FDA 510(k) Premarket Notification

K770213 BARIOR

CLEARED

Feb 7, 1977
5 days to decision

K770213 · Product code: **KOC** · Gastroenterology & Urology
Source: 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