

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

Description	Lot #	Exp Date	NDC	UPC
FENTAN FTV 0.05MG/ML HW 25X2ML	08134DK	02/01/2021	00409909422	30409909422
	08133DK	02/01/2021		

Hospira is recalling the above lots due to confirmed customer reports for vials with the potential for loose metal overseas crimp defects. This recall is to the retail/pharmacy level. Affected product started shipping October 2019.

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.