California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Cartiva Inc. Recalls Cartiva Synthetic Cartilage Implants

Recall Date	Product Description	Recalling Firm	Recall Reason
12/11/2024	Cartiva Synthetic Cartilage Implant (SCI) Catalog: CAR-06-US (6mm), CAR-08-US (8mm), CAR-10-US (10mm), CAR-12-US (12mm)	Cartiva Inc. Alpharetta, GA	Patients implanted with synthetic cartilage implant, may experience a higher-than expected occurrence rate of the following hazards: revision, removal, implant subsidence, displacement, pain, nerve damage or fragmentation.

Recall Class	Product Identification	Distribution	Affected Dates
II	Catalog/UDI-DI: CAR-06-US/00852897002328, CAR-08-US/00852897002021, CAR-10-US/ 00852897002038, CAR-12-US/00852897002335	634 units in California	December 2024 and prior.

For additional information, please visit the FDA Website.