FDA 510(k) Premarket Notification

K811859 BACTASHIELD

CLEARED

Aug 7, 1981
37 days to decision

K811859 · Product code: **FIB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k811859/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Protector, Transducer, Dialysis (FIB)
Date received	Jul 1, 1981
Decision date	Aug 7, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

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