California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Becton Dickinson & Co. Recalls BD BBL Sensi Disc Ampicillin 2 for Susceptibility Testing

Recall Date	Product Description	Recalling Firm	Recall Reason
12/11/2024	BD BBL Sensi Disc Ampicillin 2 ¿g (AM-2) - In-Vitro BD BBL Sensi Disc Ampicillin 2 ¿g (AM-2) are used for semi-quantitative in vitro susceptibility testing by the agar disc diffusion test procedure of common, rapidly growing and certain fastidious bacterial pathogens.	Becton Dickinson & Co.	BD identified through potency testing as part of a stability test request to monitor Ampicillin AM-2 due to decrease in potency results of 65% at 18 months, may result in falsely resistant result for ampicillin susceptibility.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI-DI/Lot Numbers: 30382902312636/2339360, 3010977, 3058508, 3184064, 3234190	10 Units in California	December 2024 and prior.

For additional information, please visit the FDA Website.