

K801829 MEDI-TRACE DISP. FLUSHING DEV./MONITORAug 27, 1980
27 days to decisionK801829 · Product code: **KRA** · CardiovascularSource: <https://www.510kdatabase.net/k801829/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Jul 31, 1980
Decision date	Aug 27, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Graphic Controls Corp.
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1998

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Device record: <https://www.510kdatabase.net/k801829/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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