

K811973 FORDER RETRACTORJul 31, 1981
21 days to decisionK811973 · Product code: **KOA** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k811973/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Surgical Instruments, G-u, Manual (and Accessories) (KOA)
Date received	Jul 10, 1981
Decision date	Jul 31, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Surgemed Marketing, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k811973/> 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 July 3, 2026