California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Smith & Nephew Inc. Recalls BIORAPTOR Suture Anchors

Recall Date	Product Description	Recalling Firm	Recall Reason
12/11/24	Bioraptor Suture Anchors Tendon/ligament, non-bioabsorbable bone anchor	SMITH & NEPHEW INC. Andover, MA	Sterile barrier breach due to inadequate packaging design that could not hold the geometry of the device.

Recall Class	Product Identification	Distribution	Affected Dates
II	Bioraptor Suture Anchors 72201541 BIRPTR 2.3 PK SUTURE ANCHOR W/ ULTRAB 2146369, 2151242 72201542 BIRPTR 2.3 PK SUT ANCHR W/ ULTRAB BLK 2151690, 2151691, 2153489 72203280 BIORAPTOR CRV 2.3 PK SA UB COBRD BLACK 2151422 72203281 BIORAPTOR CRV 2.3 PK SA UB COBRD BLUE 215169	4 Units in California	December 2024 and prior

For additional information, please visit the FDA Website.