FDA 510(k) Premarket Notification

K031718 LIFEVALVE CENTRAL VENOUS CATHTER

CLEARED

Jul 2, 2003
29 days to decision

K031718 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k031718/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 3, 2003
Decision date	Jul 2, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Horizon Medical Products, Inc.**
Location	Atlanta, GA, US
Contact	SCOTT MOELLER
510(k) history	16 submissions · 11 cleared · 1994-2003