

**K030502 14.5F X 55CM HEMO-FLOW DOUBLE LUMEN
CATHETER, MODELS HFS-55, HFT-55**May 20, 2003
90 days to decisionK030502 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030502/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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