

**K022678 MEDCOMP SC4**Feb 24, 2003  
196 days to decisionK022678 · Product code: **MSD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k022678/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022678/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 7, 2026