

K770864 HEMODIALYSIS FISTULA SETJun 24, 1977
44 days to decisionK770864 · Product code: **FIE** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k770864/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026