FDA 510(k) Premarket Notification

K821372 DESERET INTRACATH INTRAVENOUS CATHETER

CLEARED

Jun 2, 1982
23 days to decision

K821372 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k821372/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 10, 1982
Decision date	Jun 2, 1982
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Warner-Lambert Co.**
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003