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.

Teva Pharmaceuticals USA Inc. is recalling three lots of AirDuo® Digihaler® (fluticasone propionate and salmeterol) Inhalation Powder to the RETAIL LEVEL. These lots were distributed under the labels for Teva Pharmaceuticals USA Inc.

NDC	**Lot**	**Exp. Date**	**Product Name**	**Size**	**Shipping Dates**
59310-111-06	AFR16A	01/2022	AirDuo® Digihaler® 55/14 (fluticasone propionate 55 mcg and salmeterol 14 mcg) Inhalation Powder	60 METERED INHALATIONS	09/15/2020 - 03/03/2020
59310-129-06	AFR17A	01/2022	AirDuo® Digihaler® 113/14 (fluticasone propionate 113 mcg and salmeterol 14 mcg) Inhalation Powder	60METERED INHALATIONS	09/15/2020 - 03/05/2021
59310-136-06	AFR18A	11/2021	AirDuo® Digihaler® 232/14 (fluticasone propionate 232 mcg and salmeterol 14 mcg) Inhalation Powder	60 METERED INHALATIONS	09/15/2020 - 03/11/2021

This recall is being initiated because during routine stability testing of these lots, results for Salmeterol were below the approved specification limits. It is important to note that Fluticasone Propionate stability testing results are within approved specification limits. At the time of commercial release to the market, these three lots met all release requirements.