California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Stryker GmbH Guide Wire with Ruler Tube, 3x800 mm and 3x1000 mm

Recall Date	Product Description	Recalling Firm	Recall Reason
04/17/2025	Guide Wire with Ruler Tube 3x800 mm DIA Catalog number/REF: 2351-3080S Guide Wire with Ruler Tube 3x1000 mm DIA Catalog number/REF: 2351-3100S	Stryker GmbH	The metal ring at the end of the Guide Wire with Ruler Tube may detach from the main body of the instrument during use in surgery.

Recall Class	Product Identification	Distribution	Affected Dates
II	Catalog number/REF: 2351-3080S UDI-DI: 07613327361797 Lot #: KU155697 - KU155712 Catalog number/REF: 2351-3100S UDI-DI: 07613327361803 Lot #: KU164143 - KU184049	139 units in California	March and prior

For additional information, please visit: FDA Website (3x800 mm) and FDA Website (3x1000 mm).