FDA 510(k) Premarket Notification

K002901 KENDALL 14.5 FR MAXID CUFFED DUAL LUMEN CATHETER

CLEARED

Mar 28, 2001
191 days to decision

K002901 · Product code: **MSD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k002901/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Sep 18, 2000
Decision date	Mar 28, 2001
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Kendall Company**
Location	Mansfield, MA, US
Contact	PAUL EVANS
510(k) history	8 submissions · 8 cleared · 2001-2006