State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Abbott Recalls The Readers Used With The FreeStyle Libre, FreeStyle Libre 14 day, and FreeStyle Libre 2 Flash Glucose Monitoring Systems for Risk of Extreme Heat and Fire

Recall Date	Product Description	Recalling Firm	Recall Reason
4/06/2023	**FreeStyle Libre Flash Glucose Monitoring System, FreeStyle Libre 14 day Flash Glucose Monitoring System, FreeStyle Libre 2 Flash Glucose Monitoring System**	**Abbott** Abbott Park, Illinois	Abbott is recalling the FreeStyle Libre, Libre 14 day, and Libre 2 Flash Glucose Management Systems because the systems' reader devices, which use rechargeable lithium-ion batteries, may get extremely hot, spark, or catch on fire if not properly stored, charged, or used with its Abbott provided USB cable and power adapter.

Recall Class	Product Identification	Distribution	Affected Dates
I	all Reader serial numbers	4,210,785 Units Nationwide	November 2017 – February 2023

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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