

K811859 BACTASHIELDAug 7, 1981
37 days to decisionK811859 · Product code: **FIB** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k811859/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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