

K802075 HOLLOW FIBER DIALYZER HOLDERSep 16, 1980
20 days to decisionK802075 · Product code: **FKI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k802075/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialyzer Holder (FKI)
Date received	Aug 27, 1980
Decision date	Sep 16, 1980
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Renal Devices, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1979-1981

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Device record: <https://www.510kdatabase.net/k802075/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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