

K823701 MOTILITY CATHETERSFeb 15, 1983
69 days to decisionK823701 · Product code: **FFX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k823701/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Gastrointestinal Motility (electrical) (FFX)
Date received	Dec 8, 1982
Decision date	Feb 15, 1983
Days to decision	69 days
Third-party review	No

APPLICANT

Company	Mui Scientific
Location	Mississauga, Ontario, CA
510(k) history	4 submissions · 4 cleared · 1983-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k823701/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 1, 2026