FDA 510(k) Premarket Notification

K821671 DISPO. AES.

CLEARED

Jul 28, 1982
51 days to decision

K821671 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k821671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 7, 1982
Decision date	Jul 28, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Hiraoka New York, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k821671/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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