

K810921 MALE LUER LOCK PLUGSMay 21, 1981
45 days to decisionK810921 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k810921/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 6, 1981
Decision date	May 21, 1981
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Procedure Products, Inc.
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1981-2017

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Device record: <https://www.510kdatabase.net/k810921/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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