Provider Newsflash
September 2024

FDA Recall – Abbott Diabetes Care Inc

Purpose of this communication:

- We are writing to inform you that the FDA has issued notice of a voluntary recall by Abbott Diabetes Care, Inc. their FreeStyle Libre 3 sensors after finding that a small number of FreeStyle Libre 3 sensors may provide incorrect high glucose readings, which if undetected may pose a potential health risk for people living with diabetes.

What do I need to do?

- Please review the following recall notice: https://www.fda.gov/medical-devices/medical-device-recalls/continuous-glucose-monitoring-cgm-sensor-recall-abbott-diabetes-care-inc-issues-recall-certain?utm_medium=email&utm_source=govdelivery
- Notify impacted patients and facilitate the replacement 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.