

K801899 KEITHLEY LYMPHOCYTE DEPLETION CARTSep 26, 1980
49 days to decisionK801899 · Product code: **KDK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k801899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Transport, Kidney (KDK)
Date received	Aug 8, 1980
Decision date	Sep 26, 1980
Days to decision	49 days
Third-party review	No

APPLICANT

Company	Keithley Development Co.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1980

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Device record: <https://www.510kdatabase.net/k801899/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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