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 Healthcare Corporation Recalls Novum IQ Syringe Infusion System For Misalarm

Recall Date	Product Description	Recalling Firm	Recall Reason
11/15/2023	**Novum IQ Syringe Infusion System** Product Code 40800BAXUS	**BAXTER HEALTHCARE CORPORATION** Deerfield, Illinois	Baxter is issuing an Urgent Medical Device Correction for the Novum IQ Syringe Pump. Baxter identified that after multiple downstream occlusion alarms, the pump may display an Infusion Complete alarm even though uninfused fluid remains in the syringe.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Novum IQ Syringe Infusion System** All Serial Numbers	2023 Units Nationwide including California	13 October 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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