FDA 510(k) Premarket Notification

K823845 PORT-LOCK

CLEARED

Mar 1, 1983
70 days to decision

K823845 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k823845/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 21, 1982
Decision date	Mar 1, 1983
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Norfolk Medical Products, Inc.**
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1983-2020