Provider Newsflash
January 2022

Company Announcement/Recall – AuroMedics Pharma, LLC

Purpose of this communication:

We are writing to inform you that the FDA has posted a voluntary recall by AuroMedics Pharma, LLC of Polymyxin B for Injection USP, 500,000 Units/Vial, lot number CPB200013 due to the presence of particulate matter, identified as hair, being discovered in a vial within this lot.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/auromedics-pharma-llc-issues-voluntary-nationwide-recall-polymyxin-b-injection-usp-500000-unit-vial?utm_medium=email&utm_source=govdelivery
- Notify impacted patients and facilitate the repair, replacement and/or resolution of the recall, according to the guidelines issued by the manufacturer in the notification.
- Confirm if you have any CareCentrix patients affected by this recall and notify the CareCentrix Quality Department by faxing information to: 919-792-6718 Attn: Quality Department.

Thank you in advance for your cooperation and continued partnership.