FDA 510(k) Premarket Notification

K811973 FORDER RETRACTOR

CLEARED

Jul 31, 1981
21 days to decision

K811973 · Product code: **KOA** · Gastroenterology & Urology
Source: 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