California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Arrow International, LLC Recalls Arrow Kits and Sets that contain Lidocaine and Bupivacaine

Recall Date	Product Description	Recalling Firm	Recall Reason
6/13/2026	ARROW INTERNATIONAL, LLC Recalls Arrow Kits and Sets that contain Lidocaine and Bupivacaine	ARROW INTERNATIONAL, LLC Morrisville, NC	Lidocaine, Bupivacaine, and 0.9% sodium chloride, the subject of a supplier recall, were included into kits and sets. The supplier was found to have quality issues and limited reports of drug ineffectiveness, and if sterility of a water-based injectable product is compromised this could lead to serious or life-threatening injuries.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Product 1**	1,516 Units Nationwide	April 2026 and prior

I	**Product 2**	1,525,831 Units Nationwide	April 2026 and prior
I	**Product 3**	62,719 Units Nationwide	April 2026 and prior
I	**Product 4**	426,041 Units Nationwide	April 2026 and prior
I	**Product 5**	180 Units Nationwide	April 2026 and prior
I	**Product 6**	126,530 Units Nationwide	April 2026 and prior
I	**Product 7**	37,738 Units Nationwide	April 2026 and prior
I	**Product 8**	90 Units Nationwide	April 2026 and prior
I	**Product 9**	1,061,069 Units Nationwide	April 2026 and prior
I	**Product 10**	8,265 Units Nationwide	April 2026 and prior

For additional information, please visit FDA website.