carecentrix

Provider Newsflash Sept 15, 2020

FDA RECALL

Purpose of this communication:

We are writing to inform you that effective immediately the FDA has issued a Class I recall of BD CareFusion 303, Inc.'s Alaris System PC unit and PC unit front case keypad replacement kits manufactured from April 7, 2017 to June 15, 2020 and distributed from April 12, 2017 to June 25, 2020 because the keypad may have one or more keys that become unresponsive or stuck. This could lead to infusion delay or prevent clinicians from changing fluid or medication infusions on the affected device.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/medical-devices/medical-device-recalls/becton-dickinson-bd-carefusion-303-inc-recalls-alaristm-system-pc-unit-and-pc-unit-front-case-keypad?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.