California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Zimmer, Inc. NexGen LPS Flex Knee Joint

Recall Date	Product Description	Recalling Firm	Recall Reason
03/24/2025	NexGen LPS Flex, Knee joint patellofemorotibial polymer/metal/polymer semi-constrained cemented prosthesis Model Number: 00-5962-042-10	Zimmer, Inc.	The "Use with plate 7, 8, 9, 10" compatibility statement on top of the label does not match with the compatibility statements "USE WITH PLATE 5, 6" in the translations section of the same label.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI-DI: 00889024666214 Lot Numbers: 66789408, 66792792	5 units in California	February and prior

For additional information, please visit the FDA Website.