FDA 510(k) Premarket Notification

K965030 BRANNON ARTERIO-VENOUS PORTSYRINGE

CLEARED

Mar 4, 1997
77 days to decision

K965030 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k965030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 17, 1996
Decision date	Mar 4, 1997
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vascular Logics, Inc.**
Location	Huntington Beach, CA, US
Contact	JAMES K BRANNON
510(k) history	1 submissions · 1 cleared · 1997-1997