FDA 510(k) Premarket Notification

K062240 CAREVENT

CLEARED

Oct 16, 2006
75 days to decision

K062240 · Product code: **KNY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k062240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Aug 2, 2006
Decision date	Oct 16, 2006
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Chief Medical, LLC**
Location	Teton Village, WY, US
Contact	SCOTT HORN
510(k) history	1 submissions · 1 cleared · 2006-2006