

**K113333 KIMBERLY-CLARK KIM VENT MICROCUFF
ENDOTRACHEAL TUBE FOR ADULTS**May 11, 2012
179 days to decisionK113333 · Product code: **BTR** · Anesthesiology
Source: <https://www.510kdatabase.net/k113333/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tracheal (w/wo Connector) (BTR)
Date received	Nov 14, 2011
Decision date	May 11, 2012
Days to decision	179 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kimberly Clark Corporation
Location	Neenah, WI, US
Contact	MARCIA JOHNSON
510(k) history	13 submissions · 12 cleared · 2010-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k113333/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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