

**K002901 KENDALL 14.5 FR MAXID CUFFED DUAL LUMEN
CATHETER**Mar 28, 2001
191 days to decisionK002901 · 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

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