

K832329 STERILE IV SET 4 SIZESAug 24, 1983
40 days to decisionK832329 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k832329/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 15, 1983
Decision date	Aug 24, 1983
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Abco Dealers, Inc.
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991

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Device record: <https://www.510kdatabase.net/k832329/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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