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

Product: Amiodarone Hydrochloride Injection, USP 150mg/3mL
NDC: 55150-180-03
Batch: CAH180009
Expiration Date: Feb 2020

Product: Amiodarone Hydrochloride Injection, USP 450mg/9mL
NDC: 55150-181-09
Batch: CAH180001
Expiration Date: Jan 2020

Product: Amiodarone Hydrochloride Injection, USP 450mg/9mL
NDC: 55150-181-09
Batch: CAH180003
Expiration Date: Feb 2020

Product: Amiodarone Hydrochloride Injection, USP 450mg/9mL
NDC: 55150-181-09
Batch: CAH180011
Expiration Date: Jun 2020

Product: Amiodarone Hydrochloride Injection, USP 450mg/9mL
NDC: 55150-181-09
Batch: CAH180012
Expiration Date: Jun 2020

Product: Amiodarone Hydrochloride Injection, USP 900mg/18mL
NDC: 55150-182-18
Batch: CAH180013
Expiration Date: Jul2020

Product: Amiodarone Hydrochloride Injection, USP 900mg/18mL
NDC: 55150-182-18
Batch: CAH180014
Expiration Date: Jul2020

This recall has been initiated due to confirmed customer reports of the presence of visible particulate matter, identified as crystallized amiodarone, within several vials from the above listed lots. To date, AuroMedics has not received reports of any adverse events or identifiable safety concerns attributed· to the use of this product.

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.