State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Teleflex Recalls Endotracheal Tube Rushelit and Magill Products For Disconnection From the Connector

Recall Date	Product Description	Recalling Firm	Recall Reason
7/5/2023	**RUSCHELIT** REF 112480025, 112480030, 112480035, 112480040, 112480045	**TELEFLEX LLC** Morrisville, NC	Reports of disconnection of the 15mm connector from the endotracheal tube (ET tube) for the affected products.
7/5/2023	**Magill—Low Pressure Cuff** REF 112080050, 112080055, 112080060, 112080065, 112080070, 112080075, 112080080, 112080085, 112080090, 112080095, 112080100	**TELEFLEX LLC** Morrisville, NC	Reports of disconnection of the 15mm connector from the endotracheal tube (ET tube) for the affected products.
7/5/2023	**Magill Uncuffed** REF 100380025, 100380030, 100380035, 100380040, 100380045, 100380050, 100380055, 100380060, 100380065	**TELEFLEX LLC** Morrisville, NC	Reports of disconnection of the 15mm connector from the endotracheal tube (ET tube) for the affected products.

Recall Class	Product Identification	Distribution	Affected Dates
I	**RUSCHELIT** UDI/DI 14026704341495, Batch Numbers: 18FG31-KME23A2313…	186,940 Units Nationwide	May 2023 and Prior

I	**Magill—Low Pressure Cuff** UDI/DI 14026704341266, Batch Numbers: 18FG07-KME22C0947…	138,124,178 Units Nationwide	May 2023 and Prior
I	**Magill Uncuffed** UDI/DI 14026704340610, Batch Numbers: 19ET51-KME21D0517…	25,921 Units Nationwide	May 2023 and Prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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