

**K862056 [CENTRAL VENOUS][MULTI-LUMEN] CATHETER
W/FLEX TIP**Aug 25, 1986
87 days to decisionK862056 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k862056/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	May 30, 1986
Decision date	Aug 25, 1986
Days to decision	87 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862056/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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