

K800638 PLASMA PRODUCTS CO. NON-VENTED INTRA/SETMay 8, 1980
45 days to decisionK800638 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k800638/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 24, 1980
Decision date	May 8, 1980
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Plasma Products Co.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800638/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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