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.

Armas Pharmaceuticals is recalling the below mentioned batches of Docetaxel Injection USP 160mg/8mL (20mg/mL) and Azacitidine for Injection 100mg/ Vial.

Product Name	**Batch No.**	**Manufacturing Date**	**Expiration Date**	**NDC Number**
Docetaxel Injection USP 160mg/8mL (20mg/mL)	7S10185A	Nov-19	Oct-21	72485-216-08
Azacitidine for Injection 100mg/vial	7S10115A	Aug-19	Jul-21	72485-201-01
Azacitidine for Injection 100mg/vial	7S10256A	Dec-19	Nov-21	72485-201-01
Azacitidine for Injection 100mg/vial	7S10182B	Oct-19	Sep-21	72485-201-01
Azacitidine for Injection 100mg/vial	7S10255A	Dec-19	Nov-21	72485-201-01
Azacitidine for Injection 100mg/vial	7T10040A	Feb-20	Jan-22	72485-201-01
Azacitidine for Injection 100mg/vial	7T10028A	Jan-20	Dec-21	72485-201-01

This recall has been initiated due to cGMP deviation originating from the warning letter issued to Manufacture Shilpa Medicare Limited by U.S Food and Drug Administration for the batches mentioned in above table. Use of these product batches may not have potential health hazard; however, the recall of above listed batches is to the **RETAIL level**. Product began shipping on November 7, 2019.