California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Qapel Medical Inc. Hippo 072 Aspiration System including Cheetah Delivery Tool and Aspiration Tubing

Recall Date	Product Description	Recalling Firm	Recall Reason
04/07/2025	Hippo 072 Aspiration System including Cheetah Delivery Tool and Aspiration Tubing REF: APT6072-132 Hippo 072 Aspiration System including Cheetah Delivery Tool REF: AP6072-132 072 Aspiration Tubing REF: APT-95	Qapel Medical Inc.	Aspiration catheter distal tip features and characteristics may not be in the scope of FDA clearance; tip detachment, vessel rupture, and vasospasm could occur.

Recall Class	Product Identification	Distribution	Affected Dates
I	REF/UDI-DI/Lot(Expiration): REF: APT6072-132 UDI-DI: 00857545008127 Lot Numbers (Expiration): FG241008C-03 (04/07/2025) - FG240905C-04 (03/06/2025) REF: AP6072-132 UDI-DI: 00857545008097 Lot Numbers (Expiration): FG241206A-03 (06/08/2025), FG240917A-01 (03/17/2025)	1,617 units nationwide	February and prior

	REF: APT-95 UDI-DI: 00857545008103 Lot Numbers (Expiration): FG241206A-04 (06/08/25)		

For additional information, please visit the FDA Website.