Abbott– Recall FreeStyle Libre® 3 Sensors

- On July 31, 2024, Abbott announced a consumer level recall of three lots of FreeStyle Libre 3 sensors because the sensors may provide incorrect high glucose readings. — The FreeStyle Libre 3 system includes a sensor, reader and app. This recall impacts the sensor only. The FreeStyle Libre 3 reader and app are not impacted.

- FreeStyle Libre 3 sensors were distributed nationwide in the first half of May 2024.

Product Description	NDC#	Lot#
FreeStyle Libre 3 Sensor	57599-0818-00	T60001948 T60001966 T60001969

- FreeStyle Libre 3 sensors provide continuous glucose monitoring levels for patients with diabetes.

- Internal testing determined that some of the sensors from among three lots may provide incorrect high glucose readings, which if undetected may pose a potential health risk for people living with diabetes and can lead to incorrect treatment decisions, such as taking insulin when not required.

- Patients are instructed to conduct a fingerstick test using any blood glucose meter if they experience symptoms that do not match the sensor glucose reading or suspect the reading may be inaccurate. The built-in blood glucose meter in the FreeStyle Libre 3 reader may be used to check glucose at any time.

- If the patient's product lot number is listed above, they need to visit www.FreeStyleConfirm.com, select "Confirm Sensor Serial Number," and enter

their serial number to confirm whether their sensor is affected by this voluntary medical device correction.

- Patients wearing an affected FreeStyle Libre 3 sensor should immediately discontinue use and dispose of any affected sensors in their possession. If the patient's sensor is impacted they will need to enter their contact information so Abbott can send them a replacement sensor at no charge.

- Contact Abbott by phone at 1-833-815-4273 for questions regarding this recall.