

K823845 PORT-LOCKMar 1, 1983
70 days to decisionK823845 · 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

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Device record: <https://www.510kdatabase.net/k823845/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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