FDA 510(k) Premarket Notification

K984189 ARGON MULTI-LUMEN CENTRAL VENOUS CATHETER

CLEARED

May 5, 1999
163 days to decision

K984189 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k984189/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Nov 23, 1998
Decision date	May 5, 1999
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Maxxim Medical**
Location	Arlington, TX, US
Contact	EDDIE MONROE
510(k) history	26 submissions · 25 cleared · 1994-2002