

K953651 DIAGNOSTIC ELECTROPHYSIOLOGY CATHETER SYSTEMNov 30, 1995
118 days to decisionK953651 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k953651/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Aug 4, 1995
Decision date	Nov 30, 1995
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953651/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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