

**K903945 EXEL A-V FISTULA SET**Nov 20, 1990  
85 days to decisionK903945 · Product code: **FIE** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k903945/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Aug 27, 1990
Decision date	Nov 20, 1990
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>Exel Intl.</b>
Location	Culver City, CA, US
Contact	ESHAGH HAMID
510(k) history	18 submissions · 18 cleared · 1986-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903945/>; 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