FDA 510(k) Premarket Notification

K972264 FIRST MIDCATH CATHETER

CLEARED

Sep 15, 1997
90 days to decision

K972264 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k972264/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jun 17, 1997
Decision date	Sep 15, 1997
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Becton Dickinson Infusion Therapy Systems, Inc.**
Location	Sandy, UT, US
Contact	C.J. WELLE
510(k) history	36 submissions · 36 cleared · 1997-2026