

**K915215 CONTINUOUS ARTERIOVENOUS HEMOFILTRATION  
CATH. KITS**Mar 27, 1992  
130 days to decisionK915215 · Product code: LFK · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k915215/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Femoral (LFK)
Date received	Nov 18, 1991
Decision date	Mar 27, 1992
Days to decision	130 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Akcess Medical Products, Inc.</b>
Location	New Brunswick, NJ, US
Contact	B KAPANY
510(k) history	36 submissions · 28 cleared · 1989-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k915215/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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