

**K781050 DLC 500, DOUBLE LUMEN CATHETER**Sep 26, 1978  
97 days to decisionK781050 · Product code: **KOC** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k781050/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Jun 21, 1978
Decision date	Sep 26, 1978
Days to decision	97 days
Third-party review	No

**APPLICANT**

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Company	<b>Hemodialysis, Division of The</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k781050/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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