State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Becton Dickinson Infusion Therapy Systems Recalls BD Nexiva for Breach

Recall Date	Product Description	Recalling Firm	Recall Reason
9/8/2021	**BD Nexiva, 20 GA 1.00 IN** (1.1 X 25 mm), 3660 ml/hr. 61ml/min; Closed IV Catheter System- Over-the-needle, intravascular catheters.	**Becton Dickinson Infusion Therapy Systems Inc.** Sandy, Utah	There is a breach in the product packaging that renders the product non-sterile.

Recall Class	Product Identification	Distribution	Affected Dates
II	Catalog number/Model number: 383536; Batch numbers/UDI numbers: Lot: 1193055/(17)240630(10)1193055(30)80/(01)50382903835367, 2024-06-30; Lot: 1166273: (17)240630(10)1166273(30)80/(01)50382903835367, 2024-06-30 Lot: 1166273: (17)240630(10)1166273(30)80/(01)50382903835367, 2024-06-30 Lot: 1188953: (17)240630(10)118895 3(30)80/(01)50382903835367, 2024-06-30	155,840 devices, 80 units in each box Nationwide including California	September 2021 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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