carecentrix

Provider Newsflash December 1, 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 Baxter Healthcare's SIGMA Spectrum Infusion Pumps with Master Drug Library (Versions 6 & 8) and Spectrum IQ Infusion Systems with Dose IQ Safety Software (V9) due to the potential for unplanned shutdown issues. This may cause an infusion delay or interruption in treatment.

What do I need to do?

- Please review the following recall notice:
- https://www.fda.gov/medical-devices/medical-device-recalls/baxter-healthcare-recalls-baxter-sigma-spectrum-infusion-pumps-master-drug-library-versions-6-and-8?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.