

K915174 JMS APHERESIS NEEDLEMar 2, 1993
473 days to decisionK915174 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k915174/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Nov 15, 1991
Decision date	Mar 2, 1993
Days to decision	473 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k915174/> 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