

K840837 TBI TRAINERMar 25, 1985
395 days to decisionK840837 · Product code: **FEJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k840837/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Attachment, Binocular, For Endoscope (FEJ)
Date received	Feb 24, 1984
Decision date	Mar 25, 1985
Days to decision	395 days
Third-party review	No

APPLICANT

Company	Manico
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1985-1985

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Device record: <https://www.510kdatabase.net/k840837/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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