FDA 510(k) Premarket Notification

K823881 ARTERIOVENOUS BLOOD TUBING SET FOR

CLEARED

Jan 17, 1983
21 days to decision

K823881 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k823881/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Dec 27, 1982
Decision date	Jan 17, 1983
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Amicon, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1976-1993