carecentrix

Provider Newsflash May 6, 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 issued by Hospira, Inc. of Lot DN9185 of their Sterile Water for Injection, USP, 100 ml single dose glass fliptop vial due to a confirmed customer report of a single vial with visible particulate.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/hospira-issues-voluntary-nationwide-recall-one-lot-sterile-water-injection-usp-due-potential?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 for your prompt attention and cooperation in this matter.