Provider Newsflash
August, 2021

FDA Recall

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has published a voluntary recall issued by Cardinal Health of all lots of their Monoject Flush Prefilled Saline Syringes (0.9% Sodium Chloride). The products have been found to reintroduce air into the syringe after the air has been expelled. This could result in the injection of air into blood vessels and create the potential for air embolism, which can cause serious adverse health outcomes or death.

The following SKU's have been recalled:

- 12 mL Syringe, 10 ml Saline Fill – 8881570121
- 12 mL Syringe, 3 mL Saline Fill – 8881570123
- 12 mL Syringe, 5 mL Saline Fill – 8881570215

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/cardinal-health-issues-nationwide-recall-select-monojecttm-flush-prefilled-saline-syringes?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.