California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Moximed, Inc. MISHA Knee System Implant Small, Left and Right

Recall Date	Product Description	Recalling Firm	Recall Reason
12/23/2024	MISHA Knee System Implant Small, Left, REF: 2-1001 MISHA Knee System Implant Small, Right, REF: 2-1002.	Moximed, Inc.	Knee implant may fracture due to a supplier-related issue with one component that has nonconforming geometry at the distal end of the absorber component that may result in a reduced wall thickness and microcracks, which may lead to new or worsening symptoms of discomfort, swelling / edema, pain or stiffness, which may necessitate removal of the device.

Recall Class	Product Identification	Distribution	Affected Dates
II	REF: 2-1001 UDI-DI: 00856047005795 Lot(Expiration): 23032901(29-Mar-25), 23092102(21-Sep-25), 24032601(26-Mar-26)	41 units in California	December and prior

	REF: 2-1002 UDI-DI: 00856047005801 Lot(Expiration): 23032902(29-Mar-25), 23092503(25-Sep-25), 24040801(8-Apr-26)		

For additional information, please visit the FDA Website.