

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

Description	Lot #	Exp Date	NDC	UPC
EPTIFIB SDV 20MG/100ML AKOR1	091277A	09/30/2019	17478090290	31747890290
EPTIFIB SDV 20MG/10ML AKOR1	101097A	10/31/2019	17478090210	31747890210
	091307A	09/30/2019		
EPTIFIB SDV 75MG/100ML AKOR1	101107A	10/31/2019	17478090390	31747890390
	091377A	09/30/2019		
	091287A	09/30/2019		

Akorn is recalling the above items/lots due to out-of-specification impurities. During routine stability testing at 18-month samples resulted in 0.3% for D-aspartic acid. The specification is NMT 0.2%. Lot 101107A is also being recalled due to fill volume was out of specification at 94 mL (spec NLT 100 ml). This recall is to the retail level. Affected product started shipping December 15, 2017.

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.