State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

California Device Recall Information Sheet

Baxter Recalls Novum Iq Large Volume Pump

Recall Date	Product Description	Recalling Firm	Recall Reason
8/20/2025	**Baxter Novum Iq Large Volume Pump** Product Code REF 40700BAXUS	**BAXTER HEALTHCARE** Deerfield, Illinois	Potential for under-infusion when transitioning from a flow rate to a higher flow rate that is more than double. (e.g., rate change or bolus). The level of under-infusion is variable based on the current infusion rate, the duration the pump has been running at this flow rate, and the magnitude of the rate change. The longer the duration the pump has been running at the current infusion rate and the larger the magnitude of the rate change, the larger the under-infusion that would be experienced. Additionally, Baxter has identified an increase in customer reports of over and under-infusion potentially due to set misloading. Failure to properly load the tubing into the pump channel may result in the

			pump infusing at a rate higher or lower than programmed. Consistent with the instructions for use, customers should ensure that: 1) The door is fully open before loading the set. 2) The tubing is taut and loaded without slack in the pumping channel.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Novum Iq Large Volume Pump** UDI/DI 05413765851797, All Serial Numbers	36705 Units Nationwide	August 2025 and prior

For additional information, please visit the FDA website

CDPH Food and Drug Branch
MS 7602 ● P.O. Box 997435 ● Sacramento, CA 95899-7435
(916) 650-6500 ● (916) 650-6650 FAX
Internet Address: www.cdph.ca.gov