

K884243 BLOOD TUBING SETDec 8, 1988
58 days to decisionK884243 · Product code: **FJK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k884243/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Tubing, Blood, With And Without Anti-regurgitation Valve (FJK)
Date received	Oct 11, 1988
Decision date	Dec 8, 1988
Days to decision	58 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884243/>; Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026