



California Device Recall Information Sheet

Food and Drug Branch - Device Recalls

Medical Depot Inc. Go 2 DC Car Adapter

Recall Date	Product Description	Recalling Firm	Recall Reason
	iGo¿2 DC Car Adapter to be used with iGo¿2 Portable Oxygen Concentrator Systems. Power cord allows the user to operate the device in a vehicle.	•	Potential for DC Power Supply housing to become hot to the touch and deform the plastic housing.

Recall Class	Product Identification	Distribution	Affected Dates
	DV6X-619; UDI 885304020585; Cord Revision: Rev E; Lot No. beginning with 2023-01-01 through 2024-12-31. iGo2 Portable Oxygen Concentrator Systems Code Info: Catalog No. 125D, 125D-XB, 125D-BT, 125D-BT-XB, 125D-ARYA-XB; UDI 885304022237, 885304032250, 885304033257, 885304033240, 885304033516; Lot No. D23125001DS to D23A26121DS, D23706001DA to D23B16056DA, F23B06001DS to F25108143DS,	89,532 units Worldwide distribution - US Nationwide and the countries of Albania, Austria, Bahrain, Belgium, Bosnia and Herzegovina, Bulgaria, Canada, China, Colombia, Croatia, Cyprus, Czech Republic, France, Germany, Great Britain, Greece, Hong Kong, Hungary, Iceland, Israel, Italy, Japan, Kenya, Kuwait, Latvia, Libya, Lithuania, Malaysia, Maldives, Nepal, Netherlands, Norway, Oman, Philippines, Poland, Portugal, Qatar, Romania, Russia,	

Saudi Arabia, Serbia, Slovak Republic, Slovenia, South Korea, Spain, Sri Lanka, Sweden, Switzerland, Taiwan, Thailand, Trinidad & Tobago, Turkey, Uganda,
Ukraine, United Arab
Emirates.

For additional information, please visit the <u>FDA Website</u>.