FDA 510(k) Premarket Notification

K791489 VITAL CATHETER

CLEARED

Oct 26, 1979
84 days to decision

K791489 · Product code: **FIE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k791489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Aug 3, 1979
Decision date	Oct 26, 1979
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Hospal Medical Corp.**
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1989