FDA 510(k) Premarket Notification

CLEARED

K950301 ANGIOCATH(R) NONVALON(R) AND ANGIO-SET(R) IV CATHETERS

Apr 18, 1995
83 days to decision

K950301 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k950301/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jan 25, 1995
Decision date	Apr 18, 1995
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Becton Dickinson Vascular Access, Inc.**
Location	Sandy, UT, US
Contact	C J WELLE
510(k) history	25 submissions · 22 cleared · 1992-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k950301/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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