

**K900841 HEMODIALYSIS TRANSDUCER PROTECTOR**May 15, 1990  
82 days to decisionK900841 · Product code: **FIB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900841/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Protector, Transducer, Dialysis (FIB)
Date received	Feb 22, 1990
Decision date	May 15, 1990
Days to decision	82 days
Third-party review	No

**APPLICANT**

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Company	<b>Mediflex Intl.</b>
Location	Mchenry, IL, US
Contact	UDINE
510(k) history	9 submissions · 8 cleared · 1984-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900841/>, 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 30, 2026