California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Baxter NovumIQ Syringe Infusion System

Recall Date	Product Description	Recalling Firm	Recall Reason
03/21/2025	Baxter NovumIQ Syringe Infusion System REF: 40800BAXUS	Baxter Healthcare Corporation	Baxter Healthcare Corporation is issuing an Urgent Medical Device Recall for the Novum IQ syringe pumps listed below due to an incorrectly installed gasket. A gasket failure could lead to potential fluid ingress due to cleaning, or as a result of an IV fluid spill.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI/DI: 05413765852428 Serial Numbers: 02A220600103 - 02A240800142	13 units in California	March and prior

For additional information, please visit the FDA Website.