

California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Integra LifeSciences Corp. (NeuroSciences) CODMAN Disposable Perforator and Craniotomy Kit

Recall Date	Product Description	Recalling Firm	Recall Reason
05/06/2025	CODMAN Disposable Perforator, 14mm Cranial perforator CODMAN Craniotomy Kit, Disposable Perforator Cranio-blade Wire Pass Drill, 14mm Cranial perforator kit	Integra LifeSciences Corp. (NeuroSciences)	Inadequate weld that can potentially cause the product to disassemble.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model Number: 261221 UDI: 10381780513599 Lot Number: 5580821 - 7400949 Model Number: 261230 UDI: 10381780513629 Lot Number: 5984530 - 7379578	172,350 units nationwide	April and prior

For additional information, please visit: [FDA Website \(Disposable Perforator\)](#) and [FDA Website \(Craniotomy Kit\)](#).