California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

mo-Vis BVBA IDM-MICRO-R. Electrical wheelchair component.

Recall Date	Product Description	Recalling Firm	Recall Reason
9/2/25	IDM-MICRO-R. Electrical wheelchair component.	mo-Vis BVBA	Devices with firmware versions 2.3 and lower may experience timing variations between software tasks, which could result in skipping the necessary neutral check and the wheelchair producing unintended movement upon power-up when the joystick is out of neutral.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model No. PRSPS0014; UDI-DI: 05407008320836; Serial No. 1000 to 1690.	15 units US Nationwide	July 2025 and prior.

For additional information, please visit the FDA Website.