carecentrix

Provider Newsflash
December, 2022

FDA Recall – Accord Healthcare Daptomycin Injection

Purpose of this communication:

- We are writing to inform you effective immediately the FDA has posted notice of a voluntary recall of a single lot of Daptomycin for Injection 500 mg/vial and Daptomycin for Injection 350 mg/vial product contained in cartons imprinted with lot #R2200232, Expiration date 1/2025 to the consumer/user level due to a product mix up. The recall was issued due to receipt of a product complaint report from a hospital pharmacy stating that vials labeled as "Daptomycin for Injection 500 mg/vial" were found in cartons labeled as "Daptomycin for Injection 350 mg/vial". The lot and expiration date printed on the outer carton and inner vials are the same for both the 500 mg/vial dose and the 350 mg/vial dose. Administration of the Daptomycin 500 mg/vial to the population most at risk which are children or patients with renal failure could lead to serious adverse health consequences.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/accord-healthcare-inc-issues-nationwide-voluntary-recall-daptomycin-injection-500-mgvial-and?utm_medium=email&utm_source=govdelivery
- Notify impacted patients and facilitate the replacement and/or resolution of the recall, according to the guidelines issued by the manufacturer in the FDA 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.