FDA 510(k) Premarket Notification

K030502 14.5F X 55CM HEMO-FLOW DOUBLE LUMEN CATHETER, MODELS HFS-55, HFT-55

CLEARED

May 20, 2003
90 days to decision

K030502 · Product code: **MSD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k030502/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Feb 19, 2003
Decision date	May 20, 2003
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medcomp**
Location	Mchenry, IL, US
Contact	MEGHAN J TINTLE
510(k) history	40 submissions · 34 cleared · 1982-2014