

K801321 PACKAGE FOR MEDI-TRACE PHYSIOLOGICAL PREJun 30, 1980
25 days to decisionK801321 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k801321/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 5, 1980
Decision date	Jun 30, 1980
Days to decision	25 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/k801321/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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