FDA 510(k) Premarket Notification

K780673 CENTRAL CATHETER KIT

CLEARED

May 16, 1978
22 days to decision

K780673 · Product code: **FPK** · General Hospital
Source: https://www.510kdatabase.net/k780673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubing, Fluid Delivery (FPK)
Date received	Apr 24, 1978
Decision date	May 16, 1978
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Argon Medical Corp.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1991