

K823881 ARTERIOVENOUS BLOOD TUBING SET FORJan 17, 1983
21 days to decisionK823881 · Product code: **KDI** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k823881/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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