FDA 510(k) Premarket Notification

K771986 CATHETER, MEDIASTINAL&PERICARDIAL

CLEARED

Oct 28, 1977
9 days to decision

K771986 · Product code: **FHF** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k771986/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Evacuator, Fluid (FHF)
Date received	Oct 19, 1977
Decision date	Oct 28, 1977
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**The Minnesota Back Brace Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977