

K840490 FLUID REMOVAL ASSEMBLYApr 17, 1984
71 days to decisionK840490 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k840490/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 6, 1984
Decision date	Apr 17, 1984
Days to decision	71 days
Third-party review	No

APPLICANT

Company	Ackrad Laboratories
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1979-2002

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Device record: <https://www.510kdatabase.net/k840490/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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