California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Remote Diagnostic Technologies Ltd. Tempus Pro Patient Monitor, non-printer, and printer

Recall Date	Product Description	Recalling Firm	Recall Reason
04/24/2025	Tempus Pro Patient Monitor REF: 00-1004-R Tempus Pro non-printer, IBP, without headset dock REF: 00-1004 Tempus Pro non-printer, IBP, with headset dock REF: 00-1007-R and 00-1007 Tempus Pro printer REF: 00-1024-R Tempus Pro printer, IBP REF: 00-1026-R	Remote Diagnostic Technologies Ltd.	Software issue that causes an error screen on a portable vital signs monitor when attaching video Laryngoscope.

Recall Class	Product Identification	Distribution	Affected Dates
II	All Lots REF: 00-1004-R UDI-DI: 05060472440020, 05060472442901 REF: 00-1007-R UDI-DI: 05060472440013, 05060472442918	59 units in California	April and prior

	REF: 00-1024-R UDI-DI: 05060472441027, 05060472442925 REF: 00-1026-R UDI-DI: 05060472441058, 05060472442932 REF: 00-1004 UDI-DI: 05060472440624 REF: 00-1007 UDI-DI: 05060472440655		

For additional information, please visit the FDA Website.