FDA 510(k) Premarket Notification

K113333 KIMBERLY-CLARK KIM VENT MICROCUFF ENDOTRACHEAL TUBE FOR ADULTS

CLEARED

May 11, 2012
179 days to decision

K113333 · 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