

K993691 ARROWG +ARD BLUE PLUS ANTIMICROBIAL MULTI-LUMEN CENTRAL VENOUS CATHETER

Mar 8, 2000
128 days to decision

K993691 · Product code: FOZ · General Hospital
Source: <https://www.510kdatabase.net/k993691/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Nov 1, 1999
Decision date	Mar 8, 2000
Days to decision	128 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.net

Device record: <https://www.510kdatabase.net/k993691/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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