FDA 510(k) Premarket Notification

K813001 FIBRI-FILTER 20

CLEARED

Nov 6, 1981
11 days to decision

K813001 · Product code: **FPB** · General Hospital
Source: https://www.510kdatabase.net/k813001/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Oct 26, 1981
Decision date	Nov 6, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Tri-Med, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004