

K801430 CONTROL SYRINGEJul 8, 1980
21 days to decisionK801430 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k801430/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 17, 1980
Decision date	Jul 8, 1980
Days to decision	21 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/k801430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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