FDA 510(k) Premarket Notification

K813488 CATHETER REGULAR

CLEARED

Dec 31, 1981
30 days to decision

K813488 · Product code: **FJS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k813488/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal, Long-term Indwelling (FJS)
Date received	Dec 1, 1981
Decision date	Dec 31, 1981
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Accurate Surgical Instruments Co.**
Location	Walker, MI, US
Contact	ACCURATE SURGICAL INSTRUMENTS CO
510(k) history	9 submissions · 9 cleared · 1981-1995