

K821096 A-LINERMay 14, 1982
24 days to decisionK821096 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k821096/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 20, 1982
Decision date	May 14, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Martell Medical Products, Inc.
Location	Walker, MI, US
510(k) history	7 submissions · 7 cleared · 1981-1995

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Device record: <https://www.510kdatabase.net/k821096/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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