

**K903789 AVI MICRO 885 DUAL CHANNEL INFUSION PUMP**Aug 30, 1990  
10 days to decisionK903789 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k903789/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Aug 20, 1990
Decision date	Aug 30, 1990
Days to decision	10 days
Third-party review	No

**APPLICANT**

---

Company	<b>3M Avi, Inc.</b>
Location	St. Paul, MN, US
Contact	VON BUSCH
510(k) history	18 submissions · 18 cleared · 1990-1993

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903789/>; 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 7, 2026