

**K921833 ENDOPATH ENDO-SURGERY PERCUTANEOUS
CATHETER INTROD**Sep 21, 1992
158 days to decisionK921833 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k921833/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 16, 1992
Decision date	Sep 21, 1992
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ethicon, Inc.
Location	Raritan, NJ, US
Contact	MELISSA WALKER
Website	https://www.jnjmedtech.com
510(k) history	204 submissions · 197 cleared · 1976-2026

Ethicon, Inc. is a subsidiary of Johnson & Johnson specializing in surgical sutures and wound closure devices. The company is headquartered in Raritan, United States. Ethicon has received FDA 510(k) clearances from total submissions since 1976. The company's regulatory focus centers on General & Plastic Surgery devices, which represent the majority of its cleared submissions. The latest FDA 510(k) clearance was granted in 2026, demonstrating continued regulatory activity. Ethicon has manufactured surgical sutures and wound closure technologies since 1887. The company hold...

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Device record: <https://www.510kdatabase.net/k921833/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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