-Jul-23	2-Oct-23	Jan-25
8231500339	50,125	28-Sep-23	31-Oct-23	Jan-25
8231500347	40,802	27-Sep-23	31-Oct-23	Jan-25