

K770213 BARIORFeb 7, 1977
5 days to decisionK770213 · Product code: **KOC** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k770213/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Feb 2, 1977
Decision date	Feb 7, 1977
Days to decision	5 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/k770213/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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