



Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

ArjoHuntleigh Polska Recalls Sara Plus Floor Lift for Risk of Smoke or Fire When Lift Is Used with Depleted Battery

Recall Date	Product Description	Recalling Firm	Recall Reason
4/08/2022	Celltrion DiaTrust COVID-19 Ag Rapid Test Reference No. CT-P60 D-2 02	ARJOHUNTLEIGH POLSKA Sp. z.o.o. Komorniki, Poland	The device may emit smoke or ignite.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model Numbers: HEP0001, HEP0001-AU, HEP0001-BR, HEP0001-CN, HEP0001-JP, HEP0001-UK, HEP0001-US, HEP1001, HEP1001-AU, HEP1001CON4869, HEP1001-US, HEP2001, HEP2001-BR, HEP2001-CN, HEP2001-UK	4,449 Units Nationwide	April 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

