California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

AVID Medical, Inc. Recalls Halyard CARDIAC CATH TRAY SHANDS JAX Kit

Recall Date	Product Description	Recalling Firm	Recall Reason
5/1/2026	Halyard CARDIAC CATH TRAY SHANDS JAX kit. Model Number: SHND401-04, SHND401-05	AVID Medical, Inc. Toano, VA	Potential risk for Medline syringe rotating adaptor unwinding during use, which may result in a loose connection and/or full disconnection between the syringe and manifold.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model Number: SHND401-04. UDI-DI (Kit Lot Numbers): 10809160395615 (1577749, 1577750, 1590780, 1590782, 1590783, 1590784, 1590785, 1590833, 1591141, 1591142, 1597561, 1600738, 1613343, 1622356, 1631890, 1643855, 1643856, 1643860, 1644853, 1645163, 1647458, 1647459, 1647460, 1650983, 1652365, 1652366)	6,956 Units US distribution to FL, IL, MO, MS, NC, NE, TX	March 2026 and prior

	Model Number: SHND401-05. UDI-DI (Kit Lot Numbers): 10809160491294 (1652367, 1668844, 1668845, 1668846, 1669188, 1669189, 1669504, 1669506, 1669507, 1671312, 1671313, 1671314, 1671315, 1671316)		

For additional information, please visit the FDA website