

**K904433 DUAL LUMEN NEEDLES**Nov 23, 1990  
56 days to decisionK904433 · Product code: **LBW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k904433/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialysis, Single Needle (co-axial Flow) (LBW)
Date received	Sep 28, 1990
Decision date	Nov 23, 1990
Days to decision	56 days
Third-party review	No

**APPLICANT**

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Company	<b>Akcess Medical Products, Inc.</b>
Location	New Brunswick, NJ, US
Contact	BALBIR 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/k904433/>; 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 July 2, 2026