FDA 510(k) Premarket Notification

K823565 MULTI-LUMEN INTRAVENOUS CATHETER

CLEARED

Dec 16, 1982
13 days to decision

K823565 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k823565/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Dec 3, 1982
Decision date	Dec 16, 1982
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Mcpherson Enterprises, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-2006