FDA 510(k) Premarket Notification

K860241 JMS AV FISTULA NEEDLE

CLEARED

Feb 5, 1986
12 days to decision

K860241 · Product code: **FIE** · Gastroenterology & Urology
Source: 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