

**K984189 ARGON MULTI-LUMEN CENTRAL VENOUS
CATHETER**May 5, 1999
163 days to decisionK984189 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k984189/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026