

K821816 ANGIOGRAPHIC CATHETERAug 24, 1982
64 days to decisionK821816 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k821816/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 21, 1982
Decision date	Aug 24, 1982
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Medi-Tech, Inc.
Location	Mchenry, IL, US
510(k) history	36 submissions · 35 cleared · 1978-1996

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Device record: <https://www.510kdatabase.net/k821816/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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