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

Description: Ranitidine Tablets, USP 150mg
NDC: 53746-0253-10
Lot #s/Expiration:
HD03119A 3/31/2021;
HD03219A 3/31/2021
HE03119A 4/30/2021
HE03219A 4/30/2021

Description: Ranitidine Tablets, USP 150mg
NDC: 65162-0253-06
Lot #s/Expiration:
AR181690A 10/31/2020
AR181691A 10/31/2020
AR181806A 11/30/2020
AR190183A 1/31/2021
AR190184A 1/31/2021

Description: Ranitidine Tablets, USP 150mg
NDC: 65162-0253-10
Lot #s/Expiration:
AR180675A 4/30/2020
AR180868B 5/31/2020
AR190366B 2/28/2021

Description: Ranitidine Tablets, USP 150mg
NDC: 65162-0253-11
Lot #s/Expiration:
AR180559A 3/31/2020
AR180560A 3/31/2020
AR180594A 3/31/2020
AR180595A 3/31/2020
AR180829A 4/30/2020
AR180831A 5/31/2020
AR180832A 5/31/2020
AR180868A 5/31/2020
AR181807C 11/30/2020
AR190085A 12/31/2020
AR190086A 12/31/2020
AR190087A 12/31/2020
AR190124A 1/31/2021
AR190542B 3/31/2021
AR190609A 3/31/2021

AR190610A 3/31/2021

Description: Ranitidine Tablets, USP 150mg
NDC: 65162-0253-18
Lot #s/Expiration:
AR180483B 3/31/2020
AR181807B 11/30/2020
AR190008B 12/31/2020

Description: Ranitidine Tablets, USP 150mg
NDC: 65162-0253-50
Lot #s/Expiration:
AR180869A 5/31/2020
AR180870A 5/31/2020
AR180871A 5/31/2020
AR180872A 5/31/2020
AR180995A 5/31/2020
AR180996A 5/31/2020
AR180997A 6/30/2020
AR180998A 6/30/2020
AR181158A 7/31/2020
AR181159A 7/31/2020
AR181160A 7/31/2020
AR181161A 7/31/2020
AR181692A 10/31/2020
AR181693A 10/31/2020
AR181694A 10/31/2020
AR181709A 10/31/2020
AR181710A 11/30/2020
AR181711A 11/30/2020
AR181808A 11/30/2020
AR190004A 12/31/2020
AR190005A 12/31/2020
AR190006A 12/31/2020
AR190007A 12/31/2020
AR190008A 12/31/2020
AR190088A 12/31/2020
AR190089A 12/31/2020
AR190090A 12/31/2020
AR190121A 12/31/2020
AR190122A 12/31/2020
AR190123A 12/31/2020
AR190125B 1/31/2021
AR190181A 1/31/2021

AR190182A 1/31/2021
AR190364A 2/28/2021
AR190365A 2/28/2021
AR190366A 2/28/2021
AR190509A 3/31/2021
AR190510A 3/31/2021

Description: Ranitidine 300 mg Tablets
NDC: 65162-0254-03
Lot #s/Expiration:
AR180519A 3/31/2020
AR180615A 3/31/2020
AR181795A 11/30/2020
AR181921B 12/31/2020
AR190705A 4/30/2021

Description: Ranitidine 300 mg Tablets
NDC: 65162-0254-10
Lot #s/Expiration:
AR180613A 3/31/2020
AR181156A 7/31/2020
AR181157A 7/31/2020
AR190418B 2/28/2021

Description: Ranitidine 300 mg Tablets
NDC: 65162-0254-25
Lot #s/Expiration:
AR180638A 3/31/2020
AR180640A 4/30/2020
AR180641A 4/30/2020
AR181920A 12/31/2020
AR181921A 12/31/2020
AR190414B 2/28/2021
AR190415A 2/28/2021
AR190416A 2/28/2021
AR190417A 2/28/2021
AR190418A 2/28/2021
AR190543A 3/31/2021
AR190544A 3/31/2021
AR190545A 3/31/2021

Amneal Pharmaceuticals LLC is voluntarily on a precautionary basis recalling Ranitidine Tablets. USP. 150 mg and 300 mg to the consumer level. Ranitidine Tablets are being recalled because of potential N-Nitrosodimethylamine (NDMA) amounts above levels established by the FDA.

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.