

**K945229 ARROW MULTIPOLAR INTRACARDIAC ELECTRODE  
CATHETERS**Aug 8, 1995  
286 days to decisionK945229 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k945229/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Oct 26, 1994
Decision date	Aug 8, 1995
Days to decision	286 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Arrow Intl., Inc.</b>
Location	Mchenry, IL, US
Contact	THOMAS D NICKEL
510(k) history	110 submissions · 105 cleared · 1976-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k945229/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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