FDA 510(k) Premarket Notification

K943770 LOW VOLUME MULTIPORT

CLEARED

Apr 10, 1995
251 days to decision

K943770 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k943770/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 2, 1994
Decision date	Apr 10, 1995
Days to decision	251 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**B. Braun of America, Inc.**
Location	Bethlehem, PA, US
Contact	MARK S ALSBERGE
510(k) history	19 submissions · 16 cleared · 1991-2000