

**K982333 MODIFICATION TO ULDALL DOUBLE LUMEN  
HEMODIALYSIS CATHETER SET AND TRAY**Feb 12, 1999  
225 days to decisionK982333 · Product code: **MSD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k982333/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Jul 2, 1998
Decision date	Feb 12, 1999
Days to decision	225 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Cook, Inc.</b>
Location	Mchenry, IL, US
Contact	APRIL LAVENDER
510(k) history	190 submissions · 179 cleared · 1976-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982333/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026