FDA 510(k) Premarket Notification

K840837 TBI TRAINER

CLEARED

Mar 25, 1985
395 days to decision

K840837 · 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