California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Integra LifeSciences Corp. (NeuroSciences) Recalls Extended Tip Applicator

Recall Date	Product Description	Recalling Firm	Recall Reason
6/20/25	Extended Tip Applicator, 15 CM, Box of 5.	Integra LifeSciences Corp. (NeuroSciences)	Devices contain potentially out-of-specification levels of endotoxins and lack of sterility assurance.
6/20/25	Extended Tip Applicator, 8CM, Box of 5.	Integra LifeSciences Corp. (NeuroSciences)	Devices contain potentially out-of-specification levels of endotoxins and lack of sterility assurance.
6/20/25	Extended Tip Applicator 8CM, Box of 1.	Integra LifeSciences Corp. (NeuroSciences)	Devices contain potentially out-of-specification levels of endotoxins and lack of sterility assurance.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model/Catalog Number: 205115; UDI: 10381780000150; All Lots.	2,524 units Globally	May 2025 and prior.
I	Model/Catalog Number: 205108; UDI: 10381780000143; All Lots.	6,216 units Globally	May 2025 and prior.
I	Model/Catalog Number: 201108; UDI 10381780000167; All Lots.	212 units Globally	May 2025 and prior.

For additional information, please visit the FDA Website Extended Tip Applicator, 15 CM, Box of 5, Extended Tip Applicator, 8CM, Box of 5, Extended Tip Applicator 8CM, Box of 1.