State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Remote Diagnostic Technologies Recalls Philips Tempus Pro Patient Monitor For Full Screen Error Message On Patient Monitors

Recall Date	Product Description	Recalling Firm	Recall Reason
9/27/2023	**Remote Diagnostic** Philips Tempus Pro Patient Monitor, REF: 00-1004-R, 00-1007-R, 00-1024-R, and 00-1026-R	**Remote Diagnostic** Farnborough United Kingdom	Full screen error message may occur on patient monitors with affected hardware version, either during or after laryngoscopes are unplugged from monitors

Recall Class	Product Identification	Distribution	Affected Dates
II	**Remote Diagnostic** Philips Tempus Pro Patient Monitor REF/UDI-DI: 00-1004-R/05060472440020, 05060472442901	5773 Units Nationwide including California	August 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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