FDA 510(k) Premarket Notification

K770864 HEMODIALYSIS FISTULA SET

CLEARED

Jun 24, 1977
44 days to decision

K770864 · Product code: **FIE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k770864/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	May 11, 1977
Decision date	Jun 24, 1977
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**Venospital, Inc.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1977-1979