FDA 510(k) Premarket Notification

K960431 PRESSURE STOPCOCK/MANIFOLD

CLEARED

Oct 29, 1996
272 days to decision

K960431 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k960431/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 31, 1996
Decision date	Oct 29, 1996
Days to decision	272 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Scientific Device Manufacturer, LLC**
Location	San Rafael, CA, US
Contact	RICHARD C BALL
510(k) history	6 submissions · 6 cleared · 1995-1997