California Device Recall Information Sheet

Smiths Medical Asd Inc. Recalls Cadd-Solis Vip Ambulatory Infusion Pumps

Recall Date	Product Description	Recalling Firm	Recall Reason
8/14/2024	**Cadd Solis Vip Ambulatory Infusion Pump** Model 2120 (\x1c21-2120\x1d, \x1c21-2125\x1d, and \x1c21-2127\x1d) Indicated for the following uses: Intra-arterial, subcutaneous, intraperitoneal, in close proximity to nerves, into an intraoperative site, epidural space or subarachnoid space. Therapies that require a continuous rate of infusion, intermittent bolus, and/or with patient-controlled demand doses.	**SMITHS MEDICAL ASD INC.** Minneapolis, Mn	The CADD Solis VIP Ambulatory Infusion Pump is indicated for the following uses: " For intravenous, intraarterial, subcutaneous, intraperitoneal, perineural, surgical site, epidural space, or subarachnoid space infusion.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Product 1** UDI-DI: 10610586040429, 30610586040430, 10610586038501, 10610586042591, 10610586042607, 10610586042614, 10610586042829, 15019517084368	Nationwide	August 2024 and prior

For additional information, please visit the FDA website.