

**K912398 ENDOPATH SURG DISP
TROCAR(SHIELD/UNSHIELDED),MODIF**Dec 17, 1991
207 days to decision

K912398 · Product code: HET · Obstetrics & Gynecology

Source: <https://www.510kdatabase.net/k912398/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	May 24, 1991
Decision date	Dec 17, 1991
Days to decision	207 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ethicon, Inc.
Location	Raritan, NJ, US
Contact	JOHN D PAULSON
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/k912398/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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