carecentrix

Provider Newsflash October 6, 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's Alaris Pump Module model 8100 and Pump Module Door Assembly Replacement Kits, part numbers 49000346, 49000239, 49000438 and 49000439 manufactured 12/1/2016 to 1/23/2019 and distributed 12/1/2016 to 1/23/2019 because the keypad may have one or more keys that become unresponsive or stuck. This may lead to an infusion delay or prevent clinicians from changing fluid or medication infusions on the affected devices.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/medical-devices/medical-device-recalls/becton-dickinson-carefusion-303-inc-recalls-alaristm-system-pump-module-and-pump-module-door?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.