

K840449 AMICON DIAFILTER 40. HEMOFILTERMar 23, 1984
51 days to decisionK840449 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k840449/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI) |
| Date received | Feb 1, 1984 |
| Decision date | Mar 23, 1984 |
| Days to decision | 51 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Medical Device Consultants, Inc. |
| Location | N. Attleboro, MA, US |
| 510(k) history | 14 submissions · 14 cleared · 1982-2002 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k840449/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026