

K820648 ARROW M.I.S. MULTI-LUMEN CATHETERIZAApr 14, 1982
36 days to decisionK820648 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k820648/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Mar 9, 1982
Decision date	Apr 14, 1982
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Arrow Intl., Inc.
Location	Mchenry, IL, US
510(k) history	110 submissions · 105 cleared · 1976-2010

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Device record: <https://www.510kdatabase.net/k820648/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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