Provider Newsflash
September, 2021

FDA Recall

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has issued notice of a voluntary nationwide recall by IntegraDose Compounding Services, LLC of 9 lots of Cefazolin 2 gram in 20 ml syringe for Injection and 2 lots of Cefazolin 3 gram in 100 mL 0.9% sodium chloride bag for injection due to a lack of sterility assurance.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/integradose-compounding-services-llc-issues-voluntary-nationwide-recall-cefazolin-injection-products?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 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.