California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

MEDLINE INDUSTRIES, LP Lab Kit, SKU DYLAB1018

Recall Date	Product Description	Recalling Firm	Recall Reason
11/12/2024	Lab Kit, SKU DYLAB1018 Component No. 503581	MEDLINE INDUSTRIES, LP - Northfield	Specimen container included in kits is labeled as sterile, but has been identified to be non-sterile. Long-term consequences may include unnecessary treatment and/or prolonged hospitalization.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI-DI: 40195327088393 (Case), 10195327088392 (Ea) Lots: 24IMC246	6 units in California.	October 23rd and prior.

For additional information, please visit the FDA Website.