California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Philips Medical Systems Nederland B.V. Philips IntelliSpace Cardiovascular software

Recall Date	Product Description	Recalling Firm	Recall Reason
02/28/2025	Philips IntelliSpace Cardiovascular software Model 830089	Philips Medical Systems Nederland B.V.	Study data is not able to be archived, copied, or exported with the cardiovascular software version.

Recall Class	Product Identification	Distribution	Affected Dates
II	Software version 7.0.0.0 UDI: (01)00884838115378(11)221214 (10)7.0.0.0.	139 units nationwide	April 2024 and prior

For additional information, please visit the FDA Website.