FDA 510(k) Premarket Notification

K022678 MEDCOMP SC4

CLEARED

Feb 24, 2003
196 days to decision

K022678 · Product code: **MSD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k022678/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Aug 12, 2002
Decision date	Feb 24, 2003
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medcomp**
Location	Mchenry, IL, US
Contact	FLORENCE A CAIKOSKI
510(k) history	40 submissions · 34 cleared · 1982-2014