California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Baxter Sigma Spectrum Infusion System and Baxter Spectrum IQ Infusion System with Dose IQ Safety Software

Recall Date	Product Description	Recalling Firm	Recall Reason
03/06/2025	Baxter Sigma Spectrum Infusion System (V6 Platform), Product Code 35700BAX Baxter Spectrum IQ Infusion System with Dose IQ Safety Software, Product Code 3570009	Baxter Healthcare Corporation	There is the potential for missing motor mounting screws, which may have occurred during the servicing process.

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI/DI: 00085412091570 Serial Numbers: 704198 - 1024109 UDI/DI: 00085412610900 Serial Numbers: 3020290 - 3718985	66 units nationwide 323 units nationwide	February and prior

For additional information, please visit: FDA Website (Sigma Spectrum) and FDA Website (Spectrum IQ).