

K860449 MODIFICATION TOP ULTRA THINWALL A-V FISTULA NEEDLEFeb 26, 1986
22 days to decisionK860449 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k860449/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Feb 4, 1986
Decision date	Feb 26, 1986
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Top Corp.
Location	Littleton, CO, US
Contact	TADAYOSHI IMAMURA
510(k) history	3 submissions · 3 cleared · 1986-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860449/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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