

K780529 MYOCATH KITMay 9, 1978
36 days to decisionK780529 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k780529/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Apr 3, 1978
Decision date	May 9, 1978
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Sorensen Research
Location	Mchenry, IL, US
510(k) history	69 submissions · 69 cleared · 1976-1984

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Device record: <https://www.510kdatabase.net/k780529/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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