carecentrix

**Provider Newsflash** Sept. 17, 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. Alaris Syringe Module-model 8110, Alaris PCA Module-model 8120, and Syringe/PCA Sizer Sensor Replacement Kit-P/N 122786 manufactured from March 1, 2010 to present and distributed from March 1, 2010 to March 12, 2020 due to incorrect display or syringe types and/or sizes. This could potentially result in delays in infusion, under-infusion or over-infusion leading to serious adverse events including death.

## 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-syringe-and-alaristm-pca-modules-due?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.