

**K900125 ARTERIOVENOUS FISTULA SETS**Jan 31, 1990  
22 days to decisionK900125 · Product code: **FIE** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k900125/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Jan 9, 1990
Decision date	Jan 31, 1990
Days to decision	22 days
Third-party review	No

**APPLICANT**

---

Company	<b>Baxter Healthcare Corp</b>
Location	Mchenry, IL, US
Contact	ROBERT L WILKINSON
510(k) history	505 submissions · 496 cleared · 1977-2019

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900125/>; 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 May 17, 2026