Provider Newsflash
July 2024

FDA Recall – Baxter International, Inc.

Purpose of this communication:

- We are writing to inform you that effective immediately the FDA has issued notice of a voluntary urgent recall of the Life2000 ventilator with an attached battery charger dongle. The recall is due to reports that the devices are not properly charging when there is damage to the battery charger dongle. Damage to the battery dongle prevents the ventilator's internal battery from charging or cause intermittent charging behavior which may leave the patient unable to use the device.

What do I need to do?

- Please review the following recall notice: Baxter Issues Urgent Medical Device Recall for Life2000 Ventilator Due to Potential Battery Charger Dongle Damage | FDA
- 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.

EDRC# 1604. 092021