

## California Device Recall Information Sheet

### Food and Drug Branch – Device Recalls

#### Medline Industries Recall for Urine Collection Vhc and Clean Catch Kit

| Recall Date | Product Description   | Recalling Firm                      | Recall Reason  |
|-------------|---|-------------------------------------|--|
| 11/12/2024  | Urine Collection Vhc, Sku Dylab1016; Component No. 503581<br><br>Clean Catch Kit, Sku Dykm1833A; 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: 40195327139941 (Case), 10195327139940 (Ea), Lots 22HLA350 22HLB291<br><br>UDI-DI: 40193489240697 (Case); 10193489240696 (Ea) Lots: 19LAA531 20BAA260 21KBJ974 22JBK243 22LBJ966 23BBU789 23FBN998 23HBW916 23IBH024 24CBE580 24DBT234 24GBA986 24GBL321 | 626,305 Total Kits in the US Nationwide including CA<br><br>0 sales were recorded for Dykm1833A<br><br>0 sales were recorded for Dylab 1016 | October 2024 and Prior |

For additional information, please visit this [FDA Website VHC Product](#) and this [FDA Website CCKit Product](#).