

K820384 GLASS SYRINGES, DRUG, LUER LOCKApr 15, 1982
65 days to decisionK820384 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k820384/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 9, 1982
Decision date	Apr 15, 1982
Days to decision	65 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/k820384/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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