FDA 510(k) Premarket Notification

K771555 PERITONEAL DIALYSIS CATHETER

CLEARED

Nov 28, 1977
108 days to decision

K771555 · Product code: **FJS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k771555/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal, Long-term Indwelling (FJS)
Date received	Aug 12, 1977
Decision date	Nov 28, 1977
Days to decision	108 days
Third-party review	No

APPLICANT

Company	**Fluoro-Mold Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977