

## **Continuous Glucose Monitoring-Sensor Initialization and Return Form**

	ID JMBER: C G M R DATE: 10/12/2023 Version 2.0		
Instructions: This form reviews additional exclusions for the Continuous Glucose Monitoring ancillary study to determine if a participant is currently eligible to have the sensor placed. When the participant is determined to be eligible and has agreed to participate, the form records the sensor serial number. Parts A and B of this form are completed immediately before the FreeStyle Libre 3 Continuous Glucose Monitoring sensor is given to the participant. Part C is completed when the device is returned to the clinic and shipped to Abbott.			
	MINISTRATIVE INFORMATION  Completion Date:   Day   Year   Ob. Staff ID:   Day   Ob. Staff ID:   Day   Ob. Staff ID:   Day   Ob. Staff ID:   Ob		
0c. Would you be interested in participating in this part of the study, as I've described?			
	y ☐ Yes → Go to item 1		
	N □ No		
	Oc1. If no, why not?Save and close form		
A. CGM Sensor Exclusion Information			
1.	Do you have an MRI scan, CT scan, X-ray or diathermy treatment scheduled in the next 14 days?  Yes□ Y → Go to item 3  No□ N		
2.	Do you have any air travel scheduled over the next 14 days?  Yes□ Y  No□ N → Go to item 4		
	The CGM sensor cannot be worn through regular airport screening machines. Are you willing to request rnative security screening procedures for travel?  Yes□ Y → Go to item 4  No□ N		
3.	Are you willing to participate in the CGM study at a later date?  Yes□ <sub>Y</sub> → Save and close form  No□ <sub>N</sub> → Save and close form		
В.	CGM Sensor Initialization Information		
4.	CGM sensor serial number max length=11		
5.	Date of CGM sensor application:		

5a.	Time of CGM sensor application:
C.	CGM Sensor Return Information
6.	Did the participant return the device?  Yes□ Y  No□ N  Save and close form
7.	How many days did the participant wear the device?
8.	Date device returned to clinic:
9	Date device shipped to Abbott: