

**K000843 MODIFICATION TO JMS APHERESIS NEEDLE**Apr 10, 2000  
26 days to decisionK000843 · Product code: **FIE** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k000843/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Fistula (FIE)
Date received	Mar 15, 2000
Decision date	Apr 10, 2000
Days to decision	26 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Jms Co., Ltd.</b>
Location	Hiroshima, JP
Contact	KEISUKE URATOMI
510(k) history	24 submissions · 24 cleared · 1985-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000843/> 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 June 29, 2026