State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Intera Oncology Recalls Intera 3000 Hepatic Artery Infusion Pump

Recall Date	Product Description	Recalling Firm	Recall Reason
8/31/2022	**INTERA 3000 Hepatic Artery Infusion Pump** 30 mL, Sterile, Rx Only. implantable infusion pump	**Intera Oncology, Inc.** Wellesley, Massachusetts	Higher than expected flow rate.

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI-DI: 00850014110147; Catalogue No. AP03000H; Serial No. 16145 – 16485	50 units in multiple states, including California	July 2022 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

CDPH Food and Drug Branch
MS 7602 ● P.O. Box 997435 ● Sacramento, CA 95899-7435
(916) 650-6500 ● (916) 650-6650 FAX
Internet Address: www.cdph.ca.gov