FDA 510(k) Premarket Notification

K022662 DIATEK CANNON-CATHETER, MODEL CC5500

CLEARED

May 9, 2003
273 days to decision

K022662 · Product code: **MSD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k022662/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Aug 9, 2002
Decision date	May 9, 2003
Days to decision	273 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k022662/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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