FDA 510(k) Premarket Notification

K895480 BLUNT NEEDLE PORT/I.V. PORT

CLEARED

Dec 1, 1989
81 days to decision

K895480 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k895480/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 11, 1989
Decision date	Dec 1, 1989
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**Applied Medical Technologies**
Location	Mchenry, IL, US
Contact	DEAN SECREST
510(k) history	23 submissions · 20 cleared · 1981-1998