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

LEVEL OF NOTIFICATION: Consumer

SUPPLIER: Avkare

Description	Lot # /Exp Date	NDC	UPC
RANITID TAB 150MG AVK 1000	HH04918A 08/31/20; HG02319A 06/30/21; HG02419A 06/30/21; HG02619A 06/30/21; HH04518A 08/31/20; HH04618A 08/31/20; HL07518A 11/30/20; HH04818A 08/31/20; HE03319A 04/30/21; HK006918A 10/31/20; HK02718A 10/31/20; HK16617A 11/30/19; HL03917A 11/30/19; HL04017A 11/30/19; HA00419A 12/31/20; HH04718A 08/31/20; HC14018A 04/30/20; HA00519A 12/31/20; HA2719A 12/31/20; HA02819A 12/31/20; HB03518A 03/31/20; HB03618A 03/31/20; HC05019A 03/31/21; HE05419A 04/30/21; HC05911A 03/31/21; HE03419A 04/30/21; HC14118A 04/30/20; HC14218A 04/30/20; HC14318A 04/30/20; HC14418A 04/30/20; HC14518A 05/31/20; HM06017A 11/30/19; HC05119A 03/31/21; HM06117A 11/30/19; HL07418A 11/30/20;	42291072410	34229172410

Description	Lot # /Exp Date	NDC	UPC
RANITID TAB 150MG AVK 180	21570 03/01/20; 21571 03/01/20; 22190 03/31/20; 22192 05/31/20; 22497 05/31/20; 22620 05/31/20; 22999 09/30/20; 23000 09/30/20; 24158 03/31/21; 24159 04/30/21	42291072418	34229172418
RANITID TAB 150MG AVK 60	22657 03/31/20; 24157 04/30/21; 23001 09/30/20; 22193 03/31/20; 21680 03/01/20; 21241 03/01/20; 23002 09/30/20	42291072460	34229172460
RANITID TAB 300MG AVK 250	22247 06/30/20; 24289 01/31/21; 24199 01/31/21; 24198 01/31/21; 23244 11/30/20; 23214 09/30/20; 21528 02/01/20; 21527 02/01/20; 21309 02/01/20; 21307 02/01/20; 23243 09/30/20	42291072525	34229172525
RANITID TAB 300MG AVK 30	23776 01/31/21; 22291 06/30/20; 23215 09/30/20	42291072530	34229172530

Avkare is recalling the above item due to potential NDMA amounts present in the items above FDA levels. This recall is to the consumer level.

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.