

K860241 JMS AV FISTULA NEEDLEFeb 5, 1986
12 days to decisionK860241 · Product code: **FIE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k860241/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Jan 24, 1986
Decision date	Feb 5, 1986
Days to decision	12 days
Third-party review	No

APPLICANT

Company	Jms Co., Ltd.
Location	Hiroshima, JP
Contact	KATSUTO TAKATSUKI
510(k) history	24 submissions · 24 cleared · 1985-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860241/>, 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 29, 2026