California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Recalls Novum IQ Syringe Pump

Recall Date	Product Description	Recalling Firm	Recall Reason
8/9/25	Novum IQ Syringe Pump, Product Code 40800BAXUS, infusion pump	Baxter Healthcare Corporation	Baxter Healthcare Corporation is issuing an Urgent Medical Device Correction for the Novum IQ Large Volume Pump (LVP) and Novum IQ Syringe Pump (SP) due to software anomalies that may result in a blank Run screen (LVP and SP) and/or false motor movement system error (SP only).

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI/DI 05413765852428, All Serial Numbers	10,101 units Worldwide	August 2025 and prior.

For additional information, please visit the FDA Website.