FDA 510(k) Premarket Notification

K860290 VIAPIC CENTRAL LINE CATHETER

CLEARED

Mar 18, 1986
49 days to decision

K860290 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k860290/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jan 28, 1986
Decision date	Mar 18, 1986
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Parke-Davis Co.**
Location	Mchenry, IL, US
Contact	E. F CLEARY
510(k) history	47 submissions · 47 cleared · 1976-1986