

K813591 HEMOFILTERSMar 31, 1982
93 days to decisionK813591 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k813591/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Dec 28, 1981
Decision date	Mar 31, 1982
Days to decision	93 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/k813591/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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