FDA 510(k) Premarket Notification

K890435 VITAACCESS CATHETER INTRODUCER KIT

CLEARED

Mar 29, 1989
61 days to decision

K890435 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k890435/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jan 27, 1989
Decision date	Mar 29, 1989
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Vitaphore Corp.**
Location	San Francisco, CA, US
Contact	JESSICA AYRES
510(k) history	13 submissions · 12 cleared · 1986-1992